#### 108TH CONGRESS 1ST SESSION

## H. R. 2473

[Report No. 108- ]

To amend title XVIII of the Social Security Act to provide for a voluntary program for prescription drug coverage under the Medicare Program, to modernize the Medicare Program, and for other purposes.

#### IN THE HOUSE OF REPRESENTATIVES

June 16, 2003

Mr. Thomas (for himself and Mr. Tauzin) introduced the following bill; which was referred jointly to the Committees on Energy and Commerce, and Ways and Means

June , 2003

Reported from the Committee on Ways and Means with an amendment [Strike out all after the enacting clause and insert the part printed in italic]
[For text of introduced bill, see copy of bill as introduced on June 16, 2003]

### A BILL

To amend title XVIII of the Social Security Act to provide for a voluntary program for prescription drug coverage under the Medicare Program, to modernize the Medicare Program, and for other purposes.

- 1 Be it enacted by the Senate and House of Representa-
- 2 tives of the United States of America in Congress assembled,



1	SECTION 1. SHORT TITLE; AMENDMENTS TO SOCIAL SECU-
2	RITY ACT; REFERENCES TO BIPA AND SEC-
3	RETARY; TABLE OF CONTENTS.
4	(a) Short Title.—This Act may be cited as the
5	"Medicare Prescription Drug and Modernization Act of
6	2003".
7	(b) Amendments to Social Security Act.—Except
8	as otherwise specifically provided, whenever in this Act an
9	amendment is expressed in terms of an amendment to or
10	repeal of a section or other provision, the reference shall
11	be considered to be made to that section or other provision
12	of the Social Security Act.
13	(c) BIPA; Secretary.—In this Act:
14	(1) BIPA.—The term "BIPA" means the Medi-
15	care, Medicaid, and SCHIP Benefits Improvement
16	and Protection Act of 2000, as enacted into law by
17	section $1(a)(6)$ of Public Law 106-554.
18	(2) Secretary.—The term "Secretary" means
19	the Secretary of Health and Human Services.
20	(d) Table of Contents of contents of
21	this Act is as follows:
	Sec. 1. Short title; amendments to Social Security Act; references to BIPA and Secretary; table of contents.
	TITLE I—MEDICARE PRESCRIPTION DRUG BENEFIT
	Sec. 101. Establishment of a medicare prescription drug benefit.
	"Part D—Voluntary Prescription Drug Benefit Program

"Sec. 1860D-1. Benefits; eligibility; enrollment; and coverage period. "Sec. 1860D-2. Requirements for qualified prescription drug coverage.



- "Sec. 1860D-3. Beneficiary protections for qualified prescription drug coverage.
- "Sec. 1860D-4. Requirements for and contracts with prescription drug plan (PDP) sponsors.
- "Sec. 1860D-5. Process for beneficiaries to select qualified prescription drug coverage.
- "Sec. 1860D-6. Submission of bids and premiums.
- "Sec. 1860D-7. Premium and cost-sharing subsidies for low-income individuals.
- "Sec. 1860D-8. Subsidies for all medicare beneficiaries for qualified prescription drug coverage.
- "Sec. 1860D-9. Medicare Prescription Drug Trust Fund.
- "Sec. 1860D-10. Definitions; application to medicare advantage and EFFS programs; treatment of references to provisions in part C.
- Sec. 102. Offering of qualified prescription drug coverage under Medicare Advantage and enhanced fee-for-service (EFFS) program.
- Sec. 103. Medicaid amendments.
- "Sec. 1935. Special provisions relating to medicare prescription drug benefit.
- Sec. 104. Medigap transition.
- Sec. 105. Medicare prescription drug discount card endorsement program.
- Sec. 106. Disclosure of return information for purposes of carrying out medicare catastrophic prescription drug program.
- Sec. 107. State pharmaceutical assistance transition commission.

## TITLE II—MEDICARE ENHANCED FEE-FOR-SERVICE AND MEDICARE ADVANTAGE PROGRAMS; MEDICARE COMPETITION

Sec. 200. Medicare modernization and revitalization.

#### Subtitle A—Medicare Enhanced Fee-for-Service Program

Sec. 201. Establishment of enhanced fee-for-service (EFFS) program under medicare.

#### "Part E—Enhanced Fee-for-Service Program

- "Sec. 1860E-1. Offering of enhanced fee-for-service plans throughout the United States.
- "Sec. 1860E-2. Offering of enhanced fee-for-service (EFFS) plans.
- "Sec. 1860E-3. Submission of bids; beneficiary savings; payment of plans.
- "Sec. 1860E-4. Premiums; organizational and financial requirements; establishment of standards; contracts with EFFS organizations.

#### Subtitle B—Medicare Advantage Program

#### Chapter 1—Implementation Of Program

- Sec. 211. Implementation of medicare advantage program.
- Sec. 212. Medicare advantage improvements.

#### Chapter 2—Implementation Of Competition Program

Sec. 221. Competition program beginning in 2006.

#### Chapter 3—Additional Reforms

Sec. 231. Making permanent change in medicare advantage reporting deadlines and annual, coordinated election period.



- Sec. 232. Avoiding duplicative State regulation.
- Sec. 233. Specialized medicare advantage plans for special needs beneficiaries.
- Sec. 234. Medicare MSAs.
- Sec. 235. Extension of reasonable cost contracts.
- Sec. 236. Extension of municipal health service demonstration projects.

#### Subtitle C—Application of FEHBP-Style Competitive Reforms

Sec. 241. Application of FEHBP-style competitive reform beginning in 2010.

#### TITLE III—COMBATTING WASTE, FRAUD, AND ABUSE

- Sec. 301. Medicare secondary payor (MSP) provisions.
- Sec. 302. Competitive acquisition of certain items and services.
- Sec. 303. Competitive acquisition of covered outpatient drugs and biologicals.
- Sec. 304. Demonstration project for use of recovery audit contractors.

#### TITLE IV—RURAL HEALTH CARE IMPROVEMENTS

- Sec. 401. Enhanced disproportionate share hospital (DSH) treatment for rural hospitals and urban hospitals with fewer than 100 beds.
- Sec. 402. Immediate establishment of uniform standardized amount in rural and small urban areas.
- Sec. 403. Establishment of essential rural hospital classification.
- Sec. 404. More frequent update in weights used in hospital market basket.
- Sec. 405. Improvements to critical access hospital program.
- Sec. 406. Redistribution of unused resident positions.
- Sec. 407. Two-year extension of hold harmless provisions for small rural hospitals and sole community hospitals under prospective payment system for hospital outpatient department services.
- Sec. 408. Exclusion of certain rural health clinic and federally qualified health center services from the prospective payment system for skilled nursing facilities.
- Sec. 409. Recognition of attending nurse practitioners as attending physicians to serve hospice patients.
- Sec. 410. Improvement in payments to retain emergency capacity for ambulance services in rural areas.
- Sec. 411. Two-year increase for home health services furnished in a rural area.
- Sec. 412. Providing safe harbor for certain collaborative efforts that benefit medically underserved populations.
- Sec. 413. GAO study of geographic differences in payments for physicians' services
- Sec. 414. Treatment of missing cost reporting periods for sole community hospitals.
- Sec. 415. Extension of telemedicine demonstration project.
- Sec. 416. Adjustment to the medicare inpatient hospital PPS wage index to revise the labor-related share of such index.
- Sec. 417. Medicare incentive payment program improvements for physician scarcity.

#### TITLE V—PROVISIONS RELATING TO PART A

#### Subtitle A—Inpatient Hospital Services

- Sec. 501. Revision of acute care hospital payment updates.
- Sec. 502. Recognition of new medical technologies under inpatient hospital PPS.
- Sec. 503. Increase in Federal rate for hospitals in Puerto Rico.



- Sec. 504. Wage index adjustment reclassification reform .
- Sec. 505. MedPAC report on specialty hospitals.

#### Subtitle B—Other Provisions

- Sec. 511. Payment for covered skilled nursing facility services.
- Sec. 512. Coverage of hospice consultation services.

#### TITLE VI—PROVISIONS RELATING TO PART B

#### Subtitle A—Physicians' Services

- Sec. 601. Revision of updates for physicians' services.
- Sec. 602. Studies on access to physicians' services.
- Sec. 603. MedPAC report on payment for physicians' services.

#### Subtitle B—Preventive Services

- Sec. 611. Coverage of an initial preventive physical examination.
- Sec. 612. Coverage of cholesterol and blood lipid screening.
- Sec. 613. Waiver of deductible for colorectal cancer screening tests.
- Sec. 614. Improved payment for certain mammography services.

#### Subtitle C—Other Services

- Sec. 621. Hospital outpatient department (HOPD) payment reform.
- Sec. 622. Payment for ambulance services.
- Sec. 623. Renal dialysis services.
- Sec. 624. One-year moratorium on therapy caps; provisions relating to reports.
- Sec. 625. Adjustment to payments for services furnished in ambulatory surgical centers.
- Sec. 626. Payment for certain shoes and inserts under the fee schedule for orthotics and prosthetics.
- Sec. 627. Waiver of part B late enrollment penalty for certain military retirees; special enrollment period.
- Sec. 628. Part B deductible.
- Sec. 629. Extension of coverage of intravenous immune globulin (IVIG) for the treatment of primary immune deficiency diseases in the home.

#### TITLE VII—PROVISIONS RELATING TO PARTS A AND B

#### Subtitle A—Home Health Services

- Sec. 701. Update in home health services.
- Sec. 702. Establishment of reduced copayment for a home health service episode of care for certain beneficiaries.
- Sec. 703. MedPAC study on medicare margins of home health agencies.

#### Subtitle B—Direct Graduate Medical Education

Sec. 711. Extension of update limitation on high cost programs.

#### Subtitle C—Chronic Care Improvement

- Sec. 721. Voluntary chronic care improvement under traditional fee-for-service.
- Sec. 722. Chronic care improvement under medicare advantage and enhanced feefor-service programs.
- Sec. 723. Institute of Medicine report.
- Sec. 724. MedPAC report.



#### Subtitle D—Other Provisions

- Sec. 731. Modifications to medicare payment advisory commission (MedPAC).
- Sec. 732. Demonstration project for medical adult day care services.
- Sec. 733. Improvements in national and local coverage determination process to respond to changes in technology.
- Sec. 734. Treatment of certain physician pathology services.

#### TITLE VIII—MEDICARE BENEFITS ADMINISTRATION

Sec. 801. Establishment of Medicare Benefits Administration.

#### TITLE IX—REGULATORY REDUCTION AND CONTRACTING REFORM

#### Subtitle A—Regulatory Reform

Sec. 901. Construction; definition of supplier.

#### "Supplier

- Sec. 902. Issuance of regulations.
- Sec. 903. Compliance with changes in regulations and policies.
- Sec. 904. Reports and studies relating to regulatory reform.

#### Subtitle B—Contracting Reform

- Sec. 911. Increased flexibility in medicare administration.
- Sec. 912. Requirements for information security for medicare administrative contractors.

#### Subtitle C—Education and Outreach

- Sec. 921. Provider education and technical assistance.
- "Sec. 1889. Provider education and technical assistance.
- Sec. 922. Small provider technical assistance demonstration program.
- Sec. 923. Medicare Provider Ombudsman; Medicare Beneficiary Ombudsman.
- Sec. 924. Beneficiary outreach demonstration program.
- Sec. 925. Inclusion of additional information in notices to beneficiaries about skilled nursing facility benefits.
- Sec. 926. Information on medicare-certified skilled nursing facilities in hospital discharge plans.

#### Subtitle D—Appeals and Recovery

- Sec. 931. Transfer of responsibility for medicare appeals.
- Sec. 932. Process for expedited access to review.
- Sec. 933. Revisions to medicare appeals process.
- Sec. 934. Prepayment review.
- Sec. 935. Recovery of overpayments.
- Sec. 936. Provider enrollment process; right of appeal.
- Sec. 937. Process for correction of minor errors and omissions without pursuing appeals process.
- Sec. 938. Prior determination process for certain items and services; advance beneficiary notices.

#### Subtitle V—Miscellaneous Provisions

Sec. 941. Policy development regarding evaluation and management (E & M) documentation guidelines.



Sec. 942. Improvement in oversight of technology and coverage. Sec. 943. Treatment of hospitals for certain services under medicare secondary payor (MSP) provisions. Sec. 944. EMTALA improvements. Sec. 945. Emergency Medical Treatment and Active Labor Act (EMTALA) technical advisory group. Sec. 946. Authorizing use of arrangements to provide core hospice services in certain circumstances. Sec. 947. Application of osha bloodborne pathogens standard to certain hospitals. Sec. 948. BIPA-related technical amendments and corrections. Sec. 949. Conforming authority to waive a program exclusion. Sec. 950. Treatment of certain dental claims. Sec. 951. Furnishing hospitals with information to compute dsh formula. Sec. 952. Revisions to reassignment provisions. Sec. 953. Other provisions. Sec. 954. Temporary suspension of OASIS requirement for collection of data on

# 1 TITLE I—MEDICARE 2 PRESCRIPTION DRUG BENEFIT

non-medicare and non-medicaid patients.

3	SEC. 101. ESTABLISHMENT OF A MEDICARE PRESCRIPTION
4	DRUG BENEFIT.
5	(a) In General.—Title XVIII is amended—
6	(1) by redesignating part D as part F; and
7	(2) by inserting after part C the following new
8	part:
9	"Part D—Voluntary Prescription Drug Benefit
10	PROGRAM
11	"SEC. 1860D-1. BENEFITS; ELIGIBILITY; ENROLLMENT; AND
12	COVERAGE PERIOD.
13	"(a) Provision of Qualified Prescription Drug
14	Coverage Through Enrollment in Plans.—Subject to
15	the succeeding provisions of this part, each individual who
16	is entitled to benefits under part A or is enrolled under part



1	B is entitled to obtain qualified prescription drug coverage
2	(described in section $1860D-2(a)$ ) as follows:
3	"(1) Medicare-related plans.—
4	"(A) MEDICARE ADVANTAGE.—If the indi-
5	vidual is eligible to enroll in a Medicare Advan-
6	tage plan that provides qualified prescription
7	drug coverage under section 1851(j), the indi-
8	vidual may enroll in such plan and obtain cov-
9	erage through such plan.
10	"(B) EFFS PLANS.—If the individual is el-
11	igible to enroll in an EFFS plan that provides
12	qualified prescription drug coverage under part
13	E under section 1860 $E$ –2(d), the individual may
14	enroll in such plan and obtain coverage through
15	such plan.
16	"(C) MA-EFFS Plan; MA-EFFS RX
17	PLAN.—For purposes of this part, the term 'MA-
18	EFFS plan' means a Medicare Advantage plan
19	under part C and an EFFS plan under part E
20	and the term 'MA-EFFS Rx plan' means a MA-
21	EFFS plan insofar as such plan provides quali-
22	fied prescription drug coverage.
23	"(2) Prescription drug plan.—If the indi-
24	vidual is not enrolled in a MA-EFFS plan, the indi-



1	vidual may enroll under this part in a prescription
2	drug plan (as defined in section $1860D-10(a)(5)$ ).
3	Such individuals shall have a choice of such plans under
4	section $1860D-5(d)$ .
5	"(b) General Election Procedures.—
6	"(1) In general.—An individual eligible to
7	make an election under subsection (a) may elect to
8	enroll in a prescription drug plan under this part, or
9	elect the option of qualified prescription drug cov-
10	erage under a MA-EFFS Rx plan under part C or
11	part E, and to change such election only in such
12	manner and form as may be prescribed by regulations
13	of the Administrator of the Medicare Benefits Admin-
14	istration (appointed under section 1809(b)) (in this
15	part referred to as the 'Medicare Benefits Adminis-
16	trator') and only during an election period prescribed
17	in or under this subsection.
18	"(2) Election periods.—
19	"(A) In general.—Except as provided in
20	this paragraph, the election periods under this
21	subsection shall be the same as the coverage elec-
22	tion periods under the Medicare Advantage and
23	EFFS programs under section 1851(e),
24	including—



1	"(i) annual coordinated election peri-
2	ods; and
3	"(ii) special election periods.
4	In applying the last sentence of section
5	1851(e)(4) (relating to discontinuance of an elec-
6	tion during the first year of eligibility) under
7	this subparagraph, in the case of an election de-
8	scribed in such section in which the individual
9	had elected or is provided qualified prescription
10	drug coverage at the time of such first enroll-
11	ment, the individual shall be permitted to enroll
12	in a prescription drug plan under this part at
13	the time of the election of coverage under the
14	original fee-for-service plan.
15	"(B) Initial election periods.—
16	"(i) Individuals currently cov-
17	ERED.—In the case of an individual who is
18	entitled to benefits under part A or enrolled
19	under part B as of October 1, 2005, there
20	shall be an initial election period of 6
21	months beginning on that date.
22	"(ii) Individual covered in fu-
23	TURE.—In the case of an individual who is
24	first entitled to benefits under part $A$ or en-

rolled under part B after such date, there



1	shall be an initial election period which is
2	the same as the initial enrollment period
3	$under\ section\ 1837(d).$
4	"(C) Additional special election peri-
5	ods.—The Administrator shall establish special
6	election periods—
7	"(i) in cases of individuals who have
8	and involuntarily lose prescription drug
9	$coverage\ described\ in\ subsection\ (c)(2)(C);$
10	"(ii) in cases described in section
11	1837(h) (relating to errors in enrollment),
12	in the same manner as such section applies
13	to part B;
14	"(iii) in the case of an individual who
15	meets such exceptional conditions (including
16	conditions provided under section
17	1851(e)(4)(D)) as the Administrator may
18	provide; and
19	"(iv) in cases of individuals (as deter-
20	mined by the Administrator) who become el-
21	igible for prescription drug assistance under
22	$title\ XIX\ under\ section\ 1935(d).$
23	"(3) Information on plans.—Information de-
24	scribed in section 1860D-3(b)(1) on prescription drug
25	plans shall be made available during election periods.



1	"(c) Guaranteed Issue; Community Rating; and
2	Nondiscrimination.—
3	"(1) Guaranteed issue.—
4	"(A) In General.—An eligible individual
5	who is eligible to elect qualified prescription
6	drug coverage under a prescription drug plan or
7	MA-EFFS Rx plan at a time during which elec-
8	tions are accepted under this part with respect
9	to the plan shall not be denied enrollment based
10	on any health status-related factor (described in
11	section 2702(a)(1) of the Public Health Service
12	Act) or any other factor.
13	"(B) Medicare advantage limitations
14	PERMITTED.—The provisions of paragraphs (2)
15	and (3) (other than subparagraph (C)(i), relat-
16	ing to default enrollment) of section 1851(g) (re-
17	lating to priority and limitation on termination
18	of election) shall apply to PDP sponsors under
19	this subsection.
20	"(2) Community-rated premium.—
21	"(A) In General.—In the case of an indi-
22	vidual who enrolls under a prescription drug
23	plan or in a MA-EFFS Rx plan during the in-
24	dividual's initial enrollment period under this

part or maintains (as determined under sub-



paragraph (C)) continuous prescription drug coverage since the date the individual first quali-fies to elect prescription drug coverage under this part, a PDP sponsor or entity offering a pre-scription drug plan or MA-EFFS Rx plan and in which the individual is enrolled may not deny, limit, or condition the coverage or provi-sion of covered prescription drug benefits or vary or increase the premium under the plan based on any health status-related factor described in sec-tion 2702(a)(1) of the Public Health Service Act or any other factor. 

"(B) Late enrollment penalty.—In the case of an individual who does not maintain such continuous prescription drug coverage (as described in subparagraph (C)), a PDP sponsor or an entity offering a MA-EFFS Rx plan may (notwithstanding any provision in this title) adjust the premium otherwise applicable or impose a pre-existing condition exclusion with respect to qualified prescription drug coverage in a manner that reflects additional actuarial risk involved. Such a risk shall be established through an appropriate actuarial opinion of the type de-



1	scribed in subparagraphs (A) through (C) of sec-
2	$tion \ 2103(c)(4).$
3	"(C) Continuous prescription drug
4	coverage.—An individual is considered for
5	purposes of this part to be maintaining contin-
6	uous prescription drug coverage on and after the
7	date the individual first qualifies to elect pre-
8	scription drug coverage under this part if the in-
9	dividual establishes that as of such date the indi-
10	vidual is covered under any of the following pre-
11	scription drug coverage and before the date that
12	is the last day of the 63-day period that begins
13	on the date of termination of the particular pre-
14	scription drug coverage involved (regardless of
15	whether the individual subsequently obtains any
16	of the following prescription drug coverage):
17	"(i) Coverage under prescription
18	drug plan or ma-effs rx plan.—Quali-
19	fied prescription drug coverage under a pre-
20	scription drug plan or under a MA-EFFS
21	$Rx \ plan.$
22	"(ii) Medicaid prescription drug
23	COVERAGE.—Prescription drug coverage
24	under a medicaid plan under title XIX, in-
25	cluding through the Program of All-inclu-



1	sive Care for the Elderly (PACE) under sec-
2	tion 1934, through a social health mainte-
3	nance organization (referred to in section
4	4104(c) of the Balanced Budget Act of
5	1997), or through a demonstration project
6	under part C that demonstrates the applica-
7	tion of capitation payment rates for frail
8	elderly medicare beneficiaries through the
9	use of an interdisciplinary team and
10	through the provision of primary care serv-
11	ices to such beneficiaries by means of such
12	a team at the nursing facility involved.
13	"(iii) Prescription drug coverage
14	UNDER GROUP HEALTH PLAN.—Any out-
15	patient prescription drug coverage under a
16	group health plan, including a health bene-
17	fits plan under the Federal Employees
18	Health Benefit Plan under chapter 89 of
19	title 5, United States Code, and a qualified
20	retiree prescription drug plan as defined in
21	$section \ 1860D-8(f)(1), \ but \ only \ if \ (subject$
22	to $subparagraph\ (E)(ii))$ the $coverage\ pro-$
23	vides benefits at least equivalent to the bene-
24	fits under a qualified prescription drug

plan.



1	"(iv) Prescription drug coverage
2	UNDER CERTAIN MEDIGAP POLICIES.—Cov-
3	erage under a medicare supplemental policy
4	under section 1882 that provides benefits for
5	prescription drugs (whether or not such cov-
6	erage conforms to the standards for pack-
7	ages of benefits under section $1882(p)(1)$ ,
8	but only if the policy was in effect on Janu-
9	ary 1, 2006, and if (subject to subpara-
10	$graph\ (E)(ii))\ the\ coverage\ provides\ benefits$
11	at least equivalent to the benefits under a
12	qualified prescription drug plan.
13	"(v) State pharmaceutical assist-
14	ANCE PROGRAM.—Coverage of prescription
15	drugs under a State pharmaceutical assist-
16	ance program, but only if (subject to sub-
17	paragraph $(E)(ii))$ the coverage provides
18	benefits at least equivalent to the benefits
19	under a qualified prescription drug plan.
20	"(vi) Veterans' coverage of pre-
21	SCRIPTION DRUGS.—Coverage of prescrip-
22	tion drugs for veterans under chapter 17 of
23	title 38, United States Code, but only if
24	(subject to subparagraph $(E)(ii)$ ) the cov-

erage provides benefits at least equivalent to



1	the benefits under a qualified prescription
2	drug plan.
3	"(D) CERTIFICATION.—For purposes of car-
4	rying out this paragraph, the certifications of the
5	type described in sections 2701(e) of the Public
6	Health Service Act and in section 9801(e) of the
7	Internal Revenue Code shall also include a state-
8	ment for the period of coverage of whether the in-
9	dividual involved had prescription drug coverage
10	described in subparagraph (C).
11	"(E) Disclosure.—
12	"(i) In general.—Each entity that
13	offers coverage of the type described in
14	clause (iii), (iv), (v), or (vi) of subpara-
15	graph (C) shall provide for disclosure, con-
16	sistent with standards established by the
17	Administrator, of whether such coverage
18	provides benefits at least equivalent to the
19	benefits under a qualified prescription drug
20	plan.
21	"(ii) Waiver of limitations.—An
22	individual may apply to the Administrator
23	to waive the requirement that coverage of
24	such type provide benefits at least equiva-

lent to the benefits under a qualified pre-



1	scription drug plan, if the individual estab-
2	lishes that the individual was not ade-
3	quately informed that such coverage did not
4	provide such level of benefits.
5	"(F) Construction.—Nothing in this sec-
6	tion shall be construed as preventing the
7	disenrollment of an individual from a prescrip-
8	tion drug plan or a MA-EFFS Rx plan based on
9	the termination of an election described in sec-
10	tion $1851(g)(3)$ , including for non-payment of
11	premiums or for other reasons specified in sub-
12	section (d)(3), which takes into account a grace
13	period described in section $1851(g)(3)(B)(i)$ .
14	"(3) Nondiscrimination.—A PDP sponsor that
15	offers a prescription drug plan in an area designated
16	under section 1860D-4(b)(5) shall make such plan
17	available to all eligible individuals residing in the
18	area without regard to their health or economic status
19	or their place of residence within the area.
20	"(d) Effective Date of Elections.—
21	"(1) In general.—Except as provided in this
22	section, the Administrator shall provide that elections
23	under subsection (b) take effect at the same time as

the Administrator provides that similar elections



1	under section 1851(e) take effect under section
2	1851(f).
3	"(2) No election effective before 2006.—In
4	no case shall any election take effect before January
5	<i>1, 2006.</i>
6	"(3) Termination.—The Administrator shall
7	provide for the termination of an election in the case
8	of
9	"(A) termination of coverage under both
10	part A and part B; and
11	"(B) termination of elections described in
12	section $1851(g)(3)$ (including failure to pay re-
13	quired premiums).
14	"SEC. 1860D-2. REQUIREMENTS FOR QUALIFIED PRESCRIP-
15	TION DRUG COVERAGE.
16	"(a) Requirements.—
17	"(1) In general.—For purposes of this part
18	and part C and part E, the term 'qualified prescrip-
19	tion drug coverage' means either of the following:
20	"(A) Standard Coverage with access
21	TO NEGOTIATED PRICES.—Standard coverage (as
22	defined in subsection (b)) and access to nego-
23	tiated prices under subsection (d).
24	"(B) Actuarially equivalent coverage
25	WITH ACCESS TO NEGOTIATED PRICES—Con-



1	erage of covered outpatient drugs which meets the
2	alternative coverage requirements of subsection
3	(c) and access to negotiated prices under sub-
4	section (d), but only if it is approved by the Ad-
5	ministrator, as provided under subsection (c).
6	"(2) Permitting additional outpatient pre-
7	SCRIPTION DRUG COVERAGE.—
8	"(A) In general.—Subject to subpara-
9	graph (B), nothing in this part shall be con-
10	strued as preventing qualified prescription drug
11	coverage from including coverage of covered out-
12	patient drugs that exceeds the coverage required
13	under paragraph (1), but any such additional
14	coverage shall be limited to coverage of covered
15	outpatient drugs.
16	"(B) Disapproval authority.—The Ad-
17	ministrator shall review the offering of qualified
18	prescription drug coverage under this part or
19	part C or E. If the Administrator finds, in the
20	case of a qualified prescription drug coverage
21	under a prescription drug plan or a MA-EFFS
22	Rx plan, that the organization or sponsor offer-
23	ing the coverage is engaged in activities intended
24	to discourage enrollment of classes of eligible

beneficiaries

obtaining

coverage

medicare



1	through the plan on the basis of their higher like-
2	lihood of utilizing prescription drug coverage, the
3	Administrator may terminate the contract with
4	the sponsor or organization under this part or
5	$part\ C\ or\ E.$
6	"(3) Application of Secondary Payor Provi-
7	SIONS.—The provisions of section 1852(a)(4) shall
8	apply under this part in the same manner as they
9	apply under part C.
10	"(b) Standard Coverage.—For purposes of this
11	part, the 'standard coverage' is coverage of covered out-
12	patient drugs (as defined in subsection (f)) that meets the
13	following requirements:
14	"(1) Deductible.—The coverage has an annual
15	deductible—
16	"(A) for 2006, that is equal to \$250; or
17	"(B) for a subsequent year, that is equal to
18	the amount specified under this paragraph for
19	the previous year increased by the percentage
20	specified in paragraph (5) for the year involved.
21	Any amount determined under subparagraph (B)
22	that is not a multiple of \$10 shall be rounded to the
23	nearest multiple of \$10.
24	"(2) 80:20 BENEFIT STRUCTURE.—



1	"(A) 20 PERCENT COINSURANCE.—The cov-
2	erage has cost-sharing (for costs above the annual
3	deductible specified in paragraph (1) and up to
4	the initial coverage limit under paragraph (3))
5	that is—
6	"(i) equal to 20 percent; or
7	"(ii) is actuarially equivalent (using
8	processes established under subsection (e)) to
9	an average expected payment of 20 percent
10	of such costs.
11	"(B) Use of tiers.—Nothing in this part
12	shall be construed as preventing a PDP sponsor
13	from applying tiered copayments, so long as such
14	tiered copayments are consistent with subpara-
15	graph(A).
16	"(3) Initial coverage limit.—Subject to para-
17	graph (4), the coverage has an initial coverage limit
18	on the maximum costs that may be recognized for
19	payment purposes—
20	"(A) for 2006, that is equal to \$2,000; or
21	"(B) for a subsequent year, that is equal to
22	the amount specified in this paragraph for the
23	previous year, increased by the annual percent-
24	age increase described in paragraph (5) for the
25	year involved.



1	Any amount determined under subparagraph (B)
2	that is not a multiple of \$25 shall be rounded to the
3	nearest multiple of \$25.
4	"(4) Catastrophic protection.—
5	"(A) In general.—Notwithstanding para-
6	graph (3), the coverage provides benefits with no
7	cost-sharing after the individual has incurred
8	costs (as described in subparagraph (C)) for cov-
9	ered outpatient drugs in a year equal to the an-
10	nual out-of-pocket threshold specified in subpara-
11	graph(B).
12	"(B) Annual out-of-pocket thresh-
13	OLD.—
14	"(i) In general.—For purposes of
15	this part, the 'annual out-of-pocket thresh-
16	old' specified in this subparagraph is equal
17	to \$3,500 (subject to adjustment under
18	clause (ii) and subparagraph (D)).
19	"(ii) Inflation increase.—For a
20	year after 2006, the dollar amount specified
21	in clause (i) shall be increased by the an-
22	nual percentage increase described in para-
23	graph (5) for the year involved. Any
24	amount determined under the previous sen-



1	tence that is not a multiple of \$100 shall be
2	rounded to the nearest multiple of \$100.
3	"(C) Application.—In applying subpara-
4	graph(A)—
5	"(i) incurred costs shall only include
6	costs incurred for the annual deductible (de-
7	scribed in paragraph (1)), cost-sharing (de-
8	scribed in paragraph (2)), and amounts for
9	which benefits are not provided because of
10	the application of the initial coverage limit
11	described in paragraph (3); and
12	"(ii) such costs shall be treated as in-
13	curred only if they are paid by the indi-
14	vidual (or by another individual, such as a
15	family member, on behalf of the individual),
16	under section 1860D-7, under title XIX, or
17	under a State pharmaceutical assistance
18	program and the individual (or other indi-
19	vidual) is not reimbursed through insurance
20	or otherwise, a group health plan, or other
21	third-party payment arrangement (other
22	than under such title or such program) for
23	such costs.
24	"(D) Adjustment of annual out-of-
25	POCKET THRESHOLDS.—



1	"(i) In general.—For each enrollee
2	in a prescription drug plan or in a MA-
3	EFFS Rx plan whose adjusted gross income
4	exceeds the income threshold as defined in
5	clause (ii) for a year, the annual out-of-
6	pocket threshold otherwise determined under
7	subparagraph (B) for such year shall be in-
8	creased by an amount equal to the percent-
9	age specified in clause (iii), multiplied by
10	the lesser of—
11	"(I) the amount of such excess; or
12	"(II) the amount by which the in-
13	come threshold limit exceeds the income
14	threshold.
15	Any amount determined under the previous
16	sentence that is not a multiple of \$100 shall
17	be rounded to the nearest multiple of \$100.
18	"(ii) Income threshold.—For pur-
19	poses of clause (i)—
20	"(I) In general.—Subject to
21	subclause (II), the term 'income thresh-
22	old' means \$60,000 and the term 'in-
23	come threshold limit' means \$200,000.
24	"(II) Income inflation adjust-
25	MENT.—In the case of a year begin-



1	ning after 2006, each of the dollar
2	amounts in subclause (I) shall be in-
3	creased by an amount equal to such
4	dollar amount multiplied by the cost-
5	of-living adjustment determined under
6	section 1(f)(3) of the Internal Revenue
7	Code of 1986 for such year, determined
8	by substituting 'calendar year 2005'
9	for 'calendar year 1992'. If any
10	amount increased under the previous
11	sentence is not a multiple of \$100, such
12	amount shall be rounded to the nearest
13	$multiple\ of\ \$100.$
14	"(iii) Percentage.—The percentage
15	specified in this clause for a year is a frac-
16	tion (expressed as a percentage) equal to—
17	"(I) the annual out-of-pocket
18	threshold for a year under subpara-
19	graph (B) (determined without regard
20	to this subparagraph), divided by
21	"(II) the income threshold under
22	clause (ii) for that year.
23	If any percentage determined under the pre-
24	vious sentence that is not a multiple of
25	1/10th of 1 percentage point, such percentage



1	shall be rounded to the nearest multiple of
2	1/10th of 1 percentage point.
3	"(iv) Use of most recent return
4	INFORMATION.—For purposes of clause (i)
5	for an enrollee for a year, except as pro-
6	vided in clause (v), the adjusted gross in-
7	come of an individual shall be based on the
8	most recent information disclosed to the
9	Secretary under section 6109(l)(19) of the
10	Internal Revenue Code of 1986 before the be-
11	ginning of that year.
12	"(v) Individual election to
13	PRESENT MOST RECENT INFORMATION RE-
14	GARDING INCOME.—The Secretary shall pro-
15	vide, in coordination with the Secretary of
16	the Treasury, a procedure under which, for
17	purposes of applying this subparagraph for
18	a calendar year, instead of using the infor-
19	mation described in clause (iv), an enrollee
20	may elect to use more recent information,
21	including information with respect to a tax-
22	able year ending in such calendar year.
23	Such process shall—
24	"(I) require the enrollee to provide
25	the Secretary with a copy of the rel-



1	evant portion of the more recent return
2	to be used under this clause;
3	"(II) provide for the Medicare
4	Beneficiary Ombudsman (under sec-
5	tion 1810) offering assistance to such
6	enrollees in presenting such informa-
7	tion and the toll-free number under
8	such section being a point of contact
9	for beneficiaries to inquire as to how to
10	present such information;
11	"(III) provide for the verification
12	of the information in such return by
13	the Secretary of the Treasury under
14	section 6103(l)(19) of the Internal Rev-
15	enue Code of 1986; and
16	"(IV) provide for the payment by
17	the Secretary (in a manner specified
18	by the Secretary) to the enrollee of an
19	amount equal to the excess of the ben-
20	efit payments that would have been
21	payable under the plan if the more re-
22	cent return information were used,
23	over the benefit payments that were

made under the plan.



1	In the case of a payment under subclause
2	(III) for an enrollee under a prescription
3	drug plan, the PDP sponsor of the plan
4	shall pay to the Secretary the amount so
5	paid, less the applicable reinsurance
6	amount that would have applied under sec-
7	tion 1860D-8(c)(1)(B) if such payment had
8	been treated as an allowable cost under such
9	section. Such plan payment shall be depos-
10	ited in the Treasury to the credit of the
11	Medicare Prescription Drug Account in the
12	Federal Supplementary Medical Insurance
13	Trust Fund (under section 1841).
13 14	Trust Fund (under section 1841). "(vi) Dissemination of information
14	"(vi) Dissemination of information
14 15	"(vi) Dissemination of information on process.—The Secretary shall provide,
<ul><li>14</li><li>15</li><li>16</li></ul>	"(vi) DISSEMINATION OF INFORMATION ON PROCESS.—The Secretary shall provide, through the annual medicare handbook
<ul><li>14</li><li>15</li><li>16</li><li>17</li></ul>	"(vi) DISSEMINATION OF INFORMATION ON PROCESS.—The Secretary shall provide, through the annual medicare handbook under section 1804(a), for a general descrip-
<ul><li>14</li><li>15</li><li>16</li><li>17</li><li>18</li></ul>	"(vi) DISSEMINATION OF INFORMATION ON PROCESS.—The Secretary shall provide, through the annual medicare handbook under section 1804(a), for a general descrip- tion of the adjustment of annual out-of-
<ul><li>14</li><li>15</li><li>16</li><li>17</li><li>18</li><li>19</li></ul>	"(vi) DISSEMINATION OF INFORMATION ON PROCESS.—The Secretary shall provide, through the annual medicare handbook under section 1804(a), for a general descrip- tion of the adjustment of annual out-of- pocket thresholds provided under this sub-
14 15 16 17 18 19 20	"(vi) DISSEMINATION OF INFORMATION ON PROCESS.—The Secretary shall provide, through the annual medicare handbook under section 1804(a), for a general descrip- tion of the adjustment of annual out-of- pocket thresholds provided under this sub- paragraph, including the process for adjust-
14 15 16 17 18 19 20 21	"(vi) DISSEMINATION OF INFORMATION ON PROCESS.—The Secretary shall provide, through the annual medicare handbook under section 1804(a), for a general descrip- tion of the adjustment of annual out-of- pocket thresholds provided under this sub- paragraph, including the process for adjust- ment based upon more recent information

forth the amount of the adjustment that is



1	made under clause (i) based on the amount
2	of an enrollee's adjusted gross income.
3	"(E) Requesting information on en-
4	ROLLEES.—
5	"(i) In General.—The Secretary
6	shall, periodically as required to carry out
7	subparagraph (D), transmit to the Sec-
8	retary of the Treasury a list of the names
9	and TINs of enrollees in prescription drug
10	plans (or in MA-EFFS Rx plans) and re-
11	quest that such Secretary disclose to the
12	Secretary information under subparagraph
13	(A) of section 6103(l)(19) of the Internal
14	Revenue Code of 1986 with respect to those
15	enrollees for a specified taxable year for ap-
16	plication in a particular calendar year.
17	"(ii) Disclosure to plan spon-
18	sors.—In the case of a specified taxpayer
19	(as defined in section $6103(l)(19)(B)$ of the
20	Internal Revenue Code of 1986) who is en-
21	rolled in a prescription drug plan or in an
22	MA-EFFS Rx plan, the Secretary shall dis-
23	close to the entity that offers the plan the
24	annual out-of-pocket threshold applicable to
25	such individual under subparagraph (D).



1	"(F) Maintaining confidentiality of in-
2	FORMATION.—
3	"(i) In general.—The amount of any
4	increase in an annual out-of-pocket thresh-
5	old under subparagraph (D) may not be
6	disclosed by the Secretary except to a PDP
7	sponsor or entity that offers a MA-EFFS
8	Rx plan to the extent necessary to carry out
9	this part.
10	"(ii) Criminal and civil penalties
11	for unauthorized disclosure.—A per-
12	son who makes an unauthorized disclosure
13	of information disclosed under section
14	6103(l)(19) of the Internal Revenue Code of
15	1986 (including disclosure of any increase
16	in an annual out-of-pocket threshold under
17	subparagraph (D)) shall be subject to pen-
18	alty to the extent provided under—
19	"(I) section 7213 of such Code (re-
20	lating to criminal penalty for unau-
21	$thorized\ disclosure\ of\ information);$
22	"(II) section 7213A of such Code
23	(relating to criminal penalty for unau-
24	thorized inspection of returns or return
25	information);



1	"(III) section 7431 of such Code
2	(relating to civil damages for unau-
3	thorized inspection or disclosure of re-
4	turns and return information);
5	"(IV) any other provision of the
6	Internal Revenue Code of 1986; or
7	"(V) any other provision of law.
8	"(iii) Application of additional
9	CIVIL MONETARY PENALTY FOR UNAUTHOR-
10	ized disclosures.—In addition to any
11	penalty otherwise provided under law, any
12	person who makes an unauthorized disclo-
13	sure of such information shall be subject to
14	a civil monetary penalty of not to exceed
15	\$10,000 for each such unauthorized disclo-
16	sure. The provisions of section 1128A (other
17	than subsections (a) and (b)) shall apply to
18	civil money penalties under this subpara-
19	graph in the same manner as they apply to
20	a penalty or proceeding under section
21	1128A(a).
22	"(5) Annual percentage increase.—For pur-
23	poses of this part, the annual percentage increase
24	specified in this paragraph for a year is equal to the
25	annual percentage increase in average per capita ag-



1	gregate expenditures for covered outpatient drugs in
2	the United States for medicare beneficiaries, as deter-
3	mined by the Administrator for the 12-month period
4	ending in July of the previous year.
5	"(c) Alternative Coverage Requirements.—A
6	prescription drug plan or MA-EFFS Rx plan may provide
7	a different prescription drug benefit design from the stand-
8	ard coverage described in subsection (b) so long as the Ad-
9	ministrator determines (based on an actuarial analysis by
10	the Administrator) that the following requirements are met
11	and the plan applies for, and receives, the approval of the
12	Administrator for such benefit design:
13	"(1) Assuring at least actuarially equiva-
14	LENT COVERAGE.—
15	"(A) Assuring equivalent value of
16	TOTAL COVERAGE.—The actuarial value of the
17	total coverage (as determined under subsection
18	(e)) is at least equal to the actuarial value (as
19	so determined) of standard coverage.
20	"(B) Assuring equivalent unsubsidized
21	VALUE OF COVERAGE.—The unsubsidized value
22	of the coverage is at least equal to the unsub-
23	sidized value of standard coverage. For purposes
24	of this subparagraph, the unsubsidized value of

coverage is the amount by which the actuarial



1	value of the coverage (as determined under sub-
2	section (e)) exceeds the actuarial value of the sub-
3	sidy payments under section 1860D-8 with re-
4	spect to such coverage.
5	"(C) Assuring standard payment for
6	COSTS AT INITIAL COVERAGE LIMIT.—The cov-
7	erage is designed, based upon an actuarially rep-
8	resentative pattern of utilization (as determined
9	under subsection (e)), to provide for the pay-
10	ment, with respect to costs incurred that are
11	equal to the initial coverage limit under sub-
12	section (b)(3), of an amount equal to at least the
13	product of—
14	"(i) the amount by which the initial
15	coverage limit described in subsection (b)(3)
16	exceeds the deductible described in sub-
17	section (b)(1); and
18	"(ii) 100 percent minus the cost-shar-
19	ing percentage specified in subsection
20	(b)(2)(A)(i).
21	"(2) Catastrophic protection.—The coverage
22	provides for beneficiaries the catastrophic protection
23	described in subsection (b)(4).
24	"(d) Access to Negotiated Prices.—



"(1) In general.—Under qualified prescription
drug coverage offered by a PDP sponsor or an entity
offering a MA-EFFS Rx plan, the sponsor or entity
shall provide beneficiaries with access to negotiated
prices (including applicable discounts) used for pay-
ment for covered outpatient drugs, regardless of the
fact that no benefits may be payable under the cov-
erage with respect to such drugs because of the appli-
cation of cost-sharing or an initial coverage limit (de-
scribed in subsection (b)(3)). Insofar as a State elects
to provide medical assistance under title XIX to a
beneficiary enrolled under such title and under a pre-
scription drug plan or MA-EFFS Rx plan for a drug
based on the prices negotiated by a prescription drug
plan or MA-EFFS Rx plan under this part, the re-
quirements of section 1927 shall not apply to such
drugs. The prices negotiated by a prescription drug
plan under this part, by a MA-EFFS Rx plan with
respect to covered outpatient drugs, or by a qualified
retiree prescription drug plan (as defined in section
1860D-8(f)(1)) with respect to such drugs on behalf
of individuals entitled to benefits under part $A$ or en-
rolled under part B, shall (notwithstanding any other
provision of law) not be taken into account for the



1	purposes of establishing the best price under section
2	1927(c)(1)(C).
3	"(2) Disclosure.—The PDP sponsor or entity
4	offering a MA-EFFS Rx plan shall disclose to the Ad-
5	ministrator (in a manner specified by the Adminis-
6	trator) the extent to which discounts or rebates or
7	other remuneration or price concessions made avail-
8	able to the sponsor or organization by a manufacturer
9	are passed through to enrollees through pharmacies
10	and other dispensers or otherwise. The provisions of
11	section $1927(b)(3)(D)$ shall apply to information dis-
12	closed to the Administrator under this paragraph in
13	the same manner as such provisions apply to infor-
14	mation disclosed under such section.
15	"(3) Audits and reports.—To protect against
16	fraud and abuse and to ensure proper disclosures and
17	accounting under this part, in addition to any pro-
18	tections against fraud and abuse provided under sec-
19	tion $1860D-4(b)(3)(C)$ , the Administrator may peri-
20	odically audit the financial statements and records of
21	PDP sponsor or entities offering a MA-EFFS Rx
22	plan.
23	"(e) Actuarial Valuation; Determination of An-
24	NUAL PERCENTAGE INCREASES.—



1	"(1) Processes.—For purposes of this section,
2	the Administrator shall establish processes and
3	methods—
4	"(A) for determining the actuarial valu-
5	ation of prescription drug coverage, including—
6	"(i) an actuarial valuation of standard
7	coverage and of the reinsurance subsidy
8	payments under section 1860D-8;
9	"(ii) the use of generally accepted actu-
10	arial principles and methodologies; and
11	"(iii) applying the same methodology
12	for determinations of alternative coverage
13	under subsection (c) as is used with respect
14	to determinations of standard coverage
15	under subsection (b); and
16	"(B) for determining annual percentage in-
17	$creases\ described\ in\ subsection\ (b) (5).$
18	"(2) USE OF OUTSIDE ACTUARIES.—Under the
19	processes under paragraph (1)(A), PDP sponsors and
20	entities offering MA-EFFS Rx plans may use actu-
21	arial opinions certified by independent, qualified ac-
22	tuaries to establish actuarial values, but the Adminis-
23	trator shall determine whether such actuarial values
24	$meet\ the\ requirements\ under\ subsection\ (c)$ (1).
25	"(f) Covered Outpatient Drugs Defined.—



1	"(1) In general.—Except as provided in this
2	subsection, for purposes of this part, the term 'covered
3	outpatient drug' means—
4	"(A) a drug that may be dispensed only
5	upon a prescription and that is described in sub-
6	paragraph (A)(i) or (A)(ii) of section 1927(k)(2);
7	or
8	"(B) a biological product described in
9	clauses (i) through (iii) of subparagraph (B) of
10	such section or insulin described in subpara-
11	graph (C) of such section,
12	and such term includes a vaccine licensed under sec-
13	tion 351 of the Public Health Service Act and any
14	use of a covered outpatient drug for a medically ac-
15	cepted indication (as defined in section $1927(k)(6)$ ).
16	"(2) Exclusions.—
17	"(A) In general.—Such term does not in-
18	clude drugs or classes of drugs, or their medical
19	uses, which may be excluded from coverage or
20	otherwise $restricted$ $under$ $section$ $1927(d)(2),$
21	other than subparagraph (E) thereof (relating to
22	smoking cessation agents), or under section
23	1927(d)(3).
24	"(B) Avoidance of duplicate cov-
25	ERAGE —A drug prescribed for an individual



1	that would otherwise be a covered outpatient
2	drug under this part shall not be so considered
3	if payment for such drug is available under part
4	A or B for an individual entitled to benefits
5	$under\ part\ A\ and\ enrolled\ under\ part\ B.$
6	"(3) Application of formulary restric-
7	TIONS.—A drug prescribed for an individual that
8	would otherwise be a covered outpatient drug under
9	this part shall not be so considered under a plan is
10	the plan excludes the drug under a formulary and
11	such exclusion is not successfully appealed under sec-
12	$tion \ 1860D-3(f)(2).$
13	"(4) Application of general exclusion pro-
14	VISIONS.—A prescription drug plan or MA-EFFS Rx
15	plan may exclude from qualified prescription drug
16	coverage any covered outpatient drug—
17	"(A) for which payment would not be made
18	if section 1862(a) applied to part D; or
19	"(B) which are not prescribed in accordance
20	with the plan or this part.
21	Such exclusions are determinations subject to recon-
22	sideration and appeal pursuant to section 1860D-
23	3(f).



1	"SEC. 1860D-3. BENEFICIARY PROTECTIONS FOR QUALIFIED
2	PRESCRIPTION DRUG COVERAGE.
3	"(a) Guaranteed Issue, Community-Rated Pre-
4	MIUMS, ACCESS TO NEGOTIATED PRICES, AND NON-
5	DISCRIMINATION.—For provisions requiring guaranteed
6	issue, community-rated premiums, access to negotiated
7	prices, and nondiscrimination, see sections 1860D-1(c)(1),
8	1860D-1(c)(2), $1860D-2(d)$ , and $1860D-6(b)$ , respectively.
9	"(b) Dissemination of Information.—
10	"(1) General information.—A PDP sponsor
11	shall disclose, in a clear, accurate, and standardized
12	form to each enrollee with a prescription drug plan
13	offered by the sponsor under this part at the time of
14	enrollment and at least annually thereafter, the infor-
15	$mation\ described\ in\ section\ 1852(c)(1)\ relating\ to$
16	such plan. Such information includes the following:
17	"(A) Access to specific covered outpatient
18	drugs, including access through pharmacy net-
19	works.
20	"(B) How any formulary used by the spon-
21	sor functions, including the drugs included in
22	$the\ formulary.$
23	"(C) Co-payments and deductible require-
24	ments, including the identification of the tiered
25	or other co-payment level applicable to each drug
26	(or class of drugs).



1	"(D) Grievance and appeals procedures.
2	Such information shall also be made available upon
3	request to prospective enrollees.
4	"(2) Disclosure upon request of general
5	COVERAGE, UTILIZATION, AND GRIEVANCE INFORMA-
6	TION.—Upon request of an individual eligible to en-
7	roll under a prescription drug plan, the PDP sponsor
8	shall provide the information described in section
9	1852(c)(2) (other than subparagraph (D)) to such in-
10	dividual.
11	"(3) Response to beneficiary questions.—
12	Each PDP sponsor offering a prescription drug plan
13	shall have a mechanism for providing specific infor-
14	mation to enrollees upon request. The sponsor shall
15	make available on a timely basis, through an Internet
16	website and in writing upon request, information on
17	specific changes in its formulary.
18	"(4) Claims information.—Each PDP sponsor
19	offering a prescription drug plan must furnish to
20	each enrollee in a form easily understandable to such
21	enrollees an explanation of benefits (in accordance
22	with section 1806(a) or in a comparable manner) and
23	a notice of the benefits in relation to initial coverage
24	limit and the annual out-of-pocket threshold applica-

ble to such enrollee for the current year, whenever pre-



1	scription drug benefits are provided under this part
2	(except that such notice need not be provided more
3	often than monthly).
4	"(c) Access to Covered Benefits.—
5	"(1) Assuring pharmacy access.—
6	"(A) Participation of any willing
7	PHARMACY.—A PDP sponsor and an entity of-
8	fering a MA-EFFS Rx plan shall permit the
9	participation of any pharmacy that meets terms
10	and conditions that the plan has established.
11	"(B) Discounts allowed for network
12	PHARMACIES.—A prescription drug plan and a
13	MA-EFFS Rx plan may, notwithstanding sub-
14	paragraph (A), reduce coinsurance or copay-
15	ments for its enrolled beneficiaries below the level
16	otherwise provided for covered outpatient drugs
17	dispensed through in-network pharmacies, but in
18	no case shall such a reduction result in an in-
19	crease in payments made by the Administrator
20	under section 1860D-8 to a plan.
21	"(C) Convenient access for network
22	PHARMACIES.—The PDP sponsor of the prescrip-
23	tion drug plan and the entity offering a MA-
24	EFFS Rx plan shall secure the participation in

its network of a sufficient number of pharmacies



1	that dispense (other than by mail order) drugs
2	directly to patients to ensure convenient access
3	(consistent with rules of the Administrator). The
4	Administrator shall establish convenient access
5	rules under this subparagraph that are no less
6	favorable to enrollees than the rules for conven-
7	ient access to pharmacies of the Secretary of De-
8	fense established as of June 1, 2003, for purposes
9	of the TRICARE Retail Pharmacy (TRRx) pro-
10	gram. Such rules shall include adequate emer-
11	gency access for enrolled beneficiaries.
12	"(D) Level playing field.—Such a spon-
13	sor shall permit enrollees to receive benefits
14	(which may include a 90-day supply of drugs or
15	biologicals) through a community pharmacy,
16	rather than through mail order, with any dif-
17	ferential in cost paid by such enrollees.
18	"(E) Not required to accept insurance
19	RISK.—The terms and conditions under subpara-
20	graph (A) may not require participating phar-
21	macies to accept insurance risk as a condition of
22	participation.
23	"(2) Use of standardized technology.—
24	"(A) In general.—The PDP sponsor of a

prescription drug plan and an entity offering a



1	MA-EFFS Rx plan shall issue (and reissue, as
2	appropriate) such a card (or other technology)
3	that may be used by an enrollee to assure access
4	to negotiated prices under section 1860D-2(d)
5	for the purchase of prescription drugs for which
6	coverage is not otherwise provided under the
7	plan.
8	"(B) Standards.—
9	"(i) Development.—The Adminis-
10	trator shall provide for the development or
11	utilization of uniform standards relating to
12	a standardized format for the card or other
13	technology referred to in subparagraph $(A)$ .
14	Such standards shall be compatible with
15	standards established under part C of title
16	XI.
17	"(ii) Application of advisory task
18	FORCE.—The advisory task force established
19	$under\ subsection\ (d)(3)(B)(ii)\ shall\ provide$
20	recommendations to the Administrator
21	under such subsection regarding the stand-
22	ards developed under clause (i).
23	"(3) Requirements on development and ap-
24	PLICATION OF FORMULARIES.—If a PDP sponsor of a
25	prescription drug plan or an entity offering a MA-



1	EFFS Rx plan uses a formulary, the following re-
2	quirements must be met:
3	"(A) Pharmacy and therapeutic (P&T)
4	committee.—The sponsor or entity must estab-
5	lish a pharmacy and therapeutic committee that
6	develops and reviews the formulary. Such com-
7	mittee shall include at least one practicing phy-
8	sician and at least one practicing pharmacist
9	independent and free of conflict with respect to
10	the committee both with expertise in the care of
11	elderly or disabled persons and a majority of its
12	members shall consist of individuals who are
13	practicing physicians or practicing pharmacists
14	$(or\ both).$
15	"(B) Formulary development.—In de-
16	veloping and reviewing the formulary, the com-
17	mittee shall—
18	"(i) base clinical decisions on the
19	strength of scientific evidence and standards
20	of practice, including assessing peer-re-
21	viewed medical literature, such as random-
22	ized clinical trials, pharmacoeconomic stud-
23	ies, outcomes research data, and such other
24	information as the committee determines to

be appropriate; and



1	"(ii) shall take into account whether
2	including in the formulary particular cov-
3	ered outpatient drugs has therapeutic ad-
4	vantages in terms of safety and efficacy.
5	"(C) Inclusion of drugs in all thera-
6	PEUTIC CATEGORIES.—The formulary must in-
7	clude drugs within each therapeutic category and
8	class of covered outpatient drugs (although not
9	necessarily for all drugs within such categories
10	and classes). In establishing such classes, the
11	committee shall take into account the standards
12	published in the United States Pharmacopeia-
13	Drug Information. The committee shall make
14	available to the enrollees under the plan through
15	the Internet or otherwise the bases for the exclu-
16	sion of coverage of any drug from the formulary.
17	"(D) Provider and patient edu-
18	CATION.—The committee shall establish policies
19	and procedures to educate and inform health
20	care providers and enrollees concerning the for-
21	mulary.
22	"(E) Notice before removing drug
23	FROM FORMULARY FOR CHANGING PREFERRED
24	OR TIER STATUS OF DRUG.—Any removal of a

covered outpatient drug from a formulary and



1	any change in the preferred or tier cost-sharing
2	status of such a drug shall take effect only after
3	appropriate notice is made available to bene-
4	ficiaries and physicians.
5	"(F) Periodic evaluation of proto-
6	cols.—In connection with the formulary, a pre-
7	scription drug plan shall provide for the periodic
8	evaluation and analysis of treatment protocols
9	and procedures.
10	"(G) Grievances and appeals relating
11	TO APPLICATION OF FORMULARIES.—For provi-
12	sions relating to grievances and appeals of cov-
13	erage, see subsections (e) and (f).
14	"(d) Cost and Utilization Management; Quality
15	Assurance; Medication Therapy Management Pro-
16	GRAM.—
17	"(1) In general.—The PDP sponsor or entity
18	offering a MA-EFFS Rx plan shall have in place, di-
19	rectly or through appropriate arrangements, with re-
20	spect to covered outpatient drugs—
21	"(A) an effective cost and drug utilization
22	management program, including medically ap-
23	propriate incentives to use generic drugs and
24	therapeutic interchange, when appropriate;



1	"(B) quality assurance measures and sys-
2	tems to reduce medical errors and adverse drug
3	interactions, including side-effects, and improve
4	medication use, including a medication therapy
5	management program described in paragraph
6	(2) and for years beginning with 2007, an elec-
7	tronic prescription program described in para-
8	graph (3); and
9	"(C) a program to control fraud, abuse, and
10	waste.
11	Nothing in this section shall be construed as impair-
12	ing a PDP sponsor or entity from utilizing cost man-
13	agement tools (including differential payments) under
14	all methods of operation.
15	"(2) Medication therapy management pro-
16	GRAM.—
17	"(A) In General.—A medication therapy
18	management program described in this para-
19	graph is a program of drug therapy management
20	and medication administration that may be fur-
21	nished by a pharmacy provider and that is de-
22	signed to assure, with respect to beneficiaries at
23	risk for potential medication problems, such as
24	beneficiaries with complex or chronic diseases

(such as diabetes, asthma, hypertension, and



1	congestive heart failure) or multiple prescrip-
2	tions, that covered outpatient drugs under the
3	prescription drug plan are appropriately used to
4	optimize therapeutic outcomes through improved
5	medication use and reduce the risk of adverse
6	events, including adverse drug interactions. Such
7	programs may distinguish between services in
8	ambulatory and institutional settings.
9	"(B) Elements.—Such program may
10	include—
11	"(i) enhanced beneficiary under-
12	standing to promote the appropriate use of
13	medications by beneficiaries and to reduce
14	the risk of potential adverse events associ-
15	ated with medications, through beneficiary
16	education, counseling, case management,
17	disease state management programs, and
18	other appropriate means;
19	"(ii) increased beneficiary adherence
20	with prescription medication regimens
21	through medication refill reminders, special
22	packaging, and other compliance programs
23	and other appropriate means; and
24	"(iii) detection of patterns of overuse
25	and underuse of prescription drugs.



1	"(C) Development of program in co-
2	OPERATION WITH LICENSED PHARMACISTS.—The
3	program shall be developed in cooperation with
4	licensed and practicing pharmacists and physi-
5	cians.
6	"(D) Considerations in pharmacy
7	FEES.—The PDP sponsor of a prescription drug
8	program and an entity offering a MA-EFFS Rx
9	plan shall take into account, in establishing fees
10	for pharmacists and others providing services
11	under the medication therapy management pro-
12	gram, the resources and time used in imple-
13	menting the program. Each such sponsor or enti-
14	ty shall disclose to the Administrator upon re-
15	quest the amount of any such management or
16	dispensing fees.
17	"(3) Electronic prescription program.—
18	"(A) In General.—An electronic prescrip-
19	tion drug program described in this paragraph
20	is a program that includes at least the following
21	components, consistent with uniform standards
22	established under subparagraph (B):
23	"(i) Electronic transmittal of
24	PRESCRIPTIONS.—Prescriptions must be

written and transmitted electronically



1	(other than by facsimile), except in emer-
2	gency cases and other exceptional cir-
3	cumstances recognized by the Adminis-
4	trator.
5	"(ii) Provision of information to
6	PRESCRIBING HEALTH CARE PROFES-
7	SIONAL.—The program provides for the elec-
8	tronic transmittal to the prescribing health
9	care professional of information that
10	includes—
11	"(I) information (to the extent
12	available and feasible) on the drug or
13	drugs being prescribed for that patient
14	and other information relating to the
15	medical history or condition of the pa-
16	tient that may be relevant to the ap-
17	propriate prescription for that patient;
18	"(II) cost-effective alternatives (if
19	any) for the use of the drug prescribed;
20	and
21	"(III) information on the drugs
22	included in the applicable formulary.
23	To the extent feasible, such program shall
24	permit the prescribing health care profes-
25	sional to provide (and be provided) related



1	information on an interactive, real-time
2	basis.
3	"(B) Standards.—
4	"(i) Development.—The Adminis-
5	trator shall provide for the development of
6	uniform standards relating to the electronic
7	prescription drug program described in sub-
8	paragraph (A). Such standards shall be
9	compatible with standards established under
10	part C of title XI.
11	"(ii) Advisory task force.—In de-
12	veloping such standards and the standards
13	described in subsection $(c)(2)(B)(i)$ the Ad-
14	ministrator shall establish a task force that
15	includes representatives of physicians, hos-
16	pitals, pharmacies, beneficiaries, pharmacy
17	benefit managers, individuals with expertise
18	in information technology, and pharmacy
19	benefit experts of the Departments of Vet-
20	erans Affairs and Defense and other appro-
21	priate Federal agencies to provide rec-
22	ommendations to the Administrator on such
23	standards, including recommendations re-
24	lating to the following:



1	"(I) The range of available com-
2	puterized prescribing software and
3	hardware and their costs to develop
4	and implement.
5	"(II) The extent to which such
6	standards and systems reduce medica-
7	tion errors and can be readily imple-
8	mented by physicians, pharmacies, and
9	hospitals.
10	"(III) Efforts to develop uniform
11	standards and a common software
12	platform for the secure electronic com-
13	munication of medication history, eli-
14	gibility, benefit, and prescription in-
15	formation.
16	"(IV) Efforts to develop and pro-
17	mote universal connectivity and inter-
18	operability for the secure electronic ex-
19	change of such information.
20	"(V) The cost of implementing
21	such systems in the range of hospital
22	and physician office settings and phar-
23	macies, including hardware, software,
24	and training costs.



1	"(VI) Implementation issues as
2	they relate to part C of title XI, and
3	current Federal and State prescribing
4	laws and regulations and their impact
5	on implementation of computerized
6	prescribing.
7	"(iii) Deadlines.—
8	"(I) The Administrator shall con-
9	stitute the task force under clause (ii)
10	by not later than April 1, 2004.
11	"(II) Such task force shall submit
12	recommendations to Administrator by
13	not later than January 1, 2005.
14	"(III) The Administrator shall
15	provide for the development and pro-
16	mulgation, by not later than January
17	1, 2006, of national standards relating
18	to the electronic prescription drug pro-
19	gram described in clause (ii). Such
20	standards shall be issued by a stand-
21	ards organization accredited by the
22	American National Standards Insti-
23	tute (ANSI) and shall be compatible
24	with standards established under part
25	C of title XL



1	"(4) Treatment of accreditation.—Section
2	1852(e)(4) (relating to treatment of accreditation)
3	shall apply to prescription drug plans under this part
4	with respect to the following requirements, in the
5	same manner as they apply to plans under part C
6	with respect to the requirements described in a clause
7	of section $1852(e)(4)(B)$ :
8	"(A) Paragraph (1) (including quality as-
9	surance), including medication therapy manage-
10	ment program under paragraph (2).
11	"(B) Subsection (c)(1) (relating to access to
12	covered benefits).
13	"(C) Subsection (g) (relating to confiden-
14	tiality and accuracy of enrollee records).
15	"(5) Public disclosure of pharmaceutical
16	PRICES FOR EQUIVALENT DRUGS.—Each PDP spon-
17	sor and each entity offering a MA-EFFS Rx plan
18	shall provide that each pharmacy or other dispenser
19	that arranges for the dispensing of a covered out-
20	patient drug shall inform the beneficiary at the time
21	of purchase of the drug of any differential between the
22	price of the prescribed drug to the enrollee and the
23	price of the lowest cost available generic drug covered
24	under the plan that is therapeutically equivalent and
25	bio equivalent.



1	"(e) Grievance Mechanism, Coverage Determina-
2	TIONS, AND RECONSIDERATIONS.—
3	"(1) In general.—Each PDP sponsor shall
4	provide meaningful procedures for hearing and resolv-
5	ing grievances between the organization (including
6	any entity or individual through which the sponsor
7	provides covered benefits) and enrollees with prescrip-
8	tion drug plans of the sponsor under this part in ac-
9	cordance with section 1852(f).
10	"(2) Application of coverage determina-
11	TION AND RECONSIDERATION PROVISIONS.—A PDP
12	sponsor shall meet the requirements of paragraphs (1)
13	through (3) of section 1852(g) with respect to covered
14	benefits under the prescription drug plan it offers
15	under this part in the same manner as such require-
16	ments apply to an organization with respect to bene-
17	fits it offers under a plan under part C.
18	"(3) Request for review of tiered for-
19	MULARY DETERMINATIONS.—In the case of a prescrip-
20	tion drug plan offered by a PDP sponsor or a MA-
21	EFFS Rx plan that provides for tiered cost-sharing
22	for drugs included within a formulary and provides
23	lower cost-sharing for preferred drugs included within
24	the formulary an individual who is enrolled in the

plan may request coverage of a nonpreferred drug



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1	under the terms applicable for preferred drugs if the
2	prescribing physician determines that the preferred
3	drug for treatment of the same condition either would
4	not be as effective for the individual or would have
5	adverse effects for the individual or both.
6	"(f) Appeals.—
7	"(1) In general.—Subject to paragraph (2), a
8	PDP sponsor shall meet the requirements of para-
9	graphs (4) and (5) of section 1852(g) with respect to
10	drugs (including a determination related to the appli-
11	cation of tiered cost-sharing described in subsection
12	(e)(3)) in the same manner as such requirements
13	apply to an organization with respect to benefits it
14	offers under a plan under part C.
15	"(2) Formulary determinations.—An indi-
16	vidual who is enrolled in a prescription drug plan of-
17	fered by a PDP sponsor or in a MA-EFFS Rx plan
18	may appeal to obtain coverage for a covered out-
19	patient drug that is not on a formulary of the sponsor
20	or entity offering the plan if the prescribing physi-

cian determines that the formulary drug for treat-

ment of the same condition either would not be as ef-

fective for the individual or would have adverse effects

for the individual or both.



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1	"(g) Confidentiality and Accuracy of Enrollee
2	Records.—A PDP sponsor that offers a prescription drug
3	plan shall meet the requirements of section 1852(h) with
4	respect to enrollees under the plan in the same manner as
5	such requirements apply to an organization with respect
6	to enrollees under part C. A PDP sponsor shall be treated
7	as a business associate for purposes of the provisions of sub-
8	part E of part 164 of title 45, Code of Federal Regulations,
9	adopted pursuant to the authority of the Secretary under
10	section 264(c) of the Health Insurance Portability and Ac-
11	countability Act of 1996 (42 U.S. C. 1320d-2 note).
12	"SEC. 1860D-4. REQUIREMENTS FOR AND CONTRACTS WITH
13	PRESCRIPTION DRUG PLAN (PDP) SPONSORS.
14	"(a) General Requirements.—Each PDP sponsor
15	of a prescription drug plan shall meet the following require-
16	ments:
17	"(1) Licensure.—Subject to subsection (c), the
18	sponsor is organized and licensed under State law as
19	a risk-bearing entity eligible to offer health insurance
20	or health benefits coverage in each State in which it
21	offers a prescription drug plan.
22	"(2) Assumption of financial risk for un-
23	SUBSIDIZED COVERAGE.—
24	"(A) In general.—Subject to subpara-
25	araph(B) and section $1860D-5(d)(2)$ , the entity



1	assumes full financial risk on a prospective basis
2	for qualified prescription drug coverage that it
3	offers under a prescription drug plan and that
4	is not covered under section 1860D-8.
5	"(B) Reinsurance permitted.—The enti-
6	ty may obtain insurance or make other arrange-
7	ments for the cost of coverage provided to any
8	enrollee.
9	"(3) Solvency for unlicensed sponsors.—In
10	the case of a sponsor that is not described in para-
11	graph (1), the sponsor shall meet solvency standards
12	established by the Administrator under subsection (d).
13	"(b) Contract Requirements.—
14	"(1) In general.—The Administrator shall not
15	permit the election under section 1860D-1 of a pre-
16	scription drug plan offered by a PDP sponsor under
17	this part, and the sponsor shall not be eligible for
18	payments under section 1860D-7 or 1860D-8, unless
19	the Administrator has entered into a contract under
20	this subsection with the sponsor with respect to the of-
21	fering of such plan. Such a contract with a sponsor
22	may cover more than one prescription drug plan.
23	Such contract shall provide that the sponsor agrees to

comply with the applicable requirements and stand-



1	ards of this part and the terms and conditions of
2	payment as provided for in this part.
3	"(2) Negotiation regarding terms and con-
4	DITIONS.—The Administrator shall have the same au-
5	thority to negotiate the terms and conditions of pre-
6	scription drug plans under this part as the Director
7	of the Office of Personnel Management has with re-
8	spect to health benefits plans under chapter 89 of title
9	5, United States Code. In negotiating the terms and
10	conditions regarding premiums for which information
11	is submitted under section 1860 $D$ -6(a)(2), the Ad-
12	ministrator shall take into account the subsidy pay-
13	$ments\ under\ section\ 1860D\!-\!8.$
14	"(3) Incorporation of certain medicare ad-
15	VANTAGE CONTRACT REQUIREMENTS.—The following
16	provisions of section 1857 shall apply, subject to sub-
17	section (c)(5), to contracts under this section in the
18	same manner as they apply to contracts under section
19	1857(a):
20	"(A) Minimum enrollment.—Paragraphs
21	(1) and (3) of section 1857(b).
22	"(B) Contract period and effective-
23	NESS.—Paragraphs (1) through (3) and (5) of
24	section $1857(c)$ .



1	"(C) Protections against fraud and
2	Beneficiary protections.—Section $1857(d)$ .
3	"(D) Additional contract terms.—Sec-
4	tion 1857(e); except that in applying section
5	1857(e)(2) under this part—
6	"(i) such section shall be applied sepa-
7	rately to costs relating to this part (from
8	costs under part C and part E);
9	"(ii) in no case shall the amount of the
10	fee established under this subparagraph for
11	a plan exceed 20 percent of the maximum
12	amount of the fee that may be established
13	under subparagraph (B) of such section;
14	and
15	"(iii) no fees shall be applied under
16	this subparagraph with respect to MA-
17	EFFS Rx plans.
18	"(E) Intermediate sanctions.—Section
19	1857(g).
20	"(F) Procedures for termination.—
21	Section 1857(h).
22	"(4) Rules of application for intermediate
23	SANCTIONS.—In applying paragraph (3)(E)—



1	"(A) the reference in section $1857(g)(1)(B)$
2	to section 1854 is deemed a reference to this
3	part; and
4	"(B) the reference in section $1857(g)(1)(F)$
5	to section $1852(k)(2)(A)(ii)$ shall not be applied.
6	"(5) Service area requirement.—For pur-
7	poses of this part, the Administrator shall designate
8	at least 10 areas covering the entire United States
9	and shall be consistent with EFFS regions established
10	under section $1860E-1(a)(2)$ .
11	"(c) Waiver of Certain Requirements to Expand
12	Choice.—
13	"(1) In general.—In the case of an entity that
14	seeks to offer a prescription drug plan in a State, the
15	Administrator shall waive the requirement of sub-
16	section (a)(1) that the entity be licensed in that State
17	if the Administrator determines, based on the applica-
18	tion and other evidence presented to the Adminis-
19	trator, that any of the grounds for approval of the ap-
20	plication described in paragraph (2) have been met.
21	"(2) Grounds for Approval.—The grounds for
22	approval under this paragraph are the grounds for
23	approval described in subparagraph (B), (C), and
24	(D) of section $1855(a)(2)$ , and also include the appli-



1	cation by a State of any grounds other than those re-
2	quired under Federal law.
3	"(3) Application of waiver procedures.—
4	With respect to an application for a waiver (or a
5	waiver granted) under this subsection, the provisions
6	of subparagraphs $(E)$ , $(F)$ , and $(G)$ of section
7	1855(a)(2) shall apply.
8	"(4) Licensure does not substitute for or
9	CONSTITUTE CERTIFICATION.—The fact that an entity
10	is licensed in accordance with subsection (a)(1) does
11	not deem the entity to meet other requirements im-
12	posed under this part for a PDP sponsor.
13	"(5) References to certain provisions.—
14	For purposes of this subsection, in applying provi-
15	sions of section 1855(a)(2) under this subsection to
16	prescription drug plans and PDP sponsors—
17	"(A) any reference to a waiver application
18	under section 1855 shall be treated as a reference
19	to a waiver application under paragraph (1);
20	and
21	"(B) any reference to solvency standards
22	shall be treated as a reference to solvency stand-
23	$ards\ established\ under\ subsection\ (d).$
24	"(d) Solvency Standards for Non-Licensed
25	Sponsors.—



1	"(1) Establishment.—The Administrator shall
2	establish, by not later than October 1, 2004, financial
3	solvency and capital adequacy standards that an en-
4	tity that does not meet the requirements of subsection
5	(a)(1) must meet to qualify as a PDP sponsor under
6	this part.
7	"(2) Compliance with standards.—Each
8	PDP sponsor that is not licensed by a State under
9	subsection (a)(1) and for which a waiver application
10	has been approved under subsection (c) shall meet sol-
11	vency and capital adequacy standards established
12	under paragraph (1). The Administrator shall estab-
13	lish certification procedures for such PDP sponsors
14	with respect to such solvency standards in the manner
15	described in section $1855(c)(2)$ .
16	"(e) Relation to State Laws.—
17	"(1) In General.—The standards established
18	under this part shall supersede any State law or reg-
19	ulation (other than State licensing laws or State laws
20	relating to plan solvency, except as provided in sub-
21	section (d)) with respect to prescription drug plans
22	which are offered by PDP sponsors under this part.
23	"(2) Prohibition of State imposition of
24	PREMIUM TAXES.—No State may impose a premium

tax or similar tax with respect to premiums paid to



1	PDP sponsors for prescription drug plans under this
2	part, or with respect to any payments made to such
3	a sponsor by the Administrator under this part.
4	"SEC. 1860D-5. PROCESS FOR BENEFICIARIES TO SELECT
5	QUALIFIED PRESCRIPTION DRUG COVERAGE.
6	$``(a)\ In\ General.$ —The Administrator shall establish
7	a process for the selection of the prescription drug plan or
8	MA-EFFS Rx plan through which eligible individuals elect
9	qualified prescription drug coverage under this part.
10	"(b) Elements.—Such process shall include the fol-
11	lowing:
12	"(1) Annual, coordinated election periods, in
13	which such individuals can change the qualifying
14	plans through which they obtain coverage, in accord-
15	ance with section $1860D-1(b)(2)$ .
16	"(2) Active dissemination of information to pro-
17	mote an informed selection among qualifying plans
18	based upon price, quality, and other features, in the
19	manner described in (and in coordination with) sec-
20	tion 1851(d), including the provision of annual com-
21	parative information, maintenance of a toll-free hot-
22	line, and the use of non-Federal entities.
23	"(3) Coordination of elections through filing
24	with the entity offering a MA-EFFS Rx plan or a



1	PDP sponsor, in the manner described in (and in co-
2	ordination with) section $1851(c)(2)$ .
3	"(4) Informing each enrollee before the beginning
4	of each year of the annual out-of-pocket threshold ap-
5	plicable to the enrollee for that year under section
6	1860D-2(b)(4) at such time.
7	"(c) MA-EFFS RX ENROLLEE MAY ONLY OBTAIN
8	Benefits Through the Plan.—An individual who is
9	enrolled under a MA-EFFS Rx plan may only elect to re-
10	ceive qualified prescription drug coverage under this part
11	through such plan.
12	"(d) Assuring Access to a Choice of Qualified
13	Prescription Drug Coverage.—
14	"(1) Choice of at least two plans in each
15	AREA.—
16	``(A) IN GENERAL.—The Administrator
17	shall assure that each individual who is entitled
18	to benefits under part A or enrolled under part
19	B and who is residing in an area in the United
20	States has available, consistent with subpara-
21	graph (B), a choice of enrollment in at least two
22	qualifying plans (as defined in paragraph (5))
23	in the area in which the individual resides, at
24	least one of which is a prescription drug plan.



1	"(B) REQUIREMENT FOR DIFFERENT PLAN
2	SPONSORS.—The requirement in subparagraph
3	(A) is not satisfied with respect to an area is
4	only one PDP sponsor or one entity that offers
5	a MA-EFFS Rx plan offers all the qualifying
6	plans in the area.
7	"(2) Guaranteeing access to coverage.—In
8	order to assure access under paragraph (1) and con-
9	sistent with paragraph (3), the Administrator may
10	provide partial underwriting of risk for a PDP spon-
11	sor to expand the service area under an existing pre-
12	scription drug plan to adjoining or additional areas
13	or to establish such a plan (including offering such a
14	plan on a regional or nationwide basis), but only so
15	long as (and to the extent) necessary to assure the ac-
16	cess guaranteed under paragraph (1).
17	"(3) Limitation on authority.—In exercising
18	authority under this subsection, the Administrator—
19	"(A) shall not provide for the full under-
20	writing of financial risk for any PDP sponsor,
21	and
22	"(B) shall seek to maximize the assumption
23	of financial risk by PDP sponsors or entities of
24	fering a MA-EFFS Rx plan.



1	"(4) Reports.—The Administrator shall, in
2	each annual report to Congress under section 1809(f),
3	include information on the exercise of authority under
4	this subsection. The Administrator also shall include
5	such recommendations as may be appropriate to min-
6	imize the exercise of such authority, including mini-
7	mizing the assumption of financial risk.
8	"(5) Qualifying plan defined.—For purposes
9	of this subsection, the term 'qualifying plan' means a
10	prescription drug plan or a MA-EFFS Rx plan.
11	"SEC. 1860D-6. SUBMISSION OF BIDS AND PREMIUMS.
12	"(a) Submission of Bids, Premiums, and Related
13	Information.—
14	"(1) In general.—Each PDP sponsor shall
15	submit to the Administrator the information de-
16	scribed in paragraph (2) in the same manner as in-
17	formation is submitted by an organization under sec-
18	$tion \ 1854(a)(1).$
19	"(2) Information submitted.—The informa-
20	tion described in this paragraph is the following:
21	"(A) Coverage provided.—Information
22	on the qualified prescription drug coverage to be
23	provided.
24	"(B) Actuarial value.—Information on
25	the actuarial value of the coverage.



1	"(C) BID AND PREMIUM.—Information on
2	the bid and the premium for the coverage, in-
3	cluding an actuarial certification of—
4	"(i) the actuarial basis for such bid
5	and premium;
6	"(ii) the portion of such bid and pre-
7	mium attributable to benefits in excess of
8	standard coverage;
9	"(iii) the reduction in such bid result-
10	ing from the reinsurance subsidy payments
11	provided under section 1860D-8(a)(2); and
12	"(iv) the reduction in such premium
13	resulting from the direct and reinsurance
14	subsidy payments provided under section
15	1860D–8.
16	"(D) Additional information.—Such
17	other information as the Administrator may re-
18	quire to carry out this part.
19	"(3) Review of information; negotiation
20	AND APPROVAL OF PREMIUMS.—
21	"(A) In general.—Subject to subpara-
22	graph (B), the Administrator shall review the in-
23	formation filed under paragraph (2) for the pur-
24	pose of conducting negotiations under section
25	1860D-4(b)(2) (relating to using OPM-like au-



1	thority under the FEHBP). The Administrator,
2	using the information provided (including the
3	actuarial certification under paragraph (2)(C))
4	shall approve the premium submitted under this
5	subsection only if the premium accurately re-
6	flects both (i) the actuarial value of the benefits
7	provided, and (ii) the 73 percent average subsidy
8	provided under section 1860D-8 for the standard
9	benefit. The Administrator shall apply actuarial
10	principles to approval of a premium under this
11	part in a manner similar to the manner in
12	which those principles are applied in estab-
13	lishing the monthly part B premium under sec-
14	tion 1839.
15	"(B) Exception.—In the case of a plan de-
16	scribed in section 1851(a)(2)(C), the provisions
17	of subparagraph (A) shall not apply and the
18	provisions of paragraph (5)(B) of section
19	1854(a), prohibiting the review, approval, or dis-
20	approval of amounts described in such para-
21	graph, shall apply to the negotiation and rejec-
22	tion of the monthly bid amounts and proportion
23	referred to in submargaranh $(A)$



	, -
1	"(1) In general.—The bid and premium for a
2	prescription drug plan under this section may not
3	vary among enrollees in the plan in the same service
4	area.
5	"(2) Construction.—Nothing in paragraph (1)
6	shall be construed as preventing the imposition of a
7	late enrollment penalty under section 1860D-
8	1(c)(2)(B).
9	"(c) Collection.—
10	"(1) Beneficiary's option of payment
11	THROUGH WITHHOLDING FROM SOCIAL SECURITY
12	PAYMENT OR USE OF ELECTRONIC FUNDS TRANSFER
13	MECHANISM.—In accordance with regulations, a PDP
14	sponsor shall permit each enrollee, at the enrollee's
15	option, to make payment of premiums under this
16	part to the sponsor through withholding from benefit
17	payments in the manner provided under section 1840
18	with respect to monthly premiums under section 1839
19	or through an electronic funds transfer mechanism
20	(such as automatic charges of an account at a finan-
21	cial institution or a credit or debit card account) or
22	otherwise. All premium payments that are withheld
23	under this paragraph shall be credited to the Medi-
24	care Prescription Drug Trust Fund and shall be paid

to the PDP sponsor involved.



1	"(2) Offsetting.—Reductions in premiums for
2	coverage under parts A and B as a result of a selec-
3	tion of a MA-EFFS Rx plan may be used to reduce
4	the premium otherwise imposed under paragraph (1).
5	"(d) Acceptance of Reference Premium Amount
6	AS FULL PREMIUM FOR SUBSIDIZED LOW-INCOME INDI-
7	VIDUALS IF NO STANDARD (OR EQUIVALENT) COVERAGE IN
8	AN AREA.—
9	"(1) In General.—If there is no standard pre-
10	scription drug coverage (as defined in paragraph (2))
11	offered in an area, in the case of an individual who
12	is eligible for a premium subsidy under section
13	1860D-7 and resides in the area, the PDP sponsor of
14	any prescription drug plan offered in the area (and
15	any entity offering a MA-EFFS Rx plan in the area)
16	shall accept the reference premium amount (under
17	paragraph (3)) as payment in full for the premium
18	charge for qualified prescription drug coverage.
19	"(2) Standard prescription drug coverage
20	DEFINED.—For purposes of this subsection, the term
21	'standard prescription drug coverage' means qualified
22	prescription drug coverage that is standard coverage
23	or that has an actuarial value equivalent to the actu-

 $arial\ value\ for\ standard\ coverage.$ 



1	"(3) Reference premium amount defined.—
2	For purposes of this subsection, the term 'reference
3	premium amount' means, with respect to qualified
4	prescription drug coverage offered under—
5	"(A) a prescription drug plan that—
6	"(i) provides standard coverage (or al-
7	ternative prescription drug coverage the ac-
8	tuarial value is equivalent to that of stand-
9	ard coverage), the plan's PDP premium; or
10	"(ii) provides alternative prescription
11	drug coverage the actuarial value of which
12	is greater than that of standard coverage,
13	the plan's PDP premium multiplied by the
14	ratio of (I) the actuarial value of standard
15	coverage, to (II) the actuarial value of the
16	$alternative\ coverage;$
17	"(B) an EFFS plan, the EFFS monthly
18	prescription drug beneficiary premium (as de-
19	fined in section $1860E-4(a)(3)(B)$ ; or
20	"(C) a Medicare Advantage, the Medicare
21	Advantage monthly prescription drug beneficiary
22	premium (as defined in section $1854(b)(2)(B)$ ).
23	For purposes of subparagraph (A), the term 'PDP
24	premium' means, with respect to a prescription drug
25	plan, the premium amount for enrollment under the



1	plan under this part (determined without regard to
2	any low-income subsidy under section 1860D-7 or
3	any late enrollment penalty under section 1860D-
4	1(c)(2)(B)).
5	"SEC. 1860D-7. PREMIUM AND COST-SHARING SUBSIDIES
6	FOR LOW-INCOME INDIVIDUALS.
7	"(a) Income-Related Subsidies for Individuals
8	WITH INCOME BELOW 150 PERCENT OF FEDERAL POVERTY
9	Level.—
10	"(1) Full premium subsidy and reduction
11	OF COST-SHARING FOR INDIVIDUALS WITH INCOME
12	BELOW 135 PERCENT OF FEDERAL POVERTY LEVEL.—
13	In the case of a subsidy eligible individual (as defined
14	in paragraph (4)) who is determined to have income
15	that does not exceed 135 percent of the Federal pov-
16	erty level, the individual is entitled under this
17	section—
18	"(A) to an income-related premium subsidy
19	equal to 100 percent of the amount described in
20	$subsection (b)(1); \ and$
21	"(B) subject to subsection (c), to the substi-
22	tution for the beneficiary cost-sharing described
23	in paragraphs (1) and (2) of section 1860D-2(b)
24	(up to the initial coverage limit specified in
25	paragraph (3) of such section) of amounts that



1	do not exceed \$2 for a multiple source or generic
2	drug (as described in section $1927(k)(7)(A)$ ) and
3	\$5 for a non-preferred drug.
4	"(2) Sliding scale premium subsidy for in-
5	DIVIDUALS WITH INCOME ABOVE 135, BUT BELOW 150
6	PERCENT, OF FEDERAL POVERTY LEVEL.—In the case
7	of a subsidy eligible individual who is determined to
8	have income that exceeds 135 percent, but does not ex-
9	ceed 150 percent, of the Federal poverty level, the in-
10	dividual is entitled under this section to an income-
11	related premium subsidy determined on a linear slid-
12	ing scale ranging from 100 percent of the amount de-
13	scribed in subsection (b)(1) for individuals with in-
14	comes at 135 percent of such level to 0 percent of such
15	amount for individuals with incomes at 150 percent
16	of such level.
17	"(3) Construction.—Nothing in this section
18	shall be construed as preventing a PDP sponsor or
19	entity offering a MA-EFFS Rx plan from reducing to
20	0 the cost-sharing otherwise applicable to generic
21	drugs.
22	"(4) Determination of eligibility.—
23	"(A) Subsidy eligible individual de-
24	FINED.—For purposes of this section, subject to



1	subparagraph $(D)$ , the term 'subsidy eligible in-
2	dividual' means an individual who—
3	"(i) is eligible to elect, and has elected,
4	to obtain qualified prescription drug cov-
5	erage under this part;
6	"(ii) has income below 150 percent of
7	the Federal poverty line; and
8	"(iii) meets the resources requirement
9	$described\ in\ subparagraph\ (D)\ .$
10	"(B) Determinations.—The determina-
11	tion of whether an individual residing in a State
12	is a subsidy eligible individual and the amount
13	of such individual's income shall be determined
14	under the State medicaid plan for the State
15	under section 1935(a) or by the Social Security
16	Administration. In the case of a State that does
17	not operate such a medicaid plan (either under
18	title XIX or under a statewide waiver granted
19	under section 1115), such determination shall be
20	made under arrangements made by the Adminis-
21	trator. There are authorized to be appropriated
22	to the Social Security Administration such sums
23	as may be necessary for the determination of eli-
24	gibility under this subparagraph.



1	"(C) Income determinations.—For pur-
2	poses of applying this section—
3	"(i) income shall be determined in the
4	$manner\ described\ in\ section\ 1905(p)(1)(B);$
5	and
6	"(ii) the term 'Federal poverty line'
7	means the official poverty line (as defined
8	by the Office of Management and Budget,
9	and revised annually in accordance with
10	section 673(2) of the Omnibus Budget Rec-
11	onciliation Act of 1981) applicable to a
12	family of the size involved.
13	"(D) Resource standard applied to be
14	BASED ON TWICE SSI RESOURCE STANDARD.—
15	The resource requirement of this subparagraph is
16	that an individual's resources (as determined
17	under section 1613 for purposes of the supple-
18	mental security income program) do not
19	exceed—
20	"(i) for 2006 twice the maximum
21	amount of resources that an individual may
22	have and obtain benefits under that pro-
23	gram; and
24	"(ii) for a subsequent year the resource
25	limitation established under this clause for



1	the previous year increased by the annual
2	percentage increase in the consumer price
3	index (all items; U.S. city average) as of
4	September of such previous year.
5	Any resource limitation established under clause
6	(ii) that is not a multiple of \$10 shall be round-
7	ed to the nearest multiple of \$10.
8	"(E) Treatment of territorial resi-
9	DENTS.—In the case of an individual who is not
10	a resident of the 50 States or the District of Co-
11	lumbia, the individual is not eligible to be a sub-
12	sidy eligible individual but may be eligible for
13	financial assistance with prescription drug ex-
14	penses under section 1935(e).
15	"(F) Treatment of conforming medigap
16	POLICIES.—For purposes of this section, the term
17	'qualified prescription drug coverage' includes a
18	medicare supplemental policy described in sec-
19	tion 1860D-8(b)(4).
20	"(5) Indexing dollar amounts.—
21	"(A) FOR 2007.—The dollar amounts ap-
22	plied under paragraphs (1)(B) for 2007 shall be
23	the dollar amounts specified in such paragraph
24	increased by the annual percentage increase de-

scribed in section 1860D-2(b)(5) for 2007.



1	"(B) For subsequent years.—The dollar
2	amounts applied under paragraph (1)(B) for a
3	year after 2007 shall be the amounts (under this
4	paragraph) applied under paragraph (1)(B) for
5	the preceding year increased by the annual per-
6	centage increase described in section 1860D-
7	2(b)(5) (relating to growth in medicare prescrip-
8	tion drug costs per beneficiary) for the year in-
9	volved.
10	"(b) Premium Subsidy Amount.—
11	"(1) In General.—The premium subsidy
12	amount described in this subsection for an individual
13	residing in an area is the benchmark premium
14	amount (as defined in paragraph (2)) for qualified
15	prescription drug coverage offered by the prescription
16	drug plan or the MA-EFFS Rx plan in which the in-
17	dividual is enrolled.
18	"(2) Benchmark premium amount defined.—
19	For purposes of this subsection, the term benchmark
20	premium amount' means, with respect to qualified
21	prescription drug coverage offered under—
22	"(A) a prescription drug plan that—
23	"(i) provides standard coverage (or al-
24	ternative prescription drug coverage the ac-
25	tuarial value of which is equivalent to that



1	of standard coverage), the premium amount
2	for enrollment under the plan under this
3	part (determined without regard to any
4	subsidy under this section or any late en-
5	rollment penalty under section 1860D-
6	1(c)(2)(B)); or
7	"(ii) provides alternative prescription
8	drug coverage the actuarial value of which
9	is greater than that of standard coverage,
10	the premium amount described in clause (i)
11	multiplied by the ratio of (I) the actuarial
12	value of standard coverage, to (II) the actu-
13	arial value of the alternative coverage; or
14	"(B) a MA-EFFS Rx plan, the portion of
15	the premium amount that is attributable to stat-
16	utory drug benefits (described in section
17	1853(a)(1)(A)(ii)(II)).
18	"(c) Rules in Applying Cost-Sharing Sub-
19	SIDIES.—
20	"(1) In General.—In applying subsection
21	(a)(1)(B), nothing in this part shall be construed as
22	preventing a plan or provider from waiving or reduc-
23	ing the amount of cost-sharing otherwise applicable.
24	"(2) Limitation on charges.—In the case of
25	an individual receiving cost-sharing subsidies under



1	subsection $(a)(1)(B)$ , the PDP sponsor or entity offer-
2	ing a MA-EFFS Rx plan may not charge more than
3	\$5 per prescription.
4	"(3) Application of indexing rules.—The
5	provisions of subsection (a)(5) shall apply to the dol-
6	lar amount specified in paragraph (2) in the same
7	manner as they apply to the dollar amounts specified
8	in subsections $(a)(1)(B)$ .
9	"(d) Administration of Subsidy Program.—The
10	Administrator shall provide a process whereby, in the case
11	of an individual who is determined to be a subsidy eligible
12	individual and who is enrolled in prescription drug plan
13	or is enrolled in a MA-EFFS Rx plan—
14	"(1) the Administrator provides for a notifica-
15	tion of the PDP sponsor or the entity offering the
16	MA-EFFS Rx plan involved that the individual is el-
17	igible for a subsidy and the amount of the subsidy
18	under subsection (a);
19	"(2) the sponsor or entity involved reduces the
20	premiums or cost-sharing otherwise imposed by the
21	amount of the applicable subsidy and submits to the
22	Administrator information on the amount of such re-
23	duction: and



1	"(3) the Administrator periodically and on a
2	timely basis reimburses the sponsor or entity for the
3	amount of such reductions.
4	The reimbursement under paragraph (3) with respect to
5	cost-sharing subsidies may be computed on a capitated
6	basis, taking into account the actuarial value of the sub-
7	sidies and with appropriate adjustments to reflect dif-
8	ferences in the risks actually involved.
9	"(e) Relation to Medicaid Program.—
10	"(1) In general.—For provisions providing for
11	eligibility determinations, and additional financing,
12	under the medicaid program, see section 1935.
13	"(2) Medicaid providing wrap around bene-
14	FITS.—The coverage provided under this part is pri-
15	mary payor to benefits for prescribed drugs provided
16	under the medicaid program under title XIX con-
17	sistent with section $1935(d)(1)$ .
18	"(3) Coordination.—The Administrator shall
19	develop and implement a plan for the coordination of
20	prescription drug benefits under this part with the
21	benefits provided under the medicaid program under
22	title XIX, with particular attention to insuring co-
23	ordination of payments and prevention of fraud and
24	abuse. In developing and implementing such plan, the
25	Administrator shall involve the Secretary, the States,



1	the data processing industry, pharmacists, and phar-
2	maceutical manufacturers, and other experts.
3	"SEC. 1860D-8. SUBSIDIES FOR ALL MEDICARE BENE-
4	FICIARIES FOR QUALIFIED PRESCRIPTION
5	DRUG COVERAGE.
6	"(a) Subsidy Payment.—In order to reduce premium
7	levels applicable to qualified prescription drug coverage for
8	all medicare beneficiaries consistent with an overall subsidy
9	level of 73 percent, to reduce adverse selection among pre-
10	scription drug plans and MA-EFFS Rx plans, and to pro-
11	mote the participation of PDP sponsors under this part,
12	the Administrator shall provide in accordance with this sec-
13	tion for payment to a qualifying entity (as defined in sub-
14	section (b)) of the following subsidies:
15	"(1) Direct subsidy.—In the case of an en-
16	rollee enrolled for a month in a prescription drug
17	plan or a MA-EFFS Rx plan, a direct subsidy equal
18	to 43 percent of the national average monthly bid
19	amount (computed under subsection (g)) for that
20	month.
21	"(2) Subsidy through reinsurance.—In the
22	case of an enrollee enrolled for a month in a prescrip-
23	tion drug plan or a MA-EFFS Rx plan, the reinsur-
24	ance payment amount (as defined in subsection (c)),
25	which in the agaregate is 30 percent of the total pay-



1	ments made by qualifying entities for standard cov-
2	erage under the respective plan, for excess costs in-
3	curred in providing qualified prescription drug
4	coverage—
5	"(A) for enrollees with a prescription drug
6	plan under this part; and
7	"(B) for enrollees with a MA-EFFS Rx
8	plan.
9	"(3) Employer and union flexibility.—In
10	the case of an individual who is a participant or ben-
11	eficiary in a qualified retiree prescription drug plan
12	(as defined in subsection $(f)(1)$ ) and who is not en-
13	rolled in a prescription drug plan or in a MA-EFFS
14	Rx plan, the special subsidy payments under sub-
15	section $(f)(3)$ .
16	This section constitutes budget authority in advance of ap-
17	propriations Acts and represents the obligation of the Ad-
18	ministrator to provide for the payment of amounts provided
19	under this section. In applying the percentages under para-
20	graphs (1) and (2), there shall be taken into account under
21	the respective paragraphs the portion of the employer and
22	union special subsidy payments under subsection (f)(3) that
23	reflect payments that would have been made under the re-
24	spective paragraphs if such paragraphs had applied to



1	qualified retiree prescription drug plans instead of para-
2	graph (3).
3	"(b) Qualifying Entity Defined.—For purposes of
4	this section, the term 'qualifying entity' means any of the
5	following that has entered into an agreement with the Ad-
6	ministrator to provide the Administrator with such infor-
7	mation as may be required to carry out this section:
8	"(1) A PDP sponsor offering a prescription drug
9	plan under this part.
10	"(2) An entity that offers a MA-EFFS Rx plan.
11	"(3) The sponsor of a qualified retiree prescrip-
12	tion drug plan (as defined in subsection (f)).
13	"(c) Reinsurance Payment Amount.—
14	"(1) In General.—Subject to subsection
15	(d)(1)(B) and paragraph (4), the reinsurance pay-
16	ment amount under this subsection for a qualifying
17	covered individual (as defined in paragraph (5)) for
18	a coverage year (as defined in subsection $(h)(2)$ ) is
19	equal to the sum of the following:
20	"(A) Reinsurance between initial rein-
21	SURANCE THRESHOLD AND THE INITIAL COV-
22	ERAGE LIMIT.—For the portion of the individ-
23	ual's gross covered prescription drug costs (as de-
24	fined in paragraph (3)) for the year that exceeds
25	the initial reinsurance threshold specified in



1	paragraph (4), but does not exceed the initial
2	coverage limit specified in section 1860D-
3	2(b)(3), an amount equal to 20 percent of the al-
4	lowable costs (as defined in paragraph (2)) at-
5	tributable to such gross covered prescription drug
6	costs.
7	"(B) Reinsurance above annual out-of-
8	POCKET THRESHOLD.—For the portion of the in-
9	dividual's gross covered prescription drug costs
10	for the year that exceeds the annual out-of-pocket
11	threshold specified in $1860D-2(b)(4)(B)$ , an
12	amount equal to 80 percent of the allowable costs
13	attributable to such gross covered prescription
14	$drug\ costs.$
15	"(2) Allowable costs.—For purposes of this
16	section, the term 'allowable costs' means, with respect
17	to gross covered prescription drug costs under a plan
18	described in subsection (b) offered by a qualifying en-
19	tity, the part of such costs that are actually paid (net
20	of discounts, chargebacks, and average percentage re-
21	bates) under the plan, but in no case more than the
22	part of such costs that would have been paid under
23	the plan if the prescription drug coverage under the

plan were standard coverage.



1	"(3) Gross covered prescription drug
2	costs.—For purposes of this section, the term 'gross
3	covered prescription drug costs' means, with respect to
4	an enrollee with a qualifying entity under a plan de-
5	scribed in subsection (b) during a coverage year, the
6	costs incurred under the plan (including costs attrib-
7	utable to administrative costs) for covered prescrip-
8	tion drugs dispensed during the year, including costs
9	relating to the deductible, whether paid by the enrollee
10	or under the plan, regardless of whether the coverage
11	under the plan exceeds standard coverage and regard-
12	less of when the payment for such drugs is made.
13	"(4) Initial reinsurance threshold.—The
14	initial reinsurance threshold specified in this
15	paragraph—
16	"(A) for 2006, is equal to \$1,000; or
17	"(B) for a subsequent year, is equal to the
18	payment threshold specified in this paragraph
19	for the previous year, increased by the annual
20	percentage increase described in section 1860D-
21	2(b)(5) for the year involved.
22	Any amount determined under subparagraph (B)
23	that is not a multiple of \$10 shall be rounded to the
24	nearest multiple of \$10.



1	"(5) Qualifying covered individual de-
2	FINED.—For purposes of this subsection, the term
3	'qualifying covered individual' means an individual
4	who—
5	"(A) is enrolled with a prescription drug
6	plan under this part; or
7	"(B) is enrolled with a MA-EFFS Rx plan.
8	"(d) Adjustment of Payments.—
9	"(1) Adjustment of Reinsurance Payments
10	TO ASSURE 30 PERCENT LEVEL OF SUBSIDY THROUGH
11	REINSURANCE.—
12	"(A) Estimation of payments.—The Ad-
13	ministrator shall estimate—
14	"(i) the total payments to be made
15	(without regard to this subsection) during a
16	year under subsections (a)(2) and (c); and
17	"(ii) the total payments to be made by
18	qualifying entities for standard coverage
19	under plans described in subsection (b) dur-
20	ing the year.
21	"(B) Adjustment.—The Administrator
22	shall proportionally adjust the payments made
23	under subsections (a)(2) and (c) for a coverage
24	year in such manner so that the total of the pay-
25	ments made under such subsections (and under



1	subsection (f)(3) insofar as such payments reflect
2	payments that would have been made under such
3	subsections if such subsections had applied to
4	qualified retiree prescription drug plans instead
5	of subsections $(a)(3)$ and $(f)(3)$ ) for the year is
6	equal to 30 percent of the total payments de-
7	$scribed\ in\ subparagraph\ (A)(ii).$
8	"(2) Risk adjustment for direct sub-
9	SIDIES.—To the extent the Administrator determines
10	it appropriate to avoid risk selection, the payments
11	made for direct subsidies under subsection (a)(1) are
12	subject to adjustment based upon risk factors specified
13	by the Administrator. Any such risk adjustment shall
14	be designed in a manner as to not result in a change
15	in the aggregate payments made under such sub-
16	section.
17	"(e) Payment Methods.—
18	"(1) In general.—Payments under this section
19	shall be based on such a method as the Administrator
20	determines. The Administrator may establish a pay-
21	ment method by which interim payments of amounts
22	under this section are made during a year based on
23	the Administrator's best estimate of amounts that will

be payable after obtaining all of the information.



1	"(2) Source of payments.—Payments under
2	this section shall be made from the Medicare Prescrip-
3	tion Drug Trust Fund.
4	"(f) Rules Relating to Qualified Retiree Pre-
5	SCRIPTION DRUG PLAN.—
6	"(1) Definition.—For purposes of this section,
7	the term 'qualified retiree prescription drug plan'
8	means employment-based retiree health coverage (as
9	defined in paragraph (4)(A)) if, with respect to an
10	individual who is a participant or beneficiary under
11	such coverage and is eligible to be enrolled in a pre-
12	scription drug plan or a MA-EFFS Rx plan under
13	this part, the following requirements are met:
14	"(A) ACTUARIAL EQUIVALENCE TO STAND-
15	ARD COVERAGE.—The Administrator determines
16	(based on an actuarial analysis by the Adminis-
17	trator) that coverage provides at least the same
18	actuarial value as standard coverage. Such de-
19	termination may be made on an annual basis.
20	"(B) AUDITS.—The sponsor (and the plan)
21	shall maintain, and afford the Administrator ac-
22	cess to, such records as the Administrator may
23	require for purposes of audits and other oversight
24	activities necessary to ensure the adequacy of



1	prescription drug coverage and the accuracy of
2	payments made.
3	"(C) Provision of Certification of Pre-
4	SCRIPTION DRUG COVERAGE.—The sponsor of the
5	plan shall provide for issuance of certifications
6	of the type described in section 1860D-
7	1(c)(2)(D).
8	"(2) Limitation on benefit eligibility.—No
9	payment shall be provided under this section with re-
10	spect to a participant or beneficiary in a qualified re-
11	tiree prescription drug plan unless the individual
12	is—
13	"(A) is covered under the plan; and
14	"(B) is eligible to obtain qualified prescrip-
15	tion drug coverage under section 1860D-1 but
16	did not elect such coverage under this part (ei-
17	ther through a prescription drug plan or through
18	a MA-EFFS $Rx$ $plan)$ .
19	"(3) Employer and union special subsidy
20	AMOUNTS.—
21	"(A) In general.—For purposes of sub-
22	section (a), the special subsidy payment amount
23	under this paragraph for a qualifying covered
24	retiree(as defined in paragraph (6)) for a cov-
25	erage year (as defined in subsection (h)) enrolled



1	in a qualifying entity described in subsection
2	(b)(3) under a qualified retiree prescription drug
3	plan is, for the portion of the individual's gross
4	covered prescription drug costs for the year that
5	exceeds the deductible amount specified in sub-
6	paragraph (B), an amount equal to, subject to
7	subparagraph (D), 28 percent of the allowable
8	costs attributable to such gross covered prescrip-
9	tion drug costs, but only to the extent such costs
10	exceed the deductible under subparagraph (B)
11	and do not exceed the cost limit under such sub-
12	paragraph in the case of any such individual for
13	the plan year.
14	"(B) Deductible and cost limit appli-
15	CABLE.—Subject to subparagraph (C)—
16	"(i) the deductible under this subpara-
17	graph is equal to \$250 for plan years that
18	end in 2006; and
19	"(ii) the cost limit under this subpara-
20	graph is equal to \$5,000 for plan years that
21	end in 2006.
22	"(C) Indexing.—The deductible and cost
23	limit amounts specified in subparagraphs (B)
24	for a plan year that ends after 2006 shall be ad-
25	justed in the same manner as the annual deduct-



1	ible under section 1860D-2(b)(1) is annually ad-
2	justed under such section.
3	"(D) Adjustment contingency.—The
4	Secretary may adjust the percentage specified in
5	subparagraph (A) with respect to plan years that
6	end in a year in a manner so that the aggregate
7	expenditures in the year under this section are
8	the same as the aggregate expenditures that
9	would have been made under this section (taking
10	into account the effect of any adjustment under
11	subsection $(d)(1)(B)$ ) if paragraphs (1) and (2)
12	of subsection (a) had applied to qualified pre-
13	scription drug coverage instead of this para-
14	graph and subsection $(a)(3)$ .
15	"(4) Related definitions.—As used in this
16	section:
17	"(A) Employment-based retiree
18	HEALTH COVERAGE.—The term 'employment-
19	based retiree health coverage' means health in-
20	surance or other coverage of health care costs for
21	individuals eligible to enroll in a prescription
22	drug plan or MA-EFFS Rx plan under this part
23	(or for such individuals and their spouses and
24	dependents) under a group health plan (includ-

ing such a plan that is established or main-



1	tained under or pursuant to one or more collec-
2	tive bargaining agreements) based on their status
3	as retired participants in such plan.
4	"(B) QUALIFYING COVERED RETIREE.—The
5	term 'qualifying covered retiree' means an indi-
6	vidual who is eligible to obtain qualified pre-
7	scription drug coverage under section 1860D-1
8	but did not elect such coverage under this part
9	(either through a prescription drug plan or
10	through a MA-EFFS Rx plan) but is covered
11	under a qualified retiree prescription drug plan.
12	"(C) Sponsor.—The term 'sponsor' means
13	a plan sponsor, as defined in section $3(16)(B)$ of
14	the Employee Retirement Income Security Act of
15	1974, except that, in the case of a single-em-
16	ployer plan (as defined in section 3(41) of such
17	Act), such term means the employer of the plan
18	participants if such employer has been des-
19	ignated as the plan sponsor in all prior sum-
20	mary plan descriptions and annual reports
21	issued with respect to the plan under part 1 of
22	$subtitle\ B\ of\ title\ I\ of\ such\ Act.$
23	"(5) Construction.—Nothing in this subsection



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ferred to in subparagraph (B)) shall be



1	treated as primary coverage to which sec-
2	tion $1862(b)(2)(A)(i)$ is deemed to apply.
3	"(g) Computation of National Average Monthly
4	BID AMOUNT.—
5	"(1) In General.—For each year (beginning
6	with 2006) the Administrator shall compute a na-
7	tional average monthly bid amount equal to the aver-
8	age of the benchmark bid amounts for each prescrip-
9	tion drug plan and for each MA-EFFS Rx plan (as
10	computed under paragraph (2), but excluding plans
11	described in $section$ $1851(a)(2)(C)))$ $adjusted$ $under$
12	paragraph (4) to take into account reinsurance pay-
13	ments.
14	"(2) Benchmark bid amount defined.—For
15	purposes of this subsection, the term benchmark bid
16	amount' means, with respect to qualified prescription
17	drug coverage offered under—
18	"(A) a prescription drug plan that—
19	"(i) provides standard coverage (or al-
20	ternative prescription drug coverage the ac-
21	tuarial value of which is equivalent to that
22	of standard coverage), the PDP bid; or
23	"(ii) provides alternative prescription
24	drug coverage the actuarial value of which
25	is greater than that of standard coverage,



1	the PDP bid multiplied by the ratio of (I)
2	the actuarial value of standard coverage, to
3	(II) the actuarial value of the alternative
4	coverage; or
5	"(B) a MA-EFFS Rx plan, the portion of
6	the bid amount that is attributable to statutory
7	drug benefits (described in section
8	1853(a)(1)(A)(ii)(II)).
9	For purposes of subparagraph (A), the term 'PDP
10	bid' means, with respect to a prescription drug plan,
11	the bid amount for enrollment under the plan under
12	this part (determined without regard to any low-in-
13	come subsidy under section 1860D-7 or any late en-
14	rollment penalty under section 1860 $D$ -1( $c$ )(2)( $B$ )).
15	"(3) Weighted average.—
16	"(A) In General.—The monthly national
17	average monthly bid amount computed under
18	paragraph (1) shall be a weighted average, with
19	the weight for each plan being equal to the aver-
20	age number of beneficiaries enrolled under such
21	plan in the previous year.
22	"(B) Special rule for 2006.—For pur-
23	poses of applying this subsection for 2006, the
24	Administrator shall establish procedures for de-



1	termining the weighted average under subpara-
2	graph (A) for 2005.
3	"(4) Adjustment to Add back in value of
4	REINSURANCE SUBSIDIES.—The adjustment under
5	this paragraph, to take into account reinsurance pay-
6	ments under subsection (c) making up 30 percent of
7	total payments, is such an adjustment as will make
8	the national average monthly bid amount represent
9	represent 100 percent, instead of representing 70 per-
10	cent, of average payments under this part.
11	"(h) Coverage Year Defined.—For purposes of this
12	section, the term 'coverage year' means a calendar year in
13	which covered outpatient drugs are dispensed if a claim for
14	payment is made under the plan for such drugs, regardless
15	of when the claim is paid.
16	"SEC. 1860D-9. MEDICARE PRESCRIPTION DRUG TRUST
17	FUND.
18	"(a) In General.—There is created on the books of
19	the Treasury of the United States a trust fund to be known
20	as the 'Medicare Prescription Drug Trust Fund' (in this
21	section referred to as the 'Trust Fund'). The Trust Fund
22	shall consist of such gifts and bequests as may be made as
23	provided in section 201(i)(1), and such amounts as may
24	be deposited in, or appropriated to, such fund as provided
25	in this part. Except as otherwise provided in this section,



1	the provisions of subsections (b) through (i) of section 1842
2	shall apply to the Trust Fund in the same manner as they
3	apply to the Federal Supplementary Medical Insurance
4	Trust Fund under such section.
5	"(b) Payments From Trust Fund.—
6	"(1) In General.—The Managing Trustee shall
7	pay from time to time from the Trust Fund such
8	amounts as the Administrator certifies are necessary
9	to make—
10	"(A) payments under section 1860D-7 (re-
11	lating to low-income subsidy payments);
12	"(B) payments under section 1860D-8 (re-
13	lating to subsidy payments); and
14	"(C) payments with respect to administra
15	tive expenses under this part in accordance with
16	section $201(g)$ .
17	"(2) Transfers to medicaid account for in-
18	CREASED ADMINISTRATIVE COSTS.—The Managing
19	Trustee shall transfer from time to time from the
20	Trust Fund to the Grants to States for Medicaid ac-
21	count amounts the Administrator certifies are attrib
22	utable to increases in payment resulting from the ap-
23	plication of a higher Federal matching percentage
24	$under\ section\ 1935(b).$
25	"(c) Deposits Into Trust Fund.—



1	"(1) Low-income transfer.—There is hereby
2	transferred to the Trust Fund, from amounts appro-
3	priated for Grants to States for Medicaid, amounts
4	equivalent to the aggregate amount of the reductions
5	in payments under section 1903(a)(1) attributable to
6	the application of section $1935(c)$ .
7	"(2) Appropriations to cover government
8	CONTRIBUTIONS.—There are authorized to be appro-
9	priated from time to time, out of any moneys in the
10	Treasury not otherwise appropriated, to the Trust
11	Fund, an amount equivalent to the amount of pay-
12	ments made from the Trust Fund under subsection
13	(b), reduced by the amount transferred to the Trust
14	Fund under paragraph (1).
15	"(d) Relation to Solvency Requirements.—Any
16	provision of law that relates to the solvency of the Trust
17	Fund under this part shall take into account the Trust
18	Fund and amounts receivable by, or payable from, the
19	Trust Fund.
20	"SEC. 1860D-10. DEFINITIONS; APPLICATION TO MEDICARE
21	ADVANTAGE AND EFFS PROGRAMS; TREAT
22	MENT OF REFERENCES TO PROVISIONS IN
23	PART C.
24	"(a) DEFINITIONS.—For purposes of this part:



1	"(1) Covered outpatient drugs.—The term
2	'covered outpatient drugs' is defined in section
3	1860D-2(f).
4	"(2) Initial coverage limit.—The term 'ini-
5	tial coverage limit' means such limit as established
6	under section 1860D-2(b)(3), or, in the case of cov-
7	erage that is not standard coverage, the comparable
8	limit (if any) established under the coverage.
9	"(3) Medicare prescription drug trust
10	FUND.—The term 'Medicare Prescription Drug Trust
11	Fund' means the Trust Fund created under section
12	1860D-9(a).
13	"(4) PDP sponsor.—The term 'PDP sponsor'
14	means an entity that is certified under this part as
15	meeting the requirements and standards of this part
16	for such a sponsor.
17	"(5) Prescription drug plan.—The term 'pre-
18	scription drug plan' means health benefits coverage
19	that—
20	"(A) is offered under a policy, contract, or
21	plan by a PDP sponsor pursuant to, and in ac-
22	cordance with, a contract between the Adminis-
23	trator and the sponsor under section 1860D-
24	4(b);



1	"(B) provides qualified prescription drug
2	coverage; and
3	"(C) meets the applicable requirements of
4	the section 1860D-3 for a prescription drug
5	plan.
6	"(6) Qualified prescription drug cov-
7	ERAGE.—The term 'qualified prescription drug cov-
8	erage' is defined in section 1860D-2(a).
9	"(7) STANDARD COVERAGE.—The term 'standard
10	coverage' is defined in section 1860D-2(b).
11	"(b) Offer of Qualified Prescription Drug Cov-
12	ERAGE UNDER MEDICARE ADVANTAGE AND EFFS PRO-
13	GRAMS.—
14	"(1) As part of medicare advantage plan.—
15	Medicare Advantage organizations are required to
16	offer Medicare Advantage plans that include qualified
17	prescription drug coverage under part C pursuant to
18	section $1851(j)$ .
19	"(2) As part of effs plan.—EFFS organiza-
20	tions are required to offer EFFS plans that include
21	$qualified\ prescription\ drug\ coverage\ under\ part\ E$
22	pursuant to section 1860E-2(d).
23	"(c) Application of Part C Provisions Under
24	This Part.—For purposes of applying provisions of part
25	C under this part with respect to a prescription drug plan



1	and a PDP sponsor, unless otherwise provided in this part
2	such provisions shall be applied as if—
3	"(1) any reference to a Medicare Advantage or
4	other plan included a reference to a prescription drug
5	plan;
6	"(2) any reference to a provider-sponsored orga-
7	nization included a reference to a PDP sponsor;
8	"(3) any reference to a contract under section
9	1857 included a reference to a contract under section
10	$1860D-4(b); \ and$
11	"(4) any reference to part C included a reference
12	to this part.
13	"(d) Report on Pharmacy Services Provided to
14	Nursing Facility Patients.—
15	"(1) Review.—Within 6 months after the date of
16	the enactment of this section, the Secretary shall re-
17	view the current standards of practice for pharmacy
18	services provided to patients in nursing facilities.
19	"(2) Evaluations and recommendations.—
20	Specifically in the review under paragraph (1), the
21	Secretary shall—
22	"(A) assess the current standards of prac-
23	tice, clinical services, and other service require-
24	ments generally utilized for pharmacy services in
25	the long-term care setting;



1	"(B) evaluate the impact of those standards
2	with respect to patient safety, reduction of medi-
3	cation errors and quality of care; and
4	"(C) recommend (in the Secretary's report
5	under paragraph (3)) necessary actions and ap-
6	propriate reimbursement to ensure the provision
7	of prescription drugs to medicare beneficiaries
8	residing in nursing facilities in a manner con-
9	sistent with existing patient safety and quality
10	of care standards under applicable State and
11	Federal laws.
12	"(3) Report.—The Secretary shall submit a re-
13	port to the Congress on the Secretary's findings and
14	recommendations under this subsection, including a
15	detailed description of the Secretary's plans to imple-
16	ment this part in a manner consistent with applica-
17	ble State and Federal laws designed to protect the
18	safety and quality of care of nursing facility pa-
19	tients.".
20	(b) Additional Conforming Changes.—
21	(1) Conforming references to previous
22	PART D.—Any reference in law (in effect before the
23	date of the enactment of this Act) to part D of title
24	XVIII of the Social Security Act is deemed a reference

to part F of such title (as in effect after such date).



1	(2) Conforming amendment permitting waiv-
2	ER OF COST-SHARING.—Section $1128B(b)(3)$ (42)
3	$U.S.C.\ 1320a-7b(b)(3)) \ is \ amended$ —
4	(A) by striking "and" at the end of sub-
5	paragraph (E);
6	(B) by striking the period at the end of sub-
7	paragraph (F) and inserting "; and"; and
8	(C) by adding at the end the following new
9	subparagraph:
10	"(G) the waiver or reduction of any cost-sharing
11	imposed under part D of title XVIII.".
12	(3) Submission of legislative proposal.—
13	Not later than 6 months after the date of the enact-
14	ment of this Act, the Secretary of Health and Human
15	Services shall submit to the appropriate committees of
16	Congress a legislative proposal providing for such
17	technical and conforming amendments in the law as
18	are required by the provisions of this subtitle.
19	(c) Study on Transitioning Part B Prescription
20	Drug Coverage.—Not later than January 1, 2005, the
21	Medicare Benefits Administrator shall submit a report to
22	Congress that makes recommendations regarding methods
23	for providing benefits under part D of title XVIII of the
24	Social Security Act for outpatient prescription drugs for
25	which benefits are provided under part B of such title.



1	SEC. 102. OFFERING OF QUALIFIED PRESCRIPTION DRUG
2	COVERAGE UNDER MEDICARE ADVANTAGE
3	AND ENHANCED FEE-FOR-SERVICE (EFFS)
4	PROGRAM.
5	(a) Medicare Advantage.—Section 1851 (42 U.S.C.
6	1395w-21) is amended by adding at the end the following
7	new subsection:
8	"(j) Availability of Prescription Drug Benefits
9	and Subsidies.—
10	"(1) Offering of qualified prescription
11	DRUG COVERAGE.—A Medicare Advantage organiza-
12	tion on and after January 1, 2006—
13	"(A) may not offer a Medicare Advantage
14	plan described in section $1851(a)(2)(A)$ in an
15	area unless either that plan (or another Medicare
16	Advantage plan offered by the organization in
17	that area) includes qualified prescription drug
18	coverage; and
19	"(B) may not offer the prescription drug
20	coverage (other than that required under parts A
21	and B) to an enrollee under a Medicare Advan-
22	tage plan, unless such drug coverage is at least
23	qualified prescription drug coverage and unless
24	the requirements of this subsection with respect
25	to such coverage are met.



1	"(2) Requirement for election of part i
2	COVERAGE TO OBTAIN QUALIFIED PRESCRIPTION
3	DRUG COVERAGE.—For purposes of this part, an in-
4	dividual who has not elected qualified prescription
5	drug coverage under section 1860D-1(b) shall be
6	treated as being ineligible to enroll in a Medicare Ad-
7	vantage plan under this part that offers such cov-
8	erage.
9	"(3) Compliance with certain additional
10	BENEFICIARY PROTECTIONS FOR PRESCRIPTION DRUG
11	COVERAGE.—With respect to the offering of qualified
12	prescription drug coverage by a Medicare Advantage
13	organization under this part on and after January 1,
14	2006, the organization and plan shall meet the re-
15	quirements of subsections (a) through (d) of section
16	1860D-3 in the same manner as they apply to a
17	PDP sponsor and a prescription drug plan under
18	part D and shall submit to the Administrator the in-
19	formation described in section $1860D-6(a)(2)$ . The
20	Administrator shall waive such requirements to the
21	extent the Administrator determines that such re-
22	quirements duplicate requirements otherwise applica-
23	ble to the organization or plan under this part.
24	"(4) Availability of premium and cost-shar-



"(4) Availability of Premium and Cost-Sharing subsidies.—In the case of low-income individ-

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1	uals who are enrolled in a Medicare Advantage plan
2	that provides qualified prescription drug coverage,
3	premium and cost-sharing subsidies are provided for
4	such coverage under section 1860D-7.
5	"(5) Availability of direct and reinsur-
6	ANCE SUBSIDIES TO REDUCE BIDS AND PREMIUMS.—
7	Medicare Advantage organizations are provided direct
8	and reinsurance subsidy payments for providing
9	qualified prescription drug coverage under this part
10	under section 1860D–8.
11	"(6) Consolidation of drug and non-drug
12	PREMIUMS.—In the case of a Medicare Advantage
13	plan that includes qualified prescription drug cov-
14	erage, with respect to an enrollee in such plan there
15	shall be a single premium for both drug and non-drug
16	coverage provided under the plan.
17	"(7) Transition in initial enrollment pe-
18	RIOD.—Notwithstanding any other provision of this
19	part, the annual, coordinated election period under
20	subsection (e)(3)(B) for 2006 shall be the 6-month pe-
21	riod beginning with November 2005.
22	"(8) Qualified prescription drug coverage;
23	STANDARD COVERAGE.—For purposes of this part, the

terms 'qualified prescription drug coverage' and



1	'standard coverage' have the meanings given such
2	terms in section 1860D-2.
3	"(9) Special rules for private fee-for-
4	Service plans.— With respect to a Medicare Advan-
5	tage plan described in section $1851(a)(2)(C)$ that of-
6	fers qualified prescription drug coverage—
7	"(A) REQUIREMENTS REGARDING NEGO-
8	TIATED PRICES.—Subsections (a)(1) and (d)(1)
9	of section 1860D-2 shall not be construed to re-
10	quire the plan to negotiate prices or discounts
11	but shall apply to the extent the plan does so.
12	"(B) Modification of Pharmacy Partici-
13	PATION REQUIREMENT.—If the plan provides ac-
14	cess, without charging additional copayments, to
15	all pharmacies without regard to whether they
16	are participating pharmacies in a network, sec-
17	tion $1860D-3(c)(1)(A)(iii)$ shall not apply to the
18	plan.
19	"(C) Drug utilization management pro-
20	GRAM NOT REQUIRED.—The requirements of sec-
21	tion $1860D-3(d)(1)(A)$ shall not apply to the
22	plan.
23	"(D) Non-participating pharmacy dis-
24	CLOSURE EXCEPTION.—If the plan provides cov-
25	erage for drugs purchased from all pharmacies,



1	without entering into contracts or agreements
2	with pharmacies to provide drugs to enrollees
3	covered by the plan, section 1860D-3(d)(5) shall
4	not apply to the plan.".
5	(b) Application to EFFS Plans.—Subsection (d) of
6	section 1860E-2, as added by section 201(a), is amended
7	to read as follows:
8	"(d) Availability of Prescription Drug Benefits
9	and Subsidies.—
10	"(1) Offering of qualified prescription
11	DRUG COVERAGE.—An EFFS organization—
12	"(A) may not offer an EFFS plan in an
13	area unless either that plan (or another EFFS
14	plan offered by the organization in that area)
15	includes qualified prescription drug coverage;
16	and
17	"(B) may not offer the prescription drug
18	coverage (other than that required under parts $A$
19	and B) to an enrollee under an EFFS plan, un-
20	less such drug coverage is at least qualified pre-
21	scription drug coverage and unless the require-
22	ments of this subsection with respect to such cov-
23	erage are met.
24	"(2) Requirement for election of part d
25	COVERAGE TO OBTAIN QUALIFIED PRESCRIPTION



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1	DRUG COVERAGE.—For purposes of this part, an in-
2	dividual who has not elected qualified prescription
3	drug coverage under section 1860D-1(b) shall be
4	treated as being ineligible to enroll in an EFFS plan
5	under this part that offers such coverage.
6	"(3) Compliance with certain additional
7	BENEFICIARY PROTECTIONS FOR PRESCRIPTION DRUG
8	COVERAGE.—With respect to the offering of qualified
9	prescription drug coverage by an EFFS organization
10	under this part, the organization and plan shall meet
11	the requirements of subsections (a) through (d) of sec-
12	tion 1860D-3 in the same manner as they apply to
13	a PDP sponsor and a prescription drug plan under
14	part D and shall submit to the Administrator the in-
15	formation described in section $1860D-6(a)(2)$ . The
16	Administrator shall waive such requirements to the
17	extent the Administrator determines that such re-
18	quirements duplicate requirements otherwise applica-
19	ble to the organization or plan under this part.
20	"(4) Availability of premium and cost-shar-
21	ING SUBSIDIES.—In the case of low-income individ-
22	uals who are enrolled in an EFFS plan that provides



ING SUBSIDIES.—In the case of low-income individuals who are enrolled in an EFFS plan that provides qualified prescription drug coverage, premium and cost-sharing subsidies are provided for such coverage under section 1860D–7.

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1	"(5) Availability of direct and reinsur-
2	ANCE SUBSIDIES TO REDUCE BIDS AND PREMIUMS.—
3	EFFS organizations are provided direct and reinsur-
4	ance subsidy payments for providing qualified pre-
5	scription drug coverage under this part under section
6	1860D-8.
7	"(6) Consolidation of drug and non-drug
8	PREMIUMS.—In the case of an EFFS plan that in-
9	cludes qualified prescription drug coverage, with re-
10	spect to an enrollee in such plan there shall be a sin-
11	gle premium for both drug and non-drug coverage
12	provided under the plan.
13	"(7) Qualified prescription drug coverage,
14	STANDARD COVERAGE.—For purposes of this part, the
15	terms 'qualified prescription drug coverage' and
16	'standard coverage' have the meanings given such
17	terms in section 1860D-2.".
18	(c) Conforming Amendments.—Section 1851 (42
19	U.S.C. 1395w-21) is amended—
20	(1) in subsection (a)(1)—
21	(A) by inserting "(other than qualified pre-
22	scription drug benefits)" after "benefits";
23	(B) by striking the period at the end of sub-
24	paragraph (B) and inserting a comma; and



1	(C) by adding after and below subpara-
2	graph (B) the following:
3	"and may elect qualified prescription drug coverage
4	in accordance with section 1860D-1."; and
5	(2) in subsection $(g)(1)$ , by inserting "and sec-
6	tion $1860D-1(c)(2)(B)$ " after "in this subsection".
7	(d) Effective Date.—The amendments made by this
8	section apply to coverage provided on or after January 1,
9	2006.
10	SEC. 103. MEDICAID AMENDMENTS.
11	(a) Determinations of Eligibility for Low-In-
12	COME SUBSIDIES.—
13	(1) Requirement.—Section 1902(a) (42 U.S.C.
14	1396a(a)) is amended—
15	(A) by striking "and" at the end of para-
16	graph (64);
17	(B) by striking the period at the end of
18	paragraph (65) and inserting "; and"; and
19	(C) by inserting after paragraph (65) the
20	following new paragraph:
21	"(66) provide for making eligibility determina-
22	tions under section 1935(a).".
23	(2) NEW SECTION.—Title XIX is further
24	amended—



1	(A) by redesignating section 1935 as section
2	1936; and
3	(B) by inserting after section 1934 the fol-
4	lowing new section:
5	"SPECIAL PROVISIONS RELATING TO MEDICARE
6	PRESCRIPTION DRUG BENEFIT
7	"Sec. 1935. (a) Requirement for Making Eligi-
8	BILITY DETERMINATIONS FOR LOW-INCOME SUBSIDIES.—
9	As a condition of its State plan under this title under sec-
10	tion 1902(a)(66) and receipt of any Federal financial as-
11	sistance under section 1903(a), a State shall—
12	"(1) make determinations of eligibility for pre-
13	mium and cost-sharing subsidies under (and in ac-
14	cordance with) section 1860D-7;
15	"(2) inform the Administrator of the Medicare
16	Benefits Administration of such determinations in
17	cases in which such eligibility is established; and
18	"(3) otherwise provide such Administrator with
19	such information as may be required to carry out
20	part D of title XVIII (including section 1860D-7).
21	"(b) Payments for Additional Administrative
22	Costs.—
23	"(1) In general.—The amounts expended by a
24	State in carrying out subsection (a) are, subject to
25	paragraph (2), expenditures reimbursable under the
26	appropriate paragraph of section 1903(a); except



1	that, notwithstanding any other provision of such sec-
2	tion, the applicable Federal matching rates with re-
3	spect to such expenditures under such section shall be
4	increased as follows (but in no case shall the rate as
5	so increased exceed 100 percent):
6	"(A) For expenditures attributable to costs
7	incurred during 2005, the otherwise applicable
8	Federal matching rate shall be increased by 10
9	percent of the percentage otherwise payable (but
10	for this subsection) by the State.
11	"(B)(i) For expenditures attributable to
12	costs incurred during 2006 and each subsequent
13	year through 2013, the otherwise applicable Fed-
14	eral matching rate shall be increased by the ap-
15	plicable percent (as defined in clause (ii)) of the
16	percentage otherwise payable (but for this sub-
17	section) by the State.
18	"(ii) For purposes of clause (i), the 'appli-
19	cable percent' for—
20	"(I) 2006 is 20 percent; or
21	"(II) a subsequent year is the applica-
22	ble percent under this clause for the pre-
23	vious year increased by 10 percentage
24	points.



1	"(C) For expenditures attributable to costs
2	incurred after 2013, the otherwise applicable
3	Federal matching rate shall be increased to 100
4	percent.
5	"(2) Coordination.—The State shall provide
6	the Administrator with such information as may be
7	necessary to properly allocate administrative expendi-
8	tures described in paragraph (1) that may otherwise
9	be made for similar eligibility determinations.".
10	(b) Phased-In Federal Assumption of Medicaid
11	Responsibility for Premium and Cost-Sharing Sub-
12	SIDIES FOR DUALLY ELIGIBLE INDIVIDUALS.—
13	(1) In General.—Section 1903(a)(1) (42 U.S.C.
14	1396b(a)(1)) is amended by inserting before the semi-
15	colon the following: ", reduced by the amount com-
16	puted under section 1935(c)(1) for the State and the
17	quarter".
18	(2) Amount described.—Section 1935, as in-
19	serted by subsection (a)(2), is amended by adding at
20	the end the following new subsection:
21	"(c) Federal Assumption of Medicaid Prescrip-
22	TION DRUG COSTS FOR DUALLY-ELIGIBLE BENE-
23	FICIARIES.—
24	"(1) In General.—For purposes of section
25	1903(a)(1), for a State that is one of the 50 States



1	or the District of Columbia for a calendar quarter in
2	a year (beginning with 2005) the amount computed
3	under this subsection is equal to the product of the
4	following:
5	"(A) Medicare subsidies.—The total
6	amount of payments made in the quarter under
7	section 1860D-7 (relating to premium and cost-
8	sharing prescription drug subsidies for low-in-
9	come medicare beneficiaries) that are attrib-
10	utable to individuals who are residents of the
11	State and are entitled to benefits with respect to
12	prescribed drugs under the State plan under this
13	title (including such a plan operating under a
14	waiver under section 1115).
15	"(B) State matching rate.—A propor-
16	tion computed by subtracting from 100 percent
17	the Federal medical assistance percentage (as de-
18	fined in section 1905(b)) applicable to the State
19	and the quarter.
20	"(C) Phase-out proportion.—The phase-
21	out proportion (as defined in paragraph (2)) for
22	the quarter.
23	"(2) Phase-out proportion.—For purposes of
24	paragraph (1)(C), the 'phase-out proportion' for a



25

calendar quarter in—

1	"(A) 2006 is 93-1/3 percent;
2	"(B) a subsequent year before 2021, is the
3	phase-out proportion for calendar quarters in the
4	previous year decreased by 6-2/3 percentage
5	points; or
6	"(C) a year after 2020 is 0 percent.".
7	(c) Medicaid Providing Wrap-Around Bene-
8	FITS.—Section 1935, as so inserted and amended, is further
9	amended by adding at the end the following new subsection:
10	"(d) Additional Provisions.—
11	"(1) MEDICAID AS SECONDARY PAYOR.—In the
12	case of an individual who is entitled to qualified pre-
13	scription drug coverage under a prescription drug
14	plan under part D of title XVIII (or under a MA-
15	EFFS Rx plan under part C or E of such title) and
16	medical assistance for prescribed drugs under this
17	title, medical assistance shall continue to be provided
18	under this title (other than for copayment amounts
19	specified in $section$ $1860D-7(a)(1)(B)$ , $notwith$
20	standing section 1916) for prescribed drugs to the ex-
21	tent payment is not made under the prescription drug
22	plan or MA-EFFS Rx plan selected by the individual.
23	"(2) Condition.—A State may require, as a
24	condition for the receipt of medical assistance under
25	this title with respect to prescription drug benefits for



1	an individual eligible to obtain qualified prescription
2	drug coverage described in paragraph (1), that the in-
3	dividual elect qualified prescription drug coverage
4	under section 1860D-1.".
5	(d) Treatment of Territories.—
6	(1) In general.—Section 1935, as so inserted
7	and amended, is further amended—
8	(A) in subsection (a) in the matter pre-
9	ceding paragraph (1), by inserting "subject to
10	subsection (e)" after "section 1903(a)";
11	(B) in subsection (c)(1), by inserting "sub-
12	ject to subsection (e)" after "1903(a)(1)"; and
13	(C) by adding at the end the following new
14	subsection:
15	"(e) Treatment of Territories.—
16	"(1) In general.—In the case of a State, other
17	than the 50 States and the District of Columbia—
18	"(A) the previous provisions of this section
19	shall not apply to residents of such State; and
20	"(B) if the State establishes a plan de-
21	scribed in paragraph (2) (for providing medical
22	assistance with respect to the provision of pre-
23	scription drugs to medicare beneficiaries), the
24	amount otherwise determined under section
25	1108(f) (as increased under section $1108(a)$ ) for



1	the State shall be increased by the amount speci-
2	fied in paragraph (3).
3	"(2) Plan.—The plan described in this para-
4	graph is a plan that—
5	"(A) provides medical assistance with re-
6	spect to the provision of covered outpatient drugs
7	(as defined in section 1860D-2(f)) to low-income
8	medicare beneficiaries; and
9	"(B) assures that additional amounts re-
10	ceived by the State that are attributable to the
11	operation of this subsection are used only for
12	such assistance.
13	"(3) Increased amount.—
14	"(A) In general.—The amount specified
15	in this paragraph for a State for a year is equal
16	to the product of—
17	"(i) the aggregate amount specified in
18	subparagraph (B); and
19	"(ii) the amount specified in section
20	1108(g)(1) for that State, divided by the
21	sum of the amounts specified in such section
22	for all such States.
23	"(B) AGGREGATE AMOUNT.—The aggregate
24	amount specified in this subparagraph for—
25	"(i) 2006, is equal to \$25,000,000; or



1	"(ii) a subsequent year, is equal to the
2	aggregate amount specified in this subpara-
3	graph for the previous year increased by
4	annual percentage increase specified in sec-
5	tion $1860D-2(b)(5)$ for the year involved.
6	"(4) Report.—The Administrator shall submit
7	to Congress a report on the application of this sub-
8	section and may include in the report such rec-
9	ommendations as the Administrator deems appro-
10	priate.".
11	(2) Conforming amendment.—Section 1108(f)
12	(42 U.S.C. 1308(f)) is amended by inserting "and sec-
13	tion $1935(e)(1)(B)$ " after "Subject to subsection $(g)$ ".
14	(e) Amendment to Best Price.—Section
15	1927(c)(1)(C)(i) (42 U.S.C. $1396r-8(c)(1)(C)(i)$ ) is
16	amended—
17	(1) by striking "and" at the end of subclause
18	(III);
19	(2) by striking the period at the end of subclause
20	(IV) and inserting "; and"; and
21	(3) by adding at the end the following new sub-
22	clause:
23	"(V) any prices charged which are
24	negotiated by a prescription drug plan
25	under part D of title XVIII, by a MA-



1	EFFS Rx plan under part C or E of
2	such title with respect to covered out-
3	patient drugs, or by a qualified retiree
4	prescription drug plan (as defined in
5	section $1860D-8(f)(1)$ ) with respect to
6	such drugs on behalf of individuals en-
7	titled to benefits under part A or en-
8	rolled under part B of such title.".
9	SEC. 104. MEDIGAP TRANSITION.
10	(a) In General.—Section 1882 (42 U.S.C. 1395ss) is
11	amended by adding at the end the following new subsection:
12	"(v) Coverage of Prescription Drugs.—
13	"(1) In general.—Notwithstanding any other
14	provision of law, except as provided in paragraph (3)
15	no new medicare supplemental policy that provides
16	coverage of expenses for prescription drugs may be
17	issued under this section on or after January 1, 2006,
18	to an individual unless it replaces a medicare supple-
19	mental policy that was issued to that individual and
20	that provided some coverage of expenses for prescrip-
21	tion drugs. Nothing in this subsection shall be con-
22	strued as preventing the policy holder of a medicare
23	supplemental policy issued before January 1, 2006,
24	from continuing to receive benefits under such notice



25

on and after such date.

1	"(2) Issuance of substitute policies for
2	BENEFICIARIES ENROLLED WITH A PLAN UNDER PART
3	D.—
4	"(A) In general.—The issuer of a medi-
5	care supplemental policy—
6	"(i) may not deny or condition the
7	issuance or effectiveness of a medicare sup-
8	plemental policy that has a benefit package
9	classified as 'A', 'B', 'C', 'D', 'E', 'F', or 'G'
10	(under the standards established under sub-
11	section $(p)(2)$ ) and that is offered and is
12	available for issuance to new enrollees by
13	such issuer;
14	"(ii) may not discriminate in the pric-
15	ing of such policy, because of health status,
16	claims experience, receipt of health care, or
17	medical condition; and
18	"(iii) may not impose an exclusion of
19	benefits based on a pre-existing condition
20	under such policy,
21	in the case of an individual described in sub-
22	paragraph (B) who seeks to enroll under the pol-
23	icy not later than 63 days after the date of the
24	termination of enrollment described in such
25	paragraph and who submits evidence of the date



1	of termination or disenrollment along with the
2	application for such medicare supplemental pol-
3	icy.
4	"(B) Individual covered.—An individual
5	described in this subparagraph is an individual
6	who—
7	"(i) enrolls in a prescription drug
8	plan under part D; and
9	"(ii) at the time of such enrollment
10	was enrolled and terminates enrollment in
11	a medicare supplemental policy which has a
12	benefit package classified as 'H', 'I', or 'J'
13	under the standards referred to in subpara-
14	graph (A)(i) or terminates enrollment in a
15	policy to which such standards do not apply
16	but which provides benefits for prescription
17	drugs.
18	"(C) Enforcement.—The provisions of
19	paragraph (4) of subsection (s) shall apply with
20	respect to the requirements of this paragraph in
21	the same manner as they apply to the require-
22	ments of such subsection.
23	"(3) New Standards.—In applying subsection
24	(p)(1)(E) (including permitting the NAIC to revise
25	its model regulations in response to changes in law)



1	with respect to the change in benefits resulting from
2	title I of the Medicare Prescription Drug and Mod-
3	ernization Act of 2003, with respect to policies issued
4	to individuals who are enrolled in a plan under part
5	D, the changes in standards shall only provide for
6	substituting (for the benefit packages described in
7	paragraph (2)(B)(ii) that included coverage for pre-
8	scription drugs) two benefit packages that may pro-
9	vide for coverage of cost-sharing (other than the pre-
10	scription drug deductible) with respect to qualified
11	prescription drug coverage under such part. The two
12	benefit packages shall be consistent with the following.
13	"(A) First new policy.—The policy de-
14	scribed in this subparagraph has the following
15	benefits, notwithstanding any other provision of
16	this section relating to a core benefit package:
17	"(i) Coverage of 50 percent of the cost-
18	sharing otherwise applicable under parts A
19	and B, except coverage of 100 percent of
20	any cost-sharing otherwise applicable for
21	preventive benefits.
22	"(ii) No coverage of the part B deduct-
23	ible.



1	"(iii) Coverage for all hospital coinsur-
2	ance for long stays (as in the current core
3	benefit package).
4	"(iv) A limitation on annual out-of-
5	pocket expenditures under parts A and B to
6	\$4,000 in 2005 (or, in a subsequent year, to
7	such limitation for the previous year in-
8	creased by an appropriate inflation adjust-
9	ment specified by the Secretary).
10	"(B) Second New Policy.—The policy de-
11	scribed in this subparagraph has the same bene-
12	fits as the policy described in subparagraph (A),
13	except as follows:
14	"(i) Substitute '75 percent' for '50 per-
15	cent' in clause (i) of such subparagraph.
16	"(ii) Substitute '\$2,000' for '\$4,000' in
17	clause (iv) of such subparagraph.
18	"(4) Construction.—Any provision in this sec-
19	tion or in a medicare supplemental policy relating to
20	guaranteed renewability of coverage shall be deemed
21	to have been met through the offering of other coverage
22	under this subsection.".
23	(b) NAIC REPORT TO CONGRESS ON MEDIGAP MOD-
24	ERNIZATION.—The Secretary shall request the National As-
25	sociation of Insurance Commissioners to submit to Con-



1	gress, not later than 18 months after the date of the enact
2	ment of this Act, a report that includes recommendations
3	on the modernization of coverage under the medigap pro-
4	gram under section 1882 of the Social Security Act (42
5	U.S.C. 1395ss).
6	SEC. 105. MEDICARE PRESCRIPTION DRUG DISCOUNT CARL
7	ENDORSEMENT PROGRAM.
8	(a) In General.—Title XVIII is amended by insert-
9	ing after section 1806 the following new sections:
10	"MEDICARE PRESCRIPTION DRUG DISCOUNT CARD
11	ENDORSEMENT PROGRAM
12	"Sec. 1807. (a) Establishment of Program.—
13	"(1) In General.—The Secretary (or the Medi
14	care Benefits Administrator pursuant to section
15	1809(c)(3)(C)) shall establish a program to endorse
16	prescription drug discount card programs (each such
17	program referred to as an 'endorsed program') that
18	meet the requirements of this section in order to pro-
19	vide access to prescription drug discounts for medi-
20	care beneficiaries throughout the United States. The
21	Secretary shall make available to medicare bene-
22	ficiaries information regarding endorsed programs
23	under this section.
24	"(2) Limited Period of Operation.—The Sec-
25	retary shall begin the program under this section as

soon as possible, but in no case later than 90 days



1	after the date of the enactment of this section. The
2	Secretary shall provide for an appropriate transition
3	and discontinuation of such program at the time
4	medicare prescription drug benefits first become
5	available under part D.
6	"(b) Requirements for Card Endorsement Pro-
7	GRAM.—The Secretary may not endorse a prescription drug
8	discount card program under this section unless the pro-
9	gram meets the following requirements:
10	"(1) Savings to medicare beneficiaries.—
11	The program passes on to medicare beneficiaries who
12	enroll in the program discounts, rebates, and other
13	price concessions on prescription drugs, including dis-
14	counts negotiated with pharmacies and manufactur-
15	ers.
16	"(2) Prohibition on application only to
17	MAIL ORDER.—The program applies to drugs that are
18	available other than solely through mail order.
19	"(3) Beneficiary services.—The program
20	provides pharmaceutical support services, such as
21	education and counseling, and services to prevent ad-
22	verse drug interactions.
23	"(4) Information.—The program makes avail-
24	able to medicare beneficiaries through the Internet
25	and otherwise information, including information on



1	enrollment fees, prices charged to beneficiaries, and
2	services offered under the program, that the Secretary
3	identifies as being necessary to provide for informed
4	choice by beneficiaries among endorsed programs.
5	"(5) Demonstrated experience.—The pro-
6	gram is operated directly, or through arrangements
7	with affiliated organization, by an entity that has
8	demonstrated experience and expertise in operating
9	such a program or a similar program.
10	"(6) Quality assurance.—Such operating en-
11	tity has in place adequate procedures for assuring
12	quality service under the program.
13	"(7) Enrollment fees.—The program may
14	charge an annual enrollment fee, but the amount of
15	such annual fee may not exceed \$30. A State may
16	pay some or all of the fee for individuals residing in
17	the State.
18	"(8) Confidentiality protections.—The pro-
19	gram implements policies and procedures to safeguard
20	the use and disclosure of program beneficiaries' indi-
21	vidually identifiable health information in a manner
22	consistent with the Federal regulations (concerning
23	the privacy of individually identifiable health infor-

mation) promulgated under section 264(c) of the



1	Health Insurance Portability and Accountability Act
2	of 1996.
3	"(9) Periodic reports to secretary.—The
4	entity operating the program shall submit to the Sec-
5	retary periodic reports on performance, utilization,
6	finances, and such other matters as the Secretary
7	may specify.
8	"(10) Additional beneficiary protec-
9	TIONS.—The program meets such additional require-
10	ments as the Secretary identifies to protect and pro-
11	mote the interest of medicare beneficiaries, including
12	requirements that ensure that beneficiaries are not
13	charged more than the lower of the negotiated retail
14	price or the usual and customary price.
15	The prices negotiated by a prescription drug discount card
16	program endorsed under this section shall (notwithstanding
17	any other provision of law) not be taken into account for
18	the purposes of establishing the best price under section
19	1927(c)(1)(C).
20	"(c) Program Operation.—The Secretary shall oper-
21	ate the program under this section consistent with the fol-
22	lowing:
23	"(1) Promotion of informed choice.—In
24	order to promote informed choice among endorsed

prescription drug discount card programs, the Sec-



1	retary shall provide for the dissemination of informa-
2	tion which compares the prices and services of such
3	programs in a manner coordinated with the dissemi-
4	nation of educational information on Medicare Ad-
5	vantage plans under part C.
6	"(2) Oversight.—The Secretary shall provide
7	appropriate oversight to ensure compliance of en-
8	dorsed programs with the requirements of this section,
9	including verification and disclosure (upon request)
10	of the discounts and services provided, the amount of
11	dispensing fees recognized, and audits under section
12	1860D-2(d)(3).
13	"(3) Use of medicare toll-free number.—
14	The Secretary shall provide through the 1-800-medi-
15	care toll free telephone number for the receipt and re-
16	sponse to inquiries and complaints concerning the
17	program and programs endorsed under this section.
18	"(4) Sanctions for abusive practices.—The
19	Secretary may implement intermediate sanctions or
20	may revoke the endorsement of a program in the case
21	of a program that the Secretary determines no longer
22	meets the requirements of this section or that has en-
23	gaged in false or misleading marketing practices.
24	"(5) Enrollment practices.—A medicare ben-



1 dorsed program at any time. A medicare beneficiary 2 may change the endorsed program in which the bene-3 ficiary is enrolled, but may not make such change 4 until the beneficiary has been enrolled in a program 5 for a minimum period of time specified by the Sec-6 retary. 7 "(d) AUTHORIZATION OF APPROPRIATIONS.—There 8 are authorized to be appropriated such sums as may be nec-9 essary to carry out this section. 10 "(e) Interim, Final Regulatory Authority.—In 11 order to carry out this section in a timely manner, the Sec-12 retary may promulgate regulations that take effect on an 13 interim basis, after notice and pending opportunity for 14 public comment. 15 "TRANSITIONAL PRESCRIPTION DRUG ASSISTANCE 16 PROGRAM FOR LOW-INCOME BENEFICIARIES 17 "Sec. 1807A. (a) PURPOSE.—The purpose of this sec-18 tion is to provide low-income medicare beneficiaries with 19 incomes below 150 percent of the Federal poverty level immediate assistance in the purchase of covered outpatient 21 prescription drugs during the period before the program 22 under part D becomes effective. 23 "(b) APPROPRIATIONS.—For the purpose of carrying out this section, there is appropriated, out of any money in the Treasury not otherwise appropriated— 25



"(1) for fiscal year 2004, \$2,000,000,000; and

1	"(2) for fiscal year 2005, \$3,000,000,000.
2	"(c) Eligibility.—
3	"(1) In general.—The Secretary shall establish
4	eligibility standards consistent with this subsection.
5	"(2) Specifics.—In no case shall an individual
6	be eligible for assistance under this section unless the
7	individual—
8	"(A) is entitled to benefits under part A or
9	enrolled under part B;
10	"(B) has income that is at or below 150
11	percent of the Federal poverty line;
12	"(C) meets the resources requirement de-
13	scribed in section 1905(p)(1)(C);
14	"(D) is enrolled under a prescription drug
15	discount card program under section 1807 (or
16	under an alternative program authorized under
17	subsection (d)(2)); and
18	"(E) is not eligible for coverage of, or assist-
19	ance for, outpatient prescription drugs under
20	any of the following:
21	"(i) A medicaid plan under title XIX
22	(including under any waiver approved
23	under section 1115).
24	"(ii) Enrollment under a group health
25	plan or health insurance coverage.



1	"(iii) Enrollment under a medicare
2	supplemental insurance policy.
3	"(iv) Chapter 55 of title 10, United
4	States Code (relating to medical and dental
5	care for members of the uniformed services).
6	"(v) Chapter 17 of title 38, United
7	States Code (relating to Veterans' medical
8	care).
9	"(vi) Enrollment under a plan under
10	chapter 89 of title 5, United States Code
11	(relating to the Federal employees' health
12	benefits program).
13	"(vii) The Indian Health Care Im-
14	provement Act (25 U.S.C. 1601 et seq.).
15	"(d) Form of Assistance.—
16	"(1) In general.—Subject to paragraph (2), the
17	assistance under this section to an eligible individual
18	shall be in such form as the Secretary shall specify,
19	including the use of a debit card mechanism to pay
20	for drugs purchased through the use of the prescrip-
21	tion drug discount card program to eligible individ-
22	uals who are enrolled in such program.
23	"(2) Through alternative state pro-
24	GRAM.—A State may apply to the Secretary for au-
25	thorization to provide the assistance under this sec-



1	tion to an eligible individual through a State phar-
2	maceutical assistance program or private program of
3	pharmaceutical assistance. The Secretary shall not
4	authorize the use of such a program unless the Sec-
5	retary finds that the program—
6	"(A) was in existence before the date of the
7	enactment of this section; and
8	"(B) is reasonably designed to provide for
9	pharmaceutical assistance for a number of indi-
10	viduals, and in a scope, that is not less than the
11	number of individuals, and minimum required
12	amount, that would occur if the provisions of
13	this paragraph had not applied in the State.
14	"(3) Relationship to discounts.—The assist-
15	ance provided under this section is in addition to the
16	discount otherwise available to individuals enrolled in
17	prescription drug discount card programs who are
18	not eligible individuals.
19	"(4) Limitation on Assistance.—
20	"(A) In General.—The assistance under
21	this section for an eligible individual shall be
22	limited to assistance—
23	"(i) for covered outpatient drugs (as
24	defined for purposes of part D) and for en-



1	rollment fees imposed under prescription
2	drug discount card programs; and
3	"(ii) for expenses incurred—
4	"(I) on and after the date the in-
5	dividual is both enrolled in the pre-
6	scription drug discount card program
7	and determined to be an eligible indi-
8	vidual under this section; and
9	"(II) before the date benefits are
10	first available under the program
11	$under\ part\ D.$
12	"(B) Authority.—The Secretary shall take
13	such steps as may be necessary to assure compli-
14	ance with the expenditure limitations described
15	in subsection (b).
16	"(e) Payment of Federal Subsidy to Sponsors.—
17	"(1) In general.—Insofar as assistance is pro-
18	vided under this section through programs under sec-
19	tion 1807, the Secretary shall make payment (within
20	the amounts under subsection (b), less the administra-
21	tive costs relating to determinations of eligibility) to
22	the sponsor of the prescription drug discount card
23	program (or to a State or other entity operating an
24	$alternative\ program\ under\ subsection\ (d)(2))$ in
25	which an eligible individual is enrolled of the amount



1	of the assistance provided by the sponsor pursuant to
2	$this\ section.$
3	"(2) PERIODIC PAYMENTS.—Payments under
4	this subsection shall be made on a monthly or other
5	periodic installment basis, based upon estimates of the
6	Secretary and shall be reduced or increased to the ex-
7	tent of any overpayment or underpayment which the
8	Secretary determines was made under this section for
9	any prior period and with respect to which adjust-
10	ment has not already been made under this para-
11	graph.
12	"(f) Definitions.—For purposes of this section:
13	"(1) Eligible individual.—The term 'eligible
14	individual' means an individual who is determined
15	by a State to be eligible for assistance under this sec-
16	tion.
17	"(2) Prescription drug discount card pro-
18	GRAM.—The term 'prescription drug discount card
19	program' means such a program that is endorsed
20	under section 1807.
21	"(3) Sponsor.—The term 'sponsor' means the
22	sponsor of a prescription drug discount card pro-
23	gram, or, in the case of an alternative program au-
24	thorized under subsection $(d)(2)$ , the State or other

entity operating the program.".



1	(b) Conforming Amendment.—Section
2	1927(c)(1)(C)(i)(V) (42 U.S.C. $1396r-8(c)(1)(C)(i)(V)$ ), as
3	added by section 103(e), is amended by striking "or by a
4	qualified retiree prescription drug plan (as defined in sec-
5	tion 1860D-8(f)(1))" and inserting "by a qualified retired
6	prescription drug plan (as defined in section 1860D-
7	8(f)(1)), or by a prescription drug discount card program
8	endorsed under section 1807".
9	SEC. 106. DISCLOSURE OF RETURN INFORMATION FOR PUR
10	POSES OF CARRYING OUT MEDICARE CATA
11	STROPHIC PRESCRIPTION DRUG PROGRAM.
12	(a) In General.—Subsection (l) of section 6103 of the
13	Internal Revenue Code of 1986 (relating to disclosure of re-
14	turns and return information for purposes other than tax
15	administration) is amended by adding at the end the fol-
16	lowing new paragraph:
17	"(19) Disclosure of return information
18	FOR PURPOSES OF CARRYING OUT MEDICARE CATA-
19	STROPHIC PRESCRIPTION DRUG PROGRAM.—
20	"(A) In General.—The Secretary may,
21	upon written request from the Secretary of
22	Health and Human Services under section
23	1860D-2(b)(4)(E)(i) of the Social Security Act,
24	disclose to officers and employees of the Depart-
25	ment of Health and Human Services with re-



1	spect to a specified taxpayer for the taxable year
2	specified by the Secretary of Health and Human
3	Services in such request—
4	"(i) the taxpayer identity information
5	with respect to such taxpayer, and
6	"(ii) the adjusted gross income of such
7	taxpayer for the taxable year (or, if less, the
8	income threshold limit specified in section
9	1860D-2(b)(4)(D)(ii) for the calendar year
10	specified by such Secretary in such request).
11	"(B) Specified taxpayer.—For purposes
12	of this paragraph, the term 'specified taxpayer'
13	means any taxpayer who—
14	"(i) is identified by the Secretary of
15	Health and Human Services in the request
16	referred to in subparagraph (A), and
17	"(ii) either—
18	"(I) has an adjusted gross income
19	for the taxable year referred to in sub-
20	paragraph (A) in excess of the income
21	threshold specified in section 1860D-
22	2(b)(4)(D)(ii) of such Act for the cal-
23	endar year referred to in such subpara-
24	graph, or



1	"(II) is identified by such Sec-
2	retary under subparagraph (A) as
3	being an individual who elected to use
4	more recent information under section
5	$1860D-2(b)(4)(D)(v) \ of \ such \ Act.$
6	"(C) Joint returns.—In the case of a
7	joint return, the Secretary shall, for purposes of
8	applying this paragraph, treat each spouse as a
9	separate taxpayer having an adjusted gross in-
10	come equal to one-half of the adjusted gross in-
11	come determined with respect to such return.
12	"(D) Restriction on use of disclosed
13	Information.—Return information disclosed
14	under subparagraph (A) may be used by officers
15	and employees of the Department of Health and
16	Human Services only for the purpose of admin-
17	istering the prescription drug benefit under title
18	XVIII of the Social Security Act. Such officers
19	and employees may disclose the annual out-of-
20	pocket threshold which applies to an individual
21	under such part to the entity that offers the plan
22	referred to in section $1860D-2(b)(4)(E)(ii)$ of
23	such Act in which such individual is enrolled.
24	Such sponsor may use such information only for

purposes of administering such benefit.".



(b) Confidentiality.—Paragraph (3) of section
6103(a) of such Code is amended by striking "or (16)" and
inserting "(16), or (19)".
(c) Procedures and Recordkeeping Related to
Disclosures.—Subsection (p)(4) of section 6103 of such
Code is amended by striking "any other person described
in subsection (l)(16) or (17)" each place it appears and in-
serting "any other person described in subsection (l)(16),
(17), or (19)".
(d) Unauthorized Disclosure.—Paragraph (2) of
section 7213(a) of such Code is amended by striking "or
(16)" and inserting "(16), or (19)".
(e) Unauthorized Inspection.—Subparagraph (B)
of section 7213A(a)(1) of such Code is amended by inserting
"or (19)" after "subsection (l)(18)".
SEC. 107. STATE PHARMACEUTICAL ASSISTANCE TRANSF
TION COMMISSION.
(a) Establishment.—
(1) In general.—There is established, as of the
first day of the third month beginning after the date
of the enactment of this Act, a State Pharmaceutical
Assistance Transition Commission (in this section re-
ferred to as the "Commission") to develop a proposal
for addressing the unique transitional issues facing

State pharmaceutical assistance programs, and pro-



1	gram participants, due to the implementation of the
2	$medicare\ prescription\ drug\ program\ under\ part\ D$ of
3	title XVIII of the Social Security Act.
4	(2) Definitions.—For purposes of this section:
5	(A) State pharmaceutical assistance
6	PROGRAM DEFINED.—The term "State pharma-
7	ceutical assistance program" means a program
8	(other than the medicaid program) operated by
9	a State (or under contract with a State) that
10	provides as of the date of the enactment of this
11	Act assistance to low-income medicare bene-
12	ficiaries for the purchase of prescription drugs.
13	(B) Program participant.—The term
14	"program participant" means a low-income
15	medicare beneficiary who is a participant in a
16	State pharmaceutical assistance program.
17	(b) Composition.—The Commission shall include the
18	following:
19	(1) A representative of each governor of each
20	State that the Secretary identifies as operating on a
21	statewide basis a State pharmaceutical assistance
22	program that provides for eligibility and benefits that
23	are comparable or more generous than the low-income
24	assistance eligibility and benefits offered under part

 $D\ of\ title\ XVIII\ of\ the\ Social\ Security\ Act.$ 

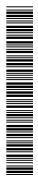


1	(2) Representatives from other States that the
2	Secretary identifies have in operation other State
3	pharmaceutical assistance programs, as appointed by
4	the Secretary.
5	(3) Representatives of organizations that have an
6	inherent interest in program participants or the pro-
7	gram itself, as appointed by the Secretary but not to
8	exceed the number of representatives under para-
9	graphs (1) and (2).
10	(4) Representatives of Medicare Advantage orga-
11	nizations and other private health insurance plans, as
12	appointed by the Secretary.
13	(5) The Secretary (or the Secretary's designee)
14	and such other members as the Secretary may specify
15	The Secretary shall designate a member to serve as chain
16	of the Commission and the Commission shall meet at the
17	call of the chair.
18	(c) Development of Proposal.—The Commission
19	shall develop the proposal described in subsection (a) in a
20	manner consistent with the following principles:
21	(1) Protection of the interests of program par-
22	ticipants in a manner that is the least disruptive to
23	such participants and that includes a single point of

contact for enrollment and processing of benefits.



1	(2) Protection of the financial and flexibility in-
2	terests of States so that States are not financially
3	worse off as a result of the enactment of this title.
4	(3) Principles of medicare modernization pro-
5	vided under title II of this Act.
6	(d) Report.—By not later than January 1, 2005, the
7	Commission shall submit to the President and the Congress
8	a report that contains a detailed proposal (including spe-
9	cific legislative or administrative recommendations, if any
10	and such other recommendations as the Commission deems
11	appropriate.
12	(e) Support.—The Secretary shall provide the Com-
13	mission with the administrative support services necessary
14	for the Commission to carry out its responsibilities under
15	this section.
16	(f) Termination.—The Commission shall terminate
17	30 days after the date of submission of the report under
18	subsection (d)



1	TITLE II—MEDICARE ENHANCED
2	FEE-FOR-SERVICE AND MEDI-
3	CARE ADVANTAGE PRO-
4	GRAMS; MEDICARE COMPETI-
5	TION
6	SEC. 200. MEDICARE MODERNIZATION AND REVITALIZA
7	TION.
8	This title provides for—
9	(1) establishment of the medicare enhanced fee
10	for-service (EFFS) program under which medicare
11	beneficiaries are provided access to a range of en
12	hanced fee-for-service (EFFS) plans that may use
13	preferred provider networks to offer an enhanced
14	range of benefits;
15	(2) establishment of a Medicare Advantage pro-
16	gram that offers improved managed care plans with
17	coordinated care; and
18	(3) competitive bidding, in the style of the Fed
19	eral Employees Health Benefits program (FEHBP)
20	among enhanced fee-for-service plans and Medicare
21	Advantage plans in order to promote greater effi
	2 2

ciency and responsiveness to medicare beneficiaries.



1	Subtitle A—Medicare Enhanced
2	Fee-for-Service Program
3	SEC. 201. ESTABLISHMENT OF ENHANCED FEE-FOR-SERV-
4	ICE (EFFS) PROGRAM UNDER MEDICARE.
5	(a) In General.—Title XVIII, as amended by section
6	101(a), is amended—
7	(1) by redesignating part E as part F; and
8	(2) by inserting after part D the following new
9	part:
10	"Part E—Enhanced Fee-for-Service Program
11	"OFFERING OF ENHANCED FEE-FOR-SERVICE PLANS
12	THROUGHOUT THE UNITED STATES
13	"Sec. 1860E-1. (a) Establishment of Program.—
14	"(1) In General.—The Administrator shall es-
15	tablish under this part beginning January 1, 2006,
16	an enhanced fee-for-service program under which en-
17	hanced fee-for-service plans (as defined in subsection
18	(b)) are offered to EFFS-eligible individuals (as so
19	defined) in EFFS regions throughout the United
20	States.
21	"(2) EFFS REGIONS.—For purposes of this part
22	the Administrator shall establish EFFS regions
23	throughout the United States by dividing the entire
24	United States into at least 10 such regions. Before es-

tablishing such regions, the Administrator shall con-



1	duct a market survey and analysis, including an ex-
2	amination of current insurance markets, to determine
3	how the regions should be established. The regions
4	shall be established in a manner to take into consider-
5	ation maximizing full access for all EFFS-eligible in-
6	dividuals, especially those residing in rural areas.
7	"(b) Definitions.—For purposes of this part:
8	"(1) EFFS organization.—The 'EFFS organi-
9	zation' means an entity that the Administrator cer-
10	tifies as meeting the requirements and standards ap-
11	plicable to such organization under this part.
12	"(2) Enhanced fee-for-service plan; effs
13	PLAN.—The terms 'enhanced fee-for-service plan' and
14	'EFFS plan' mean health benefits coverage offered
15	under a policy, contract, or plan by an EFFS organi-
16	zation pursuant to and in accordance with a contract
17	pursuant to section 1860E-4(c), but only if the plan
18	provides either fee-for-service coverage described in the
19	following subparagraph (A) or preferred provider cov-
20	erage described in the following subparagraph (B):
21	"(A) FEE-FOR-SERVICE COVERAGE.—The
22	plan—
23	"(i) reimburses hospitals, physicians,
24	and other providers at a rate determined by



1	the plan on a fee-for-service basis without
2	placing the provider at financial risk;
3	"(ii) does not vary such rates for such
4	a provider based on utilization relating to
5	such provider; and
6	"(iii) does not restrict the selection of
7	providers among those who are lawfully au-
8	thorized to provide the covered services and
9	agree to accept the terms and conditions of
10	payment established by the plan.
11	"(B) Preferred provider coverage.—
12	The plan—
13	"(i) has a network of providers that
14	have agreed to a contractually specified re-
15	imbursement for covered benefits with the
16	organization offering the plan; and
17	"(ii) provides for reimbursement for all
18	covered benefits regardless of whether such
19	benefits are provided within such network of
20	providers.
21	"(3) EFFS ELIGIBLE INDIVIDUAL.—The term
22	'EFFS eligible individual' means an eligible indi-
23	$vidual\ described\ in\ section\ 1851(a)(3).$
24	"(4) EFFS region.—The term 'EFFS region'
25	means a region established under subsection $(a)(2)$ .



1	"(c) Application of Certain Eligibility, Enroll-
2	MENT, ETC. REQUIREMENTS.—The provisions of section
3	1851 (other than subsection (h)(4)(A)) shall apply to EFFS
4	plans offered by an EFFS organization in an EFFS region,
5	including subsection (g) (relating to guaranteed issue and
6	renewal).
7	"OFFERING OF ENHANCED FEE-FOR-SERVICE (EFFS) PLANS
8	"Sec. 1860E-2. (a) Plan Requirements.—No
9	EFFS plan may be offered under this part in an EFFS
10	region unless the requirements of this part are met with
11	respect to the plan and EFFS organization offering the
12	plan.
13	"(b) Available to All EFFS Beneficiaries in the
14	Entire Region.—With respect to an EFFS plan offered
15	in an EFFS region—
16	"(1) In general.—The plan must be offered to
17	all EFFS-eligible individuals residing in the region.
18	"(2) Assuring access to services.—The plan
19	shall comply with the requirements of section
20	1852(d)(4).
21	"(c) Benefits.—
22	"(1) In general.—Each EFFS plan shall pro-
23	vide to members enrolled in the plan under this part
24	benefits, through providers and other persons that
25	meet the applicable requirements of this title and part



26

A of title XI—

1	"(A) for the items and services described in
2	$section \ 1852(a)(1);$
3	"(B) that are uniform for the plan for all
4	EFFS eligible individuals residing in the same
5	EFFS region;
6	"(C) that include a single deductible appli-
7	cable to benefits under parts A and B and in-
8	clude a catastrophic limit on out-of-pocket ex-
9	penditures for such covered benefits; and
10	"(D) that include benefits for prescription
11	drug coverage for each enrollee who elects under
12	part D to be provided qualified prescription
13	drug coverage through the plan.
14	"(2) Disapproval authority.—The Adminis-
15	trator shall not approve a plan of an EFFS organi-
16	zation if the Administrator determines (pursuant to
17	the last sentence of section $1852(b)(1)(A)$ ) that the
18	benefits are designed to substantially discourage en-
19	rollment by certain EFFS eligible individuals with
20	$the \ organization.$
21	"(d) Outpatient Prescription Drug Coverage.—
22	For rules concerning the offering of prescription drug cov-
23	erage under EFFS plans, see the amendment made by sec-
24	tion 102(b) of the Medicare Prescription Drug and Mod-
25	ernization Act of 2003.



1	"(e) Other Additional Provisions.—The provi-
2	sions of section 1852 (other than subsection $(a)(1)$ ) shall
3	apply under this part to EFFS plans. For the application
4	of chronic care improvement provisions, see the amendment
5	made by section 722(b).
6	"SUBMISSION OF BIDS; BENEFICIARY SAVINGS; PAYMENT
7	OF PLANS
8	"Sec. 1860E-3. (a) Submission of Bids.—
9	"(1) Requirement.—
10	"(A) EFFS MONTHLY BID AMOUNT.—For
11	each year (beginning with 2006), an EFFS orga-
12	nization shall submit to the Administrator an
13	EFFS monthly bid amount for each EFFS plan
14	offered in each region. Each such bid is referred
15	to in this section as the 'EFFS monthly bid
16	amount'.
17	"(B) FORM.—Such bid amounts shall be
18	submitted for each such plan and region in a
19	form and manner and time specified by the Ad-
20	ministrator, and shall include information de-
21	scribed in paragraph (3)(A).
22	"(2) Uniform bid amounts.—Each EFFS
23	monthly bid amount submitted under paragraph (1)
24	by an EFFS organization under this part for an
25	EFFS plan in an EFFS region may not vary among



1	EFFS eligible individuals residing in the EFFS re-
2	gion involved.
3	"(3) Submission of bid amount information
4	BY EFFS ORGANIZATIONS.—
5	"(A) Information to be submitted.—
6	The information described in this subparagraph
7	is as follows:
8	"(i) The EFFS monthly bid amount
9	for provision of all items and services under
10	this part, which amount shall be based on
11	average costs for a typical beneficiary resid-
12	ing in the region, and the actuarial basis
13	for determining such amount.
14	"(ii) The proportions of such bid
15	amount that are attributable to—
16	"(I) the provision of statutory
17	non-drug benefits (such portion re-
18	ferred to in this part as the
19	'unadjusted EFFS statutory non-drug
20	monthly bid amount');
21	"(II) the provision of statutory
22	prescription drug benefits; and
23	"(III) the provision of non-statu-
24	tory benefits;



1	and the actuarial basis for determining
2	such proportions.
3	"(iii) Such additional information as
4	the Administrator may require to verify the
5	actuarial bases described in clauses (i) and
6	(ii).
7	"(B) Statutory benefits defined.—For
8	purposes of this part:
9	"(i) The term 'statutory non-drug ben-
10	efits' means benefits under section
11	1852(a)(1).
12	"(ii) The term 'statutory prescription
13	drug benefits' means benefits under part D.
14	"(iii) The term 'statutory benefits'
15	means statutory prescription drug benefits
16	and statutory non-drug benefits.
17	"(C) Acceptance and negotiation of
18	BID AMOUNTS.—The Administrator has the au-
19	thority to negotiate regarding monthly bid
20	amounts submitted under subparagraph (A)
21	(and the proportion described in subparagraph
22	(A)(ii)), and for such purpose, the Administrator
23	has negotiation authority that the Director of the
24	Office of Personnel Management has with respect
25	to health benefits plans under chapter 89 of title



1	5, United States Code. The Administrator may
2	reject such a bid amount or proportion if the Ad-
3	ministrator determines that such amount or pro-
4	portion is not supported by the actuarial bases
5	provided under subparagraph (A).
6	"(D) Contract authority.—The Admin-
7	istrator may, taking into account the unadjusted
8	EFFS statutory non-drug monthly bid amounts
9	accepted under subparagraph (C), enter into con-
10	tracts for the offering of up to 3 EFFS plans in
11	any region.
12	"(b) Provision of Beneficiary Savings for Cer-
13	TAIN PLANS.—
14	"(1) Beneficiary rebate rule.—
15	"(A) Requirement.—The EFFS plan shall
16	provide to the enrollee a monthly rebate equal to
17	75 percent of the average per capita savings (if
18	any) described in paragraph (2) applicable to
19	the plan and year involved.
20	"(B) Form of Rebate.—A rebate required
21	under this paragraph shall be provided—
22	"(i) through the crediting of the
23	amount of the rebate towards the EFFS
24	monthly prescription drug beneficiary pre-
25	mium (as defined in section 1860E-



1	4(a)(3)(B)) and the EFFS monthly supple-
2	mental beneficiary premium (as defined in
3	section $1860E-4(a)(3)(C)$ ;
4	"(ii) through a direct monthly pay-
5	ment (through electronic funds transfer or
6	$otherwise);\ or$
7	"(iii) through other means approved by
8	$the\ Medicare\ Benefits\ Administrator,$
9	or any combination thereof.
10	"(2) Computation of Average per capita
11	MONTHLY SAVINGS.—For purposes of paragraph
12	(1)(A), the average per capita monthly savings re-
13	ferred to in such paragraph for an EFFS plan and
14	year is computed as follows:
15	"(A) Determination of region-wide Av-
16	ERAGE RISK ADJUSTMENT.—
17	"(i) In General.—The Medicare Ben-
18	efits Administrator shall determine, at the
19	same time rates are promulgated under sec-
20	$tion \ 1853(b)(1)$ (beginning with 2006), for
21	each EFFS region the average of the risk
22	adjustment factors described in subsection
23	(c)(3) to be applied to enrollees under this
24	part in that region. In the case of an EFFS
25	region in which an EFFS plan was offered



1	in the previous year, the Administrator
2	may compute such average based upon risk
3	adjustment factors applied under subsection
4	(c)(3) in that region in a previous year.
5	"(ii) Treatment of New Regions.—
6	In the case of a region in which no EFFS
7	plan was offered in the previous year, the
8	Administrator shall estimate such average.
9	In making such estimate, the Administrator
10	may use average risk adjustment factors ap-
11	plied to comparable EFFS regions or ap-
12	plied on a national basis.
13	"(B) Determination of risk adjusted
14	BENCHMARK AND RISK-ADJUSTED BID.—For
15	each EFFS plan offered in an EFFS region, the
16	Administrator shall—
17	"(i) adjust the EFFS region-specific
18	non-drug monthly benchmark amount (as
19	defined in paragraph (3)) by the applicable
20	average risk adjustment factor computed
21	under subparagraph (A); and
22	"(ii) adjust the unadjusted EFFS stat-
23	utory non-drug monthly bid amount by
24	such applicable average risk adjustment fac-
25	tor.



1	"(C) Determination of average per
2	CAPITA MONTHLY SAVINGS.—The average per
3	capita monthly savings described in this sub-
4	paragraph is equal to the amount (if any) by
5	which—
6	"(i) the risk-adjusted benchmark
7	amount computed under subparagraph
8	$(B)(i),\ exceeds$
9	"(ii) the risk-adjusted bid computed
10	$under\ subparagraph\ (B)(ii).$
11	"(3) Computation of EFFS region-specific
12	NON-DRUG MONTHLY BENCHMARK AMOUNT.—For pur-
13	poses of this part, the term 'EFFS region-specific
14	non-drug monthly benchmark amount' means, with
15	respect to an EFFS region for a month in a year, an
16	amount equal to $1/12$ of the average (weighted by num-
17	ber of EFFS eligible individuals in each payment
18	area described in section 1853(d)) of the annual capi-
19	tation rate as calculated under section $1853(c)(1)$ for
20	that area.
21	"(c) Payment of Plans Based on Bid Amounts.—
22	"(1) Non-drug benefits.—Under a contract
23	under section $1860E-4(c)$ and subject to section
24	1853(g) (as made applicable under subsection (d)),
25	the Administrator shall make monthly payments



1	under this subsection in advance to each EFFS orga-
2	nization, with respect to coverage of an individual
3	under this part in an EFFS region for a month, in
4	an amount determined as follows:
5	"(A) Plans with bids below bench-
6	MARK.—In the case of a plan for which there are
7	average per capita monthly savings described in
8	subsection $(b)(2)(C)$ , the payment under this sub-
9	section is equal to the unadjusted EFFS statu-
10	tory non-drug monthly bid amount, adjusted
11	under paragraphs (3) and (4), plus the amount
12	of the monthly rebate computed under subsection
13	(b)(1)(A) for that plan and year.
14	"(B) Plans with bids at or above
15	BENCHMARK.—In the case of a plan for which
16	there are no average per capita monthly savings
17	described in subsection $(b)(2)(C)$ , the payment
18	amount under this subsection is equal to the
19	EFFS region-specific non-drug monthly bench-
20	mark amount, adjusted under paragraphs (3)
21	and $(4)$ .
22	"(2) For federal drug subsidies.—In the
23	case in which an enrollee who elects under part D to

be provided qualified prescription drug coverage



1	through the plan, the EFFS organization offering
2	such plan also is entitled—
3	"(A) to direct subsidy payment under sec-
4	tion 1860D-8(a)(1);
5	"(B) to reinsurance subsidy payments
6	under section $1860D-8(a)(2)$ ; and
7	"(C) to reimbursement for premium and
8	cost-sharing reductions for low-income individ-
9	uals under section $1860D-7(c)(3)$ .
10	"(3) Demographic risk adjustment, includ-
11	ING ADJUSTMENT FOR HEALTH STATUS.—The Admin-
12	istrator shall adjust under paragraph (1)(A) the
13	unadjusted EFFS statutory non-drug monthly bid
14	amount and under paragraph (1)(B) the EFFS re-
15	gion-specific non-drug monthly benchmark amount
16	for such risk factors as age, disability status, gender,
17	institutional status, and such other factors as the Ad-
18	ministrator determines to be appropriate, including
19	adjustment for health status under section 1853(a)(3)
20	(as applied under subsection (d)), so as to ensure ac-
21	tuarial equivalence. The Administrator may add to,
22	modify, or substitute for such adjustment factors if
23	such changes will improve the determination of actu-
24	arial equivalence.



1	"(4) ADJUSTMENT FOR INTRA-REGIONAL GEO-
2	GRAPHIC VARIATIONS.—The Administrator shall also
3	adjust such amounts in a manner to take into ac-
4	count variations in payments rates under part C
5	among the different payment areas under such part
6	included in each EFFS region.
7	"(d) Application of Additional Payment
8	Rules.—The provisions of section 1853 (other than sub-
9	sections (a)(1)(A), (d), and (e)) shall apply to an EFFS
10	plan under this part, except as otherwise provided in this
11	section.
12	"PREMIUMS; ORGANIZATIONAL AND FINANCIAL REQUIRE-
13	MENTS; ESTABLISHMENT OF STANDARDS; CONTRACTS
14	WITH EFFS ORGANIZATIONS
15	"Sec. 1860E-4. (a) Premiums.—
16	"(1) In general.—The provisions of section
17	1854 (other than subsections (a)(6)(C) and (h)), in-
18	cluding subsection (b)(5) relating to the consolidation
19	of drug and non-drug beneficiary premiums and sub-
20	section (c) relating to uniform bids and premiums,
21	shall apply to an EFFS plan under this part, subject
22	to paragraph (2).
23	"(2) Cross-Walk.—In applying paragraph (1),
24	any reference in section $1854(b)(1)(A)$ or $1854(d)$
25	to—



1	"(A) a Medicare Advantage monthly basic
2	beneficiary premium is deemed a reference to the
3	EFFS monthly basic beneficiary premium (as
4	defined in paragraph $(3)(A)$ ;
5	"(B) a Medicare Advantage monthly pre-
6	scription drug beneficiary premium is deemed a
7	reference to the EFFS monthly prescription drug
8	beneficiary premium (as defined in paragraph
9	(3)(B); and
10	"(C) a Medicare Advantage monthly supple-
11	mental beneficiary premium is deemed a ref-
12	erence to the EFFS monthly supplemental bene-
13	ficiary premium (as defined in paragraph
14	(3)(C)).
15	"(3) Definitions.—For purposes of this part:
16	"(A) EFFS MONTHLY BASIC BENEFICIARY
17	PREMIUM.—The term 'EFFS monthly basic ben-
18	eficiary premium' means, with respect to an
19	EFFS plan—
20	"(i) described in section 1860E-
21	3(c)(1)(A) (relating to plans providing re-
22	bates), zero; or
23	"(ii) described in section 1860E-
24	3(c)(1)(B), the amount (if any) by which
25	the unadjusted EFFS statutory non-drug



1	monthly bid amount exceeds the EFFS re-
2	gion-specific non-drug monthly benchmark
3	amount (as defined in section 1860E-
4	3(b)(3)).
5	"(B) EFFS MONTHLY PRESCRIPTION DRUG
6	BENEFICIARY PREMIUM.—The term 'EFFS
7	monthly prescription drug beneficiary premium
8	means, with respect to an EFFS plan, the por-
9	tion of the aggregate monthly bid amount sub-
10	mitted under clause (i) of section 1860E-
11	3(a)(3)(A) for the year that is attributable under
12	such section to the provision of statutory pre-
13	scription drug benefits.
14	"(C) EFFS MONTHLY SUPPLEMENTAL BEN-
15	EFICIARY PREMIUM.—The term 'EFFS monthly
16	supplemental beneficiary premium' means, with
17	respect to an EFFS plan, the portion of the ag-
18	gregate monthly bid amount submitted under
19	clause (i) of section $1860E-3(a)(3)(A)$ for the
20	year that is attributable under such section to
21	the provision of nonstatutory benefits.
22	"(b) Organizational and Financial Require-
23	MENTS.—The provisions of section 1855 shall apply to an
24	EFFS plan offered by an EFFS organization under this
25	part.



- 1 "(c) Contracts with EFFS Organizations.—The
- 2 provisions of section 1857 shall apply to an EFFS plan
- 3 offered by an EFFS organization under this part, except
- 4 that any reference in such section to part C is deemed a
- 5 reference to this part.".
- 6 (b) Prohibition on Coverage Under Medigap
- 7 Plans of Deductible Imposed Under EFFS Plans.—
- 8 Section 1882 (42 U.S.C. 1395ss), as amended by section
- 9 104(a), is amended by adding at the end the following new
- 10 subsection:
- 11 "(w) Prohibition on Coverage of Deductible
- 12 and Certain Cost-Sharing Imposed Under EFFS
- 13 Plans.—Notwithstanding any other provision of law, no
- 14 medicare supplemental policy (other than the 2 benefit
- 15 packages described in subsection (v)(3)) may provide for
- 16 coverage of the single deductible or more than 50 percent
- 17 of other cost-sharing imposed under an EFFS plan under
- 18 part E.".
- 19 (c) Conforming Provisions.—Section 1882 of the
- 20 Social Security Act (42 U.S.C. 1395ss) shall be adminis-
- 21 tered as if any reference to a Medicare+Choice organization
- 22 offering a Medicare+Choice plan under part C of title
- 23 XVIII of such Act were a reference both to a Medicare Ad-
- 24 vantage organization offering a Medicare Advantage plan



1	under such part and an EFFS organization offering an
2	EFFS plan under part E of such title.
3	Subtitle B—Medicare Advantage
4	Program
5	CHAPTER 1—IMPLEMENTATION OF
6	PROGRAM
7	SEC. 211. IMPLEMENTATION OF MEDICARE ADVANTAGE
8	PROGRAM.
9	(a) In General.—There is hereby established the
10	Medicare Advantage program. The Medicare Advantage
11	program shall consist of the program under part C of title
12	XVIII of the Social Security Act, as amended by this title.
13	(b) References.—Any reference to the program
14	under part C of title XVIII of the Social Security Act shall
15	be deemed a reference to the Medicare Advantage program
16	and, with respect to such part, any reference to
17	"Medicare+Choice" is deemed a reference to "Medicare Ad-
18	vantage".
19	SEC. 212. MEDICARE ADVANTAGE IMPROVEMENTS.
20	(a) Equalizing Payments With Fee-For-Serv-
21	ICE.—
22	(1) In General.—Section 1853(c)(1) (42 U.S.C.
23	1395w-23(c)(1)) is amended by adding at the end the
24	followina:



1	"(D) Based on 100 percent of fee-for-
2	SERVICE COSTS.—
3	"(i) In General.—For 2004, the ad-
4	justed average per capita cost for the year
5	involved, determined under section
6	1876(a)(4) for the Medicare Advantage pay-
7	ment area for services covered under parts
8	A and B for individuals entitled to benefits
9	under part A and enrolled under part B
10	who are not enrolled in a Medicare Advan-
11	tage under this part for the year, but ad-
12	justed to exclude costs attributable to pay-
13	ments under section 1886(h).
14	"(ii) Inclusion of costs of va and
15	DOD MILITARY FACILITY SERVICES TO MEDI-
16	Care-eligible beneficiaries.—In deter-
17	mining the adjusted average per capita cost
18	under clause (i) for a year, such cost shall
19	be adjusted to include the Secretary's esti-
20	mate, on a per capita basis, of the amount
21	of additional payments that would have
22	been made in the area involved under this
23	title if individuals entitled to benefits under
24	this title had not received services from fa-



1	cilities of the Department of Veterans Af-
2	fairs or the Department of Defense.".
3	(2) Conforming amendment.—Such section is
4	further amended, in the matter before subparagraph
5	(A), by striking "or (C)" and inserting "(C), or (D)".
6	(b) Change in Budget Neutrality for Blend.—
7	Section 1853(c) (42 U.S.C. 1395w-23(c)) is amended—
8	(1) in paragraph (1)(A), by inserting "(for a
9	year other than 2004)" after "multiplied"; and
10	(2) in paragraph (5), by inserting "(other than
11	2004)" after "for each year".
12	(c) Increasing Minimum Percentage Increase to
13	National Growth Rate.—
14	(1) In General.—Section 1853(c)(1) (42 U.S.C.
15	1395w-23(c)(1)) is amended—
16	(A) in subparagraph $(B)(iv)$ , by striking
17	"and each succeeding year" and inserting ",
18	2003, and 2004";
19	(B) in subparagraph $(C)(iv)$ , by striking
20	"and each succeeding year" and inserting "and
21	2003"; and
22	(C) by adding at the end of subparagraph
23	(C) the following new clause:
24	"(v) For 2004 and each succeeding
25	year, the greater of—



1	"(I) 102 percent of the annual
2	Medicare Advantage capitation rate
3	under this paragraph for the area for
4	the previous year; or
5	"(II) the annual Medicare Advan-
6	tage capitation rate under this para-
7	graph for the area for the previous
8	year increased by the national per cap-
9	ita Medicare Advantage growth per-
10	centage, described in paragraph (6) for
11	that succeeding year, but not taking
12	into account any adjustment under
13	paragraph (6)(C) for a year before
14	2004.".
15	(2) Conforming Amendment.—Section
16	1853(c)(6)(C) (42 U.S.C. $1395w-23(c)(6)(C)$ ) is
17	amended by inserting before the period at the end the
18	following: ", except that for purposes of paragraph
19	(1)(C)(v)(II), no such adjustment shall be made for a
20	year before 2004".
21	(d) Inclusion of Costs of DOD and VA Military
22	FACILITY SERVICES TO MEDICARE-ELIGIBLE BENE-
23	FICIARIES IN CALCULATION OF MEDICARE+CHOICE PAY-
24	MENT RATES.—Section 1853(c)(3) (42 U.S.C. 1395w-
25	23(c)(3)) is amended—



1	(1) in subparagraph (A), by striking "subpara-
2	graph (B)" and inserting "subparagraphs (B) and
3	(E)", and
4	(2) by adding at the end the following new sub-
5	paragraph:
6	"(E) Inclusion of costs of dod and va
7	MILITARY FACILITY SERVICES TO MEDICARE-ELI-
8	GIBLE BENEFICIARIES.—In determining the
9	$area-specific \ Medicare+Choice \ capitation \ rate$
10	under subparagraph (A) for a year (beginning
11	with 2004), the annual per capita rate of pay-
12	ment for 1997 determined under section
13	1876(a)(1)(C) shall be adjusted to include in the
14	rate the Secretary's estimate, on a per capita
15	basis, of the amount of additional payments that
16	would have been made in the area involved
17	under this title if individuals entitled to benefits
18	under this title had not received services from fa-
19	cilities of the Department of Defense or the De-
20	partment of Veterans Affairs.".
21	(e) Extending Special Rule for Certain Inpa-
22	TIENT HOSPITAL STAYS TO REHABILITATION HOS-
23	PITALS.—
24	(1) In General.—Section 1853(g) (42 U.S.C.
25	1395w-23(g)) is amended—



1	(A) by inserting "or from a rehabilitation
2	facility (as defined in section $1886(j)(1)(A)$ )"
3	after " $1886(d)(1)(B)$ "; and
4	(B) in paragraph (2)(B), by inserting "or
5	section 1886(j), as the case may be," after
6	"1886(d)".
7	(2) Effective date.—The amendments made
8	by paragraph (1) shall apply to contract years begin-
9	ning on or after January 1, 2004.
10	(f) MedPAC Study of AAPCC.—
11	(1) Study.—The Medicare Payment Advisory
12	Commission shall conduct a study that assesses the
13	method used for determining the adjusted average per
14	capita cost (AAPCC) under section 1876(a)(4) of the
15	Social Security Act (42 U.S.C. 1395mm(a)(4)) as ap-
16	plied under section $1853(c)(1)(A)$ of such $Act$ (as
17	amended by subsection (a)). Such study shall include
18	an examination of—
19	(A) the bases for variation in such costs be-
20	tween different areas, including differences in
21	input prices, utilization, and practice patterns;
22	(B) the appropriate geographic area for
23	payment under the Medicare Advantage program
24	under part C of title XVIII of such Act; and



1	(C) the accuracy of risk adjustment methods
2	in reflecting differences in costs of providing care
3	to different groups of beneficiaries served under
4	such program.
5	(2) Report.—Not later than 18 months after the
6	date of the enactment of this Act, the Commission
7	shall submit to Congress a report on the study con-
8	ducted under paragraph (1).
9	(g) Report on Impact of Increased Financial As-
10	SISTANCE TO MEDICARE ADVANTAGE PLANS.—Not later
11	than July 1, 2006, the Medicare Benefits Administrator
12	shall submit to Congress a report that describes the impact
13	of additional financing provided under this Act and other
14	Acts (including the Medicare, Medicaid, and SCHIP Bal-
15	anced Budget Refinement Act of 1999 and BIPA) on the
16	availability of Medicare Advantage plans in different areas
17	and its impact on lowering premiums and increasing bene-
18	fits under such plans.
19	CHAPTER 2—IMPLEMENTATION OF
20	COMPETITION PROGRAM
21	SEC. 221. COMPETITION PROGRAM BEGINNING IN 2006.
22	(a) Submission of EFFS-Like Bidding Informa-
23	TION BEGINNING IN 2006.—Section 1854 (42 U.S.C.
24	1395w-24) is amended—



1	(1) by amending the section heading to read as
2	follows:
3	"PREMIUMS AND BID AMOUNT";
4	(2) in subsection $(a)(1)(A)$ —
5	(A) by striking "(A)" and inserting "(A)(i)
6	if the following year is before 2006,"; and
7	(B) by inserting before the semicolon at the
8	end the following: "or (ii) if the following year
9	is 2006 or later, the information described in
10	paragraph (3) or (6)(A) for the type of plan in-
11	volved"; and
12	(3) by adding at the end of subsection (a) the fol-
13	lowing:
14	"(6) Submission of bid amounts by medicare
15	ADVANTAGE ORGANIZATIONS.—
16	"(A) Information to be submitted.—
17	The information described in this subparagraph
18	is as follows:
19	"(i) The monthly aggregate bid
20	amount for provision of all items and serv-
21	ices under this part, which amount shall be
22	based on average costs for a typical bene-
23	ficiary residing in the area, and the actu-
24	arial basis for determining such amount.
25	"(ii) The proportions of such bid
26	amount that are attributable to—



1	"(I) the provision of statutory
2	non-drug benefits (such portion re-
3	ferred to in this part as the
4	'unadjusted Medicare Advantage statu-
5	tory non-drug monthly bid amount');
6	"(II) the provision of statutory
7	prescription drug benefits; and
8	"(III) the provision of non-statu-
9	tory benefits;
10	and the actuarial basis for determining
11	such proportions.
12	"(iii) Such additional information as
13	the Administrator may require to verify the
14	actuarial bases described in clauses (i) and
15	(ii).
16	"(B) Statutory benefits defined.—For
17	purposes of this part:
18	"(i) The term 'statutory non-drug ben-
19	efits' means benefits under section
20	1852(a)(1).
21	"(ii) The term 'statutory prescription
22	drug benefits' means benefits under part D.
23	"(iii) The term 'statutory benefits'
24	means statutory prescription drug benefits
25	and statutory non-drug benefits.



1	"(C) Acceptance and negotiation of
2	BID AMOUNTS.—
3	"(i) In general.—Subject to clause
4	(ii)—
5	"(I) the Administrator has the au-
6	thority to negotiate regarding monthly
7	bid amounts submitted under subpara-
8	graph (A) (and the proportion de-
9	scribed in subparagraph (A)(ii)), and
10	for such purpose and subject to such
11	clause, the Administrator has negotia-
12	tion authority that the Director of the
13	Office of Personnel Management has
14	with respect to health benefits plans
15	under chapter 89 of title 5, United
16	States Code; and
17	"(II) the Administrator may re-
18	ject such a bid amount or proportion if
19	the Administrator determines that such
20	amount or proportion is not supported
21	by the actuarial bases provided under
22	subparagraph (A).
23	"(ii) Exception.—In the case of a
24	plan described in section $1851(a)(2)(C)$ , the
25	provisions of clause (i) shall not apply and



1	the provisions of paragraph $(5)(B)$ , prohib-
2	iting the review, approval, or disapproval of
3	amounts described in such paragraph, shall
4	apply to the negotiation and rejection of the
5	monthly bid amounts and proportion re-
6	ferred to in subparagraph (A).".
7	(b) Providing for Beneficiary Savings for Cer-
8	TAIN PLANS.—
9	(1) In general.—Section 1854(b) (42 U.S.C.
10	1395w-24(b)) is amended—
11	(A) by adding at the end of paragraph (1)
12	the following new subparagraph:
13	"(C) Beneficiary rebate rule.—
14	"(i) Requirement.—The Medicare
15	Advantage plan shall provide to the enrollee
16	a monthly rebate equal to 75 percent of the
17	average per capita savings (if any) de-
18	scribed in paragraph (3) applicable to the
19	plan and year involved.
20	"(iii) Form of rebate.—A rebate re-
21	quired under this subparagraph shall be
22	provided—
23	"(I) through the crediting of the
24	amount of the rebate towards the Medi-
25	care Advantage monthly supple-



1	mentary beneficiary premium or the
2	premium imposed for prescription
3	$drug\ coverage\ under\ part\ D;$
4	"(II) through a direct monthly
5	payment (through electronic funds
6	transfer or otherwise); or
7	"(III) through other means ap-
8	proved by the Medicare Benefits Ad-
9	ministrator,
10	or any combination thereof."; and
11	(B) by adding at the end the following new
12	paragraphs:
13	"(3) Computation of Average per capita
14	MONTHLY SAVINGS.—For purposes of paragraph
15	(1)(C)(i), the average per capita monthly savings re-
16	ferred to in such paragraph for a Medicare Advantage
17	plan and year is computed as follows:
18	"(A) Determination of state-wide av-
19	ERAGE RISK ADJUSTMENT.—
20	"(i) In General.—The Medicare Ben-
21	efits Administrator shall determine, at the
22	same time rates are promulgated under sec-
23	tion 1853(b)(1) (beginning with 2006), for
24	each State the average of the risk adjust-
25	ment factors to be applied under section



1	1853(a)(1)(A) to payment for enrollees in
2	that State. In the case of a State in which
3	a Medicare Advantage plan was offered in
4	the previous year, the Administrator may
5	compute such average based upon risk ad-
6	justment factors applied in that State in a
7	previous year.
8	"(ii) Treatment of New States.—In
9	the case of a State in which no Medicare
10	Advantage plan was offered in the previous
11	year, the Administrator shall estimate such
12	average. In making such estimate, the Ad-
13	ministrator may use average risk adjust-
14	ment factors applied to comparable States
15	or applied on a national basis.
16	"(B) Determination of risk adjusted
17	BENCHMARK AND RISK-ADJUSTED BID.—For
18	each Medicare Advantage plan offered in a State,
19	$the \ Administrator \ shall$ —
20	"(i) adjust the Medicare Advantage
21	area-specific non-drug monthly benchmark
22	amount (as defined in subsection (j)) by the
23	applicable average risk adjustment factor
24	computed under subparagraph (A); and



1	"(ii) adjust the unadjusted Medicare
2	Advantage statutory non-drug monthly bid
3	amount by such applicable average risk ad-
4	justment factor.
5	"(C) Determination of Average per
6	CAPITA MONTHLY SAVINGS.—The average per
7	capita monthly savings described in this sub-
8	paragraph is equal to the amount (if any) by
9	which—
10	"(i) the risk-adjusted benchmark
11	amount computed under subparagraph
12	$(B)(i),\ exceeds$
13	"(ii) the risk-adjusted bid computed
14	$under\ subparagraph\ (B)(ii).$
15	"(D) Authority to determine risk ad-
16	JUSTMENT FOR AREAS OTHER THAN STATES.—
17	The Administrator may provide for the deter-
18	mination and application of risk adjustment fac-
19	tors under this paragraph on the basis of areas
20	other than States.
21	"(4) Beneficiary's option of payment
22	THROUGH WITHHOLDING FROM SOCIAL SECURITY
23	PAYMENT OR USE OF ELECTRONIC FUNDS TRANSFER
24	MECHANISM.—In accordance with regulations, a
25	Medicare Advantage organization shall permit each



	110
1	enrollee, at the enrollee's option, to make payment of
2	premiums under this part to the organization indi-
3	rectly through withholding from benefit payments in
4	the manner provided under section 1840 with respect
5	to monthly premiums under section 1839 or through
6	an electronic funds transfer mechanism (such as auto-
7	matic charges of an account at a financial institution
8	or a credit or debit card account) or otherwise. All
9	premium payments that are withheld under this
10	paragraph that are credited to the Federal Supple-
11	mentary Medical Insurance Drug Trust Fund shall be
12	paid to the Medicare Advantage organization in-
13	volved.".
14	(2) Provision of single consolidated pre-
15	MIUM.—Section 1854(b) (42 U.S.C. 1395w-24(b)), as
16	amended by paragraph (1), is further amended by
17	adding at the end the following new paragraph:
18	"(5) Single consolidated premium.—In the
19	case of an enrollee in a Medicare Advantage plan who



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1	prescription drug coverage under part D provided
2	through the plan.".
3	(3) Computation of medicare advantage
4	AREA-SPECIFIC NON-DRUG BENCHMARK.—Section
5	1853 (42 U.S.C. 1395w-23) is amended by adding at
6	the end the following new subsection:
7	"(j) Computation of Medicare Advantage Area-
8	Specific Non-Drug Monthly Benchmark Amount.—
9	For purposes of this part, the term 'Medicare Advantage
10	area-specific non-drug monthly benchmark amount' means,
11	with respect to a Medicare Advantage payment area for a
12	month in a year, an amount equal to 1/12 of the annual
13	Medicare Advantage capitation rate under section
14	1853(c)(1) for the area for the year.".
15	(c) Payment of Plans Based on Bid Amounts.—
16	(1) In General.—Section $1853(a)(1)(A)$ (42)
17	U.S.C. 1395w-23) is amended by striking "in an
18	amount" and all that follows and inserting the fol-
19	lowing: "in an amount determined as follows:
20	"(i) Payment before 2006.—For
21	years before 2006, the payment amount
22	shall be equal to $1/12$ of the annual Medicare
23	Advantage capitation rate (as calculated
24	under subsection $(c)(1)$ ) with respect to that
25	individual for that area, reduced by the



1	amount of any reduction elected under sec-
2	tion $1854(f)(1)(E)$ and adjusted under
3	clause (iv).
4	"(ii) Payment for statutory non-
5	DRUG BENEFITS BEGINNING WITH 2006.—
6	For years beginning with 2006—
7	"(I) Plans with bids below
8	BENCHMARK.—In the case of a plan
9	for which there are average per capita
10	monthly savings described in section
11	1854(b)(3)(C), the payment under this
12	subsection is equal to the unadjusted
13	Medicare Advantage statutory non-
14	drug monthly bid amount, adjusted
15	under clause (iv), plus the amount of
16	the monthly rebate computed under
17	section $1854(b)(1)(C)(i)$ for that plan
18	and year.
19	"(II) Plans with bids at or
20	ABOVE BENCHMARK.—In the case of a
21	plan for which there are no average
22	per capita monthly savings described
23	in section $1854(b)(3)(C)$ , the payment
24	amount under this subsection is equal
25	to the Medicare Advantage area-spe-



1	cific non-drug monthly benchmark
2	amount, adjusted under clause (iv).
3	"(iii) For federal drug sub-
4	SIDIES.—In the case in which an enrollee
5	who elects under part D to be provided
6	qualified prescription drug coverage through
7	the plan, the Medicare Advantage organiza-
8	tion offering such plan also is entitled—
9	"(I) to direct subsidy payment
10	$under\ section\ 1860D-8(a)(1);$
11	"(II) to reinsurance subsidy pay-
12	ments $under$ $section$ $1860D-8(a)(2);$
13	and
14	"(III) to reimbursement for pre-
15	mium and cost-sharing reductions for
16	low-income individuals under section
17	1860D-7(c)(3).
18	"(iv) Demographic adjustment, in-
19	CLUDING ADJUSTMENT FOR HEALTH STA-
20	TUS.—The Administrator shall adjust the
21	payment amount under clause (i), the
22	unadjusted Medicare Advantage statutory
23	non-drug monthly bid amount under clause
24	(ii)(I), and the Medicare Advantage area-
25	specific non-drug monthly benchmark



1	amount under clause (ii)(II) for such risk
2	factors as age, disability status, gender, in-
3	stitutional status, and such other factors as
4	the Administrator determines to be appro-
5	priate, including adjustment for health sta-
6	tus under paragraph (3), so as to ensure ac-
7	tuarial equivalence. The Administrator may
8	add to, modify, or substitute for such ad-
9	justment factors if such changes will im-
10	prove the determination of actuarial equiva-
11	lence.".
12	(d) Conforming Amendments.—
13	(1) Protection against beneficiary selec-
14	TION.—Section 1852(b)(1)(A) (42 U.S.C. 1395w-
15	22(b)(1)(A)) is amended by adding at the end the fol-
16	lowing: "The Administrator shall not approve a plan
17	of an organization if the Administrator determines
18	that the benefits are designed to substantially discour-
19	age enrollment by certain Medicare Advantage eligible
20	individuals with the organization.".
21	(2) Conforming amendment to premium ter-
22	MINOLOGY.—Section 1854(b)(2) (42 U.S.C. 1395w-
23	24(b)(2)) is amended by redesignating subparagraph
24	(C) as subparagraph (D) and by striking subpara-

graphs (A) and (B) and inserting the following:



1	"(A) Medicare advantage monthly
2	Basic beneficiary premium.—The term 'Medi-
3	care Advantage monthly basic beneficiary pre-
4	mium' means, with respect to a Medicare Advan-
5	tage plan—
6	"(i) described in section
7	1853(a)(1)(A)(ii)(I) (relating to plans pro-
8	viding rebates), zero; or
9	"(ii) described in section
10	1853(a)(1)(A)(ii)(II), the amount (if any)
11	by which the unadjusted Medicare Advan-
12	tage statutory non-drug monthly bid
13	amount exceeds the Medicare Advantage
14	area-specific non-drug monthly benchmark
15	amount.
16	"(B) Medicare advantage monthly pre-
17	SCRIPTION DRUG BENEFICIARY PREMIUM.—The
18	term 'Medicare Advantage monthly prescription
19	drug beneficiary premium' means, with respect
20	to a Medicare Advantage plan, that portion of
21	the bid amount submitted under clause (i) of
22	subsection (a)(6)(A) for the year that is attrib-
23	utable under such section to the provision of stat-
24	utory prescription drug benefits.



1	"(C) Medicare advantage monthly sup-
2	PLEMENTAL BENEFICIARY PREMIUM.—The term
3	'Medicare Advantage monthly supplemental ben-
4	eficiary premium' means, with respect to a
5	Medicare Advantage plan, the portion of the ag-
6	gregate monthly bid amount submitted under
7	clause (i) of subsection $(a)(6)(A)$ for the year
8	that is attributable under such section to the pro-
9	vision of nonstatutory benefits.".
10	(3) Requirement for uniform premium and
11	BID AMOUNTS.—Section 1854(c) (42 U.S.C. 1395w-
12	24(c)) is amended to read as follows:
13	"(c) Uniform Premium and Bid Amounts.—The
14	Medicare Advantage monthly bid amount submitted under
15	subsection (a)(6), the Medicare Advantage monthly basic,
16	prescription drug, and supplemental beneficiary premiums,
17	and the Medicare Advantage monthly MSA premium
18	charged under subsection (b) of a Medicare Advantage orga-
19	nization under this part may not vary among individuals
20	enrolled in the plan.".
21	(4) Permitting beneficiary rebates.—
22	(A) Section $1851(h)(4)(A)$ (42 U.S.C.
23	1395w-21(h)(4)(A)) is amended by inserting
24	"except as provided under section 1854(b)(1)(C)"
25	after "or otherwise".



1	(B) Section 1854(d) (42 U.S.C. 1395w-
2	24(d)) is amended by inserting ", except as pro-
3	$vided\ under\ subsection\ (b)(1)(C),"\ after\ "and$
4	may not provide".
5	(5) Other conforming amendments relating
6	TO BIDS.—Section 1854 (42 U.S.C. 1395w-24) is
7	amended—
8	(A) in the heading of subsection (a), by in-
9	serting "AND BID AMOUNTS" after "PREMIUMS";
10	and
11	(B) in subsection $(a)(5)(A)$ , by inserting
12	"paragraphs (2), (3), and (4) of" after "filed
13	under".
14	(e) Additional Conforming Amendments.—
15	(1) Annual determination and announce-
16	MENT OF CERTAIN FACTORS.—Section 1853(b)(1) (42
17	$U.S.C.\ 1395w-23(b)(1))$ is amended by striking "the
18	respective calendar year" and all that follows and in-
19	serting the following: "the calendar year concerned
20	with respect to each Medicare Advantage payment
21	area, the following:
22	"(A) Pre-competition information.—
23	For years before 2006, the following:
24	"(i) Medicare advantage capita-
25	TION RATES.—The annual Medicare Advan-



1	tage capitation rate for each Medicare Ad-
2	vantage payment area for the year.
3	"(ii) Adjustment factors.—The risk
4	and other factors to be used in adjusting
5	such rates under subsection $(a)(1)(A)$ for
6	payments for months in that year.
7	"(B) Competition information.—For
8	years beginning with 2006, the following:
9	"(i) Benchmark.—The Medicare Ad-
10	vantage area-specific non-drug benchmark
11	$under\ section\ 1853(j).$
12	"(ii) Adjustment factors.—The ad-
13	justment factors applied under section
14	1853(a)(1)(A)(iv) (relating to demographic
15	$adjustment), \ section \ 1853(a)(1)(B) \ (relating$
16	to adjustment for end-stage renal disease),
17	and section 1853(a)(3) (relating to health
18	status adjustment).".
19	(2) Repeal of provisions relating to ad-
20	JUSTED COMMUNITY RATE (ACR).—
21	(A) In general.—Subsections (e) and (f)
22	of section 1854 (42 U.S.C. 1395w-24) are re-
23	pealed.
24	(B) Conforming amendments.—(i) Sec-
25	tion $1839(a)(2)$ (42 U.S.C. $1395r(a)(2)$ ) is



1	amended by striking ", and to reflect" and all
2	that follows and inserting a period.
3	(ii) Section 1852(a)(1) (42 U.S.C. 1395w-
4	22(a)(1)) is amended by striking "title XI" and
5	all that follows and inserting the following: "title
6	XI those items and services (other than hospice
7	care) for which benefits are available under parts
8	A and B to individuals residing in the area
9	served by the plan.".
10	(iii) Section 1857(d)(1) (42 U.S.C. 1395w-
11	27(d)(1)) is amended by striking ", costs, and
12	computation of the adjusted community rate"
13	and inserting "and costs".
14	(f) References under Part E.—Section 1859 (42
15	U.S.C. 1395w-29) is amended by adding at the end the fol-
16	lowing new subsection:
17	"(f) Application under Part E.—In the case of any
18	reference under part E to a requirement or provision of this
19	part in the relation to an EFFS plan or organization under
20	such part, except as otherwise specified any such require-
21	ment or provision shall be applied to such organization or
22	plan in the same manner as such requirement or provision
23	applies to a Medicare Advantage private fee-for-service plan
24	(and the Medicare Advantage organization that offers such
25	plan) under this part.".



1	(g) Effective Date.—The amendments made by this
2	section shall apply to payments and premiums for months
3	beginning with January 2006.
4	CHAPTER 3—ADDITIONAL REFORMS
5	SEC. 231. MAKING PERMANENT CHANGE IN MEDICARE AD-
6	VANTAGE REPORTING DEADLINES AND AN-
7	NUAL, COORDINATED ELECTION PERIOD.
8	(a) Change in Reporting Deadline.—Section
9	1854(a)(1) (42 U.S.C. 1395w-24(a)(1)), as amended by sec-
10	tion 532(b)(1) of the Public Health Security and Bioter-
11	rorism Preparedness and Response Act of 2002, is amended
12	by striking "2002, 2003, and 2004 (or July 1 of each other
13	year)" and inserting "2002 and each subsequent year".
14	(b) Delay in Annual, Coordinated Election Pe-
15	RIOD.—Section $1851(e)(3)(B)$ (42 U.S.C. $1395w$ -
16	21(e)(3)(B)), as amended by section $532(c)(1)(A)$ of the
17	Public Health Security and Bioterrorism Preparedness and
18	Response Act of 2002, is amended—
19	(1) by striking "and after 2005"; and
20	(2) by striking ", 2004, and 2005" and inserting
21	"and any subsequent year".
22	(c) Annual Announcement of Payment Rates.—
23	Section 1853(b)(1) (42 U.S.C. 1395w-23(b)(1)), as amend-
24	ed by section 532(d)(1) of the Public Health Security and



1	Bioterrorism Preparedness and Response Act of 2002, is
2	amended—
3	(1) by striking "and after 2005"; and
4	(2) by striking "and 2005" and inserting "and
5	each subsequent year".
6	(d) Requiring Provision of Available Informa-
7	TION COMPARING PLAN OPTIONS.—The first sentence of sec-
8	tion 1851(d)(2)(A)(ii) (42 U.S.C. 1395w-21(d)(2)(A)(ii)) is
9	amended by inserting before the period the following: "to
10	the extent such information is available at the time of prep-
11	aration of materials for the mailing".
12	SEC. 232. AVOIDING DUPLICATIVE STATE REGULATION.
13	(a) In General.—Section 1856(b)(3) (42 U.S.C.
14	1395w-26(b)(3)) is amended to read as follows:
15	"(3) Relation to state laws.—The standards
16	established under this subsection shall supersede any
17	State law or regulation (other than State licensing
18	laws or State laws relating to plan solvency) with re-
19	spect to Medicare Advantage plans which are offered
20	by Medicare Advantage organizations under this
21	part.".
22	(b) Effective Date.—The amendment made by sub-
23	section (a) shall take effect on the date of the enactment
24	of this Act.



1	SEC. 233. SPECIALIZED MEDICARE ADVANTAGE PLANS FOR
2	SPECIAL NEEDS BENEFICIARIES.
3	(a) Treatment as Coordinated Care Plan.—Sec-
4	tion $1851(a)(2)(A)$ (42 U.S.C. $1395w-21(a)(2)(A)$ ) is
5	amended by adding at the end the following new sentence:
6	"Specialized Medicare Advantage plans for special needs
7	beneficiaries (as defined in section 1859(b)(4)) may be any
8	type of coordinated care plan.".
9	(b) Specialized Medicare Advantage Plan for
10	Special Needs Beneficiaries Defined.—Section
11	1859(b) (42 U.S.C. 1395w-29(b)) is amended by adding at
12	the end the following new paragraph:
13	"(4) Specialized medicare advantage plans
14	FOR SPECIAL NEEDS BENEFICIARIES.—
15	"(A) In General.—The term 'specialized
16	Medicare Advantage plan for special needs bene-
17	ficiaries' means a Medicare Advantage plan that
18	exclusively serves special needs beneficiaries (as
19	defined in subparagraph $(B)$ ).
20	"(B) Special needs beneficiary.—The
21	term 'special needs beneficiary' means a Medi-
22	care Advantage eligible individual who—
23	"(i) is institutionalized (as defined by
24	$the \ Secretary);$
25	"(ii) is entitled to medical assistance
26	under a State plan under title XIX; or



1	"(iii) meets such requirements as the
2	Secretary may determine would benefit
3	from enrollment in such a specialized Medi-
4	care Advantage plan described in subpara-
5	graph (A) for individuals with severe or
6	disabling chronic conditions.".
7	(c) Restriction on Enrollment Permitted.—Sec-
8	tion 1859 (42 U.S.C. 1395w-29) is amended by adding at
9	the end the following new subsection:
10	"(f) Restriction on Enrollment for Specialized
11	Medicare Advantage Plans for Special Needs Bene-
12	FICIARIES.—In the case of a specialized Medicare Advan-
13	tage plan (as defined in subsection (b)(4)), notwithstanding
14	any other provision of this part and in accordance with
15	regulations of the Secretary and for periods before January
16	1, 2007, the plan may restrict the enrollment of individuals
17	under the plan to individuals who are within one or more
18	classes of special needs beneficiaries.".
19	(d) Report to Congress.—Not later than December
20	31, 2005, the Medicare Benefits Administrator shall submit
21	to Congress a report that assesses the impact of specialized
22	Medicare Advantage plans for special needs beneficiaries on
23	the cost and quality of services provided to enrollees. Such
24	report shall include an assessment of the costs and savings



1	to the medicare program as a result of amendments made
2	by subsections (a), (b), and (c).
3	(e) Effective Dates.—
4	(1) In General.—The amendments made by
5	subsections (a), (b), and (c) shall take effect upon the
6	date of the enactment of this Act.
7	(2) Deadline for issuance of requirements
8	FOR SPECIAL NEEDS BENEFICIARIES; TRANSITION.—
9	No later than 6 months after the date of the enact-
10	ment of this Act, the Secretary shall issue interim
11	final regulations to establish requirements for special
12	$needs\ beneficiaries\ under\ section\ 1859(b)(4)(B)(iii)\ og$
13	the Social Security Act, as added by subsection (b).
14	SEC. 234. MEDICARE MSAS.
15	(a) Exemption from Reporting Enrollee En-
16	COUNTER DATA.—
17	(1) In General.—Section 1852(e)(1) (42 U.S.C.
18	1395w-22(e)(1)) is amended by inserting "(other than
19	MSA plans)" after "plans".
20	(2) Conforming amendments.—Section 1852
21	(42 U.S.C. 1395w-22) is amended—
22	(A) in subsection $(c)(1)(I)$ , by inserting be-
23	fore the period at the end the following: "if re-
24	quired under such section"; and



1	(B) in subparagraphs (A) and (B) of sub-
2	section (e)(2), by striking ", a non-network MSA
3	plan," and ", NON-NETWORK MSA PLANS," each
4	place it appears.
5	(b) Making Program Permanent and Eliminating
6	CAP.—Section 1851(b)(4) (42 U.S.C. 1395w-21(b)(4)) is
7	amended—
8	(1) in the heading, by striking "ON A DEM-
9	ONSTRATION BASIS";
10	(2) by striking the first sentence of subparagraph
11	(A); and
12	(3) by striking the second sentence of subpara-
13	graph(C).
14	(c) Applying Limitations on Balance Billing.—
15	Section 1852(k)(1) (42 U.S.C. 1395w-22(k)(1)) is amended
16	by inserting "or with an organization offering a MSA
17	plan" after "section $1851(a)(2)(A)$ ".
18	(d) Additional Amendment.—Section 1851(e)(5)(A)
19	(42 U.S.C. 1395w-21(e)(5)(A)) is amended—
20	(1) by adding "or" at the end of clause (i);
21	(2) by striking ", or" at the end of clause (ii)
22	and inserting a semicolon; and
23	(3) by striking clause (iii).



## SEC. 235. EXTENSION OF REASONABLE COST CONTRACTS. 2 Subparagraph (C) of section 1876(h)(5) (42 U.S.C. 3 1395mm(h)(5)) is amended to read as follows: 4 "(C)(i) Subject to clause (ii), may be extended or re-5 newed under this subsection indefinitely. 6 "(ii) For any period beginning on or after January 1, 2008, a reasonable cost reimbursement contract under this subsection may not be extended or renewed for a service area insofar as such area, during the entire previous year, was within the service area of 2 or more plans which were 10 11 coordinated care Medicare Advantage plans under part C or 2 or more enhanced fee-for-service plans under part E 13 and each of which plan for that previous year for the area involved meets the following minimum enrollment require-15 ments: 16 "(I) With respect to any portion of the area in-17 volved that is within a Metropolitan Statistical Area 18 with a population of more than 250,000 and counties 19 contiguous to such Metropolitan Statistical Area,

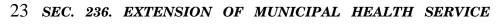


20

"(II) With respect to any other portion of such

22 area, 1,500 individuals.".

5,000 individuals.



24 **DEMONSTRATION PROJECTS.** 

25 Section 9215(a) of the Consolidated Omnibus Budget

**26** Reconciliation Act of 1985 (42 U.S.C. 1395b–1 note), as



1	amended by section 6135 of the Omnibus Budget Reconcili-
2	ation Act of 1989, section 13557 of the Omnibus Budget
3	Reconciliation Act of 1993, section 4017 of BBA, section
4	534 of BBRA (113 Stat. 1501A-390), and section 633 of
5	BIPA, is amended by striking "December 31, 2004" and
6	inserting "December 31, 2009".
7	Subtitle C—Application of FEHBP-
8	Style Competitive Reforms
9	SEC. 241. APPLICATION OF FEHBP-STYLE COMPETITIVE RE-
10	FORM BEGINNING IN 2010.
11	(a) Identification of Competitive EFFS Regions,
12	Computation of Competitive EFFS Non-Drug Bench-
13	MARKS UNDER EFFS PROGRAM.—
14	(1) In general.—Section 1860E-3, as added by
15	section 201(a), is amended by adding at the end the
16	following new subsection:
17	"(e) Application of Competition.—
18	"(1) Determination of competitive effs re-
19	GIONS.—
20	"(A) In General.—For purposes of this
21	part, the term 'competitive EFFS region' means,
22	for a year beginning with 2010, an EFFS region
23	that the Administrator finds—
24	"(i) there will be offered in the region
25	during the annual, coordinated election pe-



1	riod under section 1851(e)(3)(B) (as ap-
2	plied under section 1860E-1(c)) before the
3	beginning of the year at least 2 EFFS plans
4	(in addition to the fee-for-service program
5	under parts A and B), each offered by a dif-
6	ferent EFFS organization and each of
7	which met the minimum enrollment re-
8	quirements of paragraph (1) of section
9	1857(b) (as applied without regard to para-
10	graph (3) thereof) as of March of the pre-
11	vious year; and
12	"(ii) during March of the previous
13	year at least the percentage specified in sub-
14	paragraph (C) of the number of EFFS eli-
15	gible individuals who reside in the region
16	were enrolled in an EFFS plan.
17	"(B) Percentage specified.—
18	"(i) In general.—For purposes of
19	subparagraph (A), subject to clause (ii), the
20	percentage specified in this subparagraph
21	for a year is equal the lesser of 20 percent
22	or to the sum of—
23	"(I) the percentage, as estimated
24	by the Administrator, of EFFS eligible

individuals in the United States who



1	are enrolled in EFFS plans during
2	March of the previous year; and
3	"(II) the percentage, as estimated
4	by the Administrator, of Medicare Ad-
5	vantage eligible individuals in the
6	United States who are enrolled in
7	Medicare Advantage plans during
8	March of the previous year.
9	"(ii) Exception.—In the case of an
10	EFFS region that was a competitive EFFS
11	region for the previous year, the Medicare
12	Benefits Administrator may continue to
13	treat the region as meeting the requirement
14	of subparagraph (A)(ii) if the region would
15	meet such requirement but for a de minimis
16	reduction below the percentage specified in
17	clause $(i)$ .
18	"(2) Competitive effs non-drug monthly
19	BENCHMARK AMOUNT.—For purposes of this part, the
20	term 'competitive EFFS non-drug monthly bench-
21	mark amount' means, with respect to an EFFS re-
22	gion for a month in a year and subject to paragraph
23	(8), the sum of the 2 components described in para-
24	graph (3) for the region and year. The Administrator
25	shall compute such benchmark amount for each com-



1	petitive EFFS region before the beginning of each an-
2	nual, coordinated election period under section
3	1851(e)(3)(B) for each year (beginning with 2010) in
4	which it is designated as such a region.
5	"(3) 2 components.—For purposes of para-
6	graph (2), the 2 components described in this para-
7	graph for an EFFS region and a year are the fol-
8	lowing:
9	"(A) EFFS COMPONENT.—The product of
10	the following:
11	"(i) Weighted average of plan
12	BIDS IN REGION.—The weighted average of
13	the EFFS plan bids for the region and year
14	(as determined under paragraph $(4)(A)$ ).
15	"(ii) Non-ffs market share.—1
16	minus the fee-for-service market share per-
17	centage determined under paragraph (5) for
18	the region and the year.
19	"(B) Fee-for-service component.—The
20	product of the following:
21	"(i) Fee-for-service region-spe-
22	CIFIC NON-DRUG AMOUNT.—The fee-for-serv-
23	ice region-specific non-drug amount (as de-
24	fined in paragraph (6)) for the region and
25	year.



1	"(ii) Fee-for-service market
2	SHARE.—The fee-for-service market share
3	percentage (determined under paragraph
4	(5)) for the region and the year.
5	"(4) Determination of weighted average
6	EFFS PLAN BIDS FOR A REGION.—
7	"(A) In general.—For purposes of para-
8	$graph\ (3)(A)(i),\ the\ weighted\ average\ of\ EFFS$
9	plan bids for an EFFS region and a year is the
10	sum of the following products for EFFS plans
11	described in subparagraph (C) in the region and
12	year:
13	"(i) Unadjusted effs statutory
14	NON-DRUG MONTHLY BID AMOUNT.—The
15	unadjusted EFFS statutory non-drug
16	monthly bid amount (as defined in sub-
17	section $(a)(3)(A)(ii)(I)$ ) for the region and
18	year.
19	"(ii) Plan's share of effs enroll-
20	MENT IN REGION.—The number of individ-
21	uals described in subparagraph (B), divided
22	by the total number of such individuals for
23	all EFFS plans described in subparagraph
24	(C) for that region and year.



1	"(B) Counting of individuals.—The Ad-
2	ministrator shall count, for each EFFS plan de
3	scribed in subparagraph (C) for an EFFS region
4	and year, the number of individuals who reside
5	in the region and who were enrolled under such
6	plan under this part during March of the pre-
7	vious year.
8	"(C) Exclusion of plans not offered
9	IN PREVIOUS YEAR.—For an EFFS region and
10	year, the EFFS plans described in this subpara
11	graph are plans that are offered in the region
12	and year and were offered in the region in
13	March of the previous year.
14	"(5) Computation of fee-for-service mar-
15	KET SHARE PERCENTAGE.—The Administrator shall
16	determine, for a year and an EFFS region, the pro-
17	portion (in this subsection referred to as the 'fee-for-
18	service market share percentage') of the EFFS eligible
19	individuals who are residents of the region during
20	March of the previous year, of such individuals who
21	were not enrolled in an EFFS plan or in a Medicare
22	Advantage plan (or, if greater, such proportion deter-
23	mined for individuals nationally).
24	"(6) Fee-for-service region-specific non-
25	DRUG AMOUNT.—



1	"(A) In general.—For purposes of para-
2	$graph\ (3)(B)(i)\ and\ section\ 1839(h)(2)(A),\ sub-$
3	ject to subparagraph (B), the term 'fee-for-service
4	region-specific non-drug amount' means, for a
5	competitive EFFS region and a year, the ad-
6	justed average per capita cost for the year in-
7	volved, determined under section 1876(a)(4) for
8	such region for services covered under parts A
9	and B for individuals entitled to benefits under
10	part A and enrolled under this part who are not
11	enrolled in an EFFS plan under part E or a
12	Medicare Advantage plan under part C for the
13	year, but adjusted to exclude costs attributable to
14	payments under section 1886(h).
15	"(B) Inclusion of costs of va and dod
16	MILITARY FACILITY SERVICES TO MEDICARE-ELI-
17	GIBLE BENEFICIARIES.—In determining the ad-
18	justed average per capita cost under subpara-
19	graph (A) for a year, such cost shall be adjusted
20	to include the Administrator's estimate, on a per
21	capita basis, of the amount of additional pay-
22	ments that would have been made in the region
23	involved under this title if individuals entitled to

benefits under this title had not received services



1	from facilities of the Department of Veterans Af-
2	fairs or the Department of Defense.
3	"(7) Application of competition.—In the case
4	of an EFFS region that is a competitive EFFS re-
5	gion for a year, for purposes of applying subsections
6	(b) and (c)(1) and section 1860E-4(a), any reference
7	to an EFFS region-specific non-drug monthly bench-
8	mark amount shall be treated as a reference to the
9	competitive EFFS non-drug monthly benchmark
10	amount under paragraph (2) for the region and year.
11	"(8) Phase-in of Benchmark for each re-
12	GION.—
13	"(A) Use of blended benchmark.—In
14	the case of a region that has not been a competi-
15	tive EFFS region for each of the previous 4
16	years, the competitive EFFS non-drug monthly
17	benchmark amount shall be equal to the sum of
18	$the\ following:$
19	"(i) New competitive component.—
20	The product of—
21	"(I) the weighted average phase-in
22	proportion for that area and year, as
23	specified in subparagraph (B); and
24	"(II) the competitive EFFS non-
25	drug monthly benchmark amount for



1	the region and year, determined under
2	paragraph (2) without regard to this
3	paragraph.
4	"(ii) Old competitive component.—
5	The product of—
6	"(I) 1 minus the weighted average
7	phase-in proportion for that region
8	and year; and
9	"(II) the EFFS region-specific
10	non-drug benchmark amount for the
11	region and the year.
12	"(B) Computation of Weighted Average
13	PHASE-IN PROPORTION.—For purposes of this
14	paragraph, the 'weighted average phase-in pro-
15	portion' for an EFFS region for a year shall be
16	determined as follows:
17	"(i) First year (and region not
18	COMPETITIVE REGION IN PREVIOUS
19	YEAR).—If the area was not a competitive
20	EFFS region in the previous year, the
21	weighted average phase-in proportion for
22	the region for the year is equal to ½.
23	"(ii) Competitive region in pre-
24	VIOUS YEAR.—If the region was a competi-
25	tive EFFS region in the previous year, the



1	weighted average phase-in proportion for
2	the region for the year is equal to the
3	weighted average phase-in proportion deter-
4	mined under this subparagraph for the re-
5	gion for the previous year plus ½, but in
6	no case more than 1.".
7	(2) Conforming amendments.—
8	(A) Such section 1860E-3 is further
9	amended—
10	(i) in subsection (b), by adding at the
11	end the following new paragraph:
12	"(4) Application in competitive re-
13	GIONS.—For special rules applying this sub-
14	section in competitive EFFS regions, see sub-
15	section (e)(7).";
16	(ii) in subsection (c)(1), by inserting
17	"and subsection (e)(7)" after "(as made ap-
18	plicable under subsection (d))"; and
19	(iii) in subsection (d) , by striking
20	"and (e)" and inserting "(e), and (k)".
21	(B) Section 1860 $E$ -4(a)(1), as inserted by
22	section 201(a)(2), is amended by inserting ", ex-
23	cept as provided in section 1860E-3(e)(7)" after
24	"paragraph (2)".



1	(b) Identification of Competitive Medicare Ad-
2	VANTAGE AREAS; APPLICATION OF COMPETITIVE MEDICARE
3	Advantage Non-Drug Benchmarks Under Medicare
4	Advantage Program.—
5	(1) In general.—Section 1853, as amended by
6	section 221(b)(3), is amended by adding at the end
7	the following new subsection:
8	"(k) Application of Competition.—
9	"(1) Determination of competitive medi-
10	CARE ADVANTAGE AREAS.—
11	"(A) In General.—For purposes of this
12	part, the terms 'competitive Medicare Advantage
13	area' and 'CMA area' mean, for a year begin-
14	ning with 2010, an area (which is a metropoli-
15	tan statistical area or other area with a substan-
16	tial number of Medicare Advantage enrollees)
17	that the Administrator finds—
18	"(i) there will be offered during the an-
19	nual, coordinated election period under sec-
20	tion $1851(e)(3)(B)$ under this part before
21	the beginning of the year at least 2 Medi-
22	care Advantage plans (in addition to the
23	fee-for-service program under parts A and
24	B), each offered by a different Medicare Ad-
25	vantage organization and each of which met



1	the minimum enrollment requirements of
2	paragraph (1) of section 1857(b) (as ap-
3	plied without regard to paragraph (3)
4	thereof) as of March of the previous year
5	with respect to the area; and
6	"(ii) during March of the previous
7	year at least the percentage specified in sub-
8	paragraph (B) of the number of Medicare
9	Advantage eligible individuals who reside in
10	the area were enrolled in a Medicare Ad-
11	vantage plan.
12	"(B) Percentage specified.—
13	"(i) In General.—For purposes of
14	subparagraph (A), subject to clause (ii), the
15	percentage specified in this subparagraph
16	for a year is equal the lesser of 20 percent
17	or to the sum of—
18	"(I) the percentage, as estimated
19	by the Administrator, of EFFS eligible
20	individuals in the United States who
21	are enrolled in EFFS plans during
22	March of the previous year; and
23	"(II) the percentage, as estimated
24	by the Administrator, of Medicare Ad-
25	vantage eligible individuals in the



1	United States who are enrolled in
2	Medicare Advantage plans during
3	March of the previous year.
4	"(ii) Exception.—In the case of an
5	area that was a competitive area for the
6	previous year, the Medicare Benefits Ad-
7	ministrator may continue to treat the area
8	as meeting the requirement of subparagraph
9	(A)(ii) if the area would meet such require-
10	ment but for a de minimis reduction below
11	the percentage specified in clause (i).
12	"(2) Competitive medicare advantage non-
13	DRUG MONTHLY BENCHMARK AMOUNT.—For purposes
14	of this part, the term 'competitive Medicare Advan-
15	tage non-drug monthly benchmark amount' means,
16	with respect to a competitive Medicare Advantage
17	area for a month in a year subject to paragraph (8),
18	the sum of the 2 components described in paragraph
19	(3) for the area and year. The Administrator shall
20	compute such benchmark amount for each competitive
21	Medicare Advantage area before the beginning of each
22	annual, coordinated election period under section
23	1851(e)(3)(B) for each year (beginning with 2010) in
24	which it is designated as such an area.



1	"(3) 2 COMPONENTS.—For purposes of para-
2	graph (2), the 2 components described in this para-
3	graph for a competitive Medicare Advantage area and
4	a year are the following:
5	"(A) Medicare advantage component.—
6	The product of the following:
7	"(i) Weighted average of medi-
8	CARE ADVANTAGE PLAN BIDS IN AREA.—The
9	weighted average of the plan bids for the
10	area and year (as determined under para-
11	graph(4)(A)).
12	"(ii) Non-ffs market share.—1
13	minus the fee-for-service market share per-
14	centage, determined under paragraph (5)
15	for the area and year.
16	"(B) Fee-for-service component.—The
17	product of the following:
18	"(i) Fee-for-service area-specific
19	NON-DRUG AMOUNT.—The fee-for-service
20	area-specific non-drug amount (as defined
21	in paragraph (6)) for the area and year.
22	"(ii) Fee-for-service market
23	SHARE.—The fee-for-service market share
24	percentage, determined under paragraph (5)
25	for the area and year.



1	"(4) Determination of weighted average
2	MEDICARE ADVANTAGE BIDS FOR AN AREA.—
3	"(A) In general.—For purposes of para-
4	$graph\ (3)(A)(i),\ the\ weighted\ average\ of\ plan$
5	bids for an area and a year is the sum of the
6	following products for Medicare Advantage plans
7	described in subparagraph (C) in the area and
8	year:
9	"(i) Monthly medicare advantage
10	STATUTORY NON-DRUG BID AMOUNT.—The
11	unadjusted Medicare Advantage statutory
12	non-drug monthly bid amount.
13	"(ii) Plan's share of medicare ad-
14	VANTAGE ENROLLMENT IN AREA.—The
15	number of individuals described in subpara-
16	graph (B), divided by the total number of
17	such individuals for all Medicare Advantage
18	plans described in subparagraph (C) for
19	that area and year.
20	"(B) Counting of individuals.—The Ad-
21	ministrator shall count, for each Medicare Ad-
22	vantage plan described in subparagraph (C) for
23	an area and year, the number of individuals
24	who reside in the area and who were enrolled



1	under such plan under this part during March
2	of the previous year.
3	"(C) Exclusion of plans not offered
4	IN PREVIOUS YEAR.—For an area and year, the
5	Medicare Advantage plans described in this sub-
6	paragraph are plans described in the first sen-
7	tence of section 1851(a)(2)(A) that are offered in
8	the area and year and were offered in the area
9	in March of the previous year.
10	"(5) Computation of fee-for-service mar-
11	KET SHARE PERCENTAGE.—The Administrator shall
12	determine, for a year and a competitive Medicare Ad-
13	vantage area, the proportion (in this subsection re-
14	ferred to as the 'fee-for-service market share percent-
15	age') of Medicare Advantage eligible individuals re-
16	siding in the area who during March of the previous
17	year were not enrolled in a Medicare Advantage plan
18	or in an EFFS plan (or, if greater, such proportion
19	determined for individuals nationally).
20	"(6) Fee-for-service area-specific non-
21	DRUG AMOUNT.—
22	"(A) In general.—For purposes of para-
23	$graph\ (3)(B)(i)\ and\ section\ 1839(h)(1)(A),\ sub-$
24	ject to subparagraph (B), the term 'fee-for-service

area-specific non-drug amount' means, for a



1	competitive Medicare Advantage area and a
2	year, the adjusted average per capita cost for the
3	year involved, determined under section
4	1876(a)(4) for such area for services covered
5	under parts A and B for individuals entitled to
6	benefits under part A and enrolled under this
7	part who are not enrolled in a Medicare Advan-
8	tage plan under part C or an EFFS plan under
9	part E for the year, but adjusted to exclude costs
10	attributable to payments under section 1886(h).
11	"(B) Inclusion of costs of va and dod
12	MILITARY FACILITY SERVICES TO MEDICARE-ELI-
13	GIBLE BENEFICIARIES.—In determining the ad-
14	justed average per capita cost under subpara-
15	graph (A) for a year, such cost shall be adjusted
16	to include the Administrator's estimate, on a per
17	capita basis, of the amount of additional pay-
18	ments that would have been made in the area in-
19	volved under this title if individuals entitled to
20	benefits under this title had not received services
21	from facilities of the Department of Veterans Af-
22	fairs or the Department of Defense.
23	"(7) APPLICATION OF COMPETITION.—In the case
24	of an area that is a competitive Medicare Advantage

 $area\ for\ a\ year,\ for\ purposes\ of\ applying\ subsection$ 



1	(a)(1)(A)(ii) and sections $1854(b)(2)(A)(ii)$ and
2	1854(b)(3)(B)(i), any reference to a Medicare Advan-
3	tage area-specific non-drug monthly benchmark
4	amount shall be treated as a reference to the competi-
5	tive Medicare Advantage non-drug monthly bench-
6	mark amount under paragraph (2) for the area and
7	year.
8	"(8) Phase-in of Benchmark for each
9	AREA.—
10	"(A) Use of blended benchmark.—In
11	the case of an area that has not been a competi-
12	tive Medicare Advantage area for each of the pre-
13	vious 4 years, the competitive Medicare Advan-
14	tage non-drug monthly benchmark amount shall
15	be equal to the sum of the following:
16	"(i) New competitive component.—
17	The product of—
18	"(I) the weighted average phase-in
19	proportion for that area and year, as
20	specified in subparagraph (B); and
21	"(II) the competitive Medicare
22	Advantage non-drug monthly bench-
23	mark amount for the area and year,
24	determined under paragraph (2) with-
25	out regard to this paragraph.



1	"(ii) Old competitive component.—
2	The product of—
3	"(I) 1 minus the weighted average
4	phase-in proportion for that area and
5	year; and
6	"(II) the Medicare Advantage
7	area-wide non-drug benchmark amount
8	for the area and the year.
9	"(B) Computation of weighted average
10	PHASE-IN PROPORTION.—For purposes of this
11	paragraph, the 'weighted average phase-in pro-
12	portion' for a Medicare Advantage payment area
13	for a year shall be determined as follows:
14	"(i) First year (and area not com-
15	PETITIVE AREA IN PREVIOUS YEAR).—If the
16	area was not a Medicare Advantage com-
17	petitive area in the previous year, the
18	weighted average phase-in proportion for
19	the area for the year is equal to ½.
20	"(ii) Competitive area in previous
21	YEAR.—If the area was a competitive Medi-
22	care Advantage area in the previous year,
23	the weighted average phase-in proportion
24	for the area for the year is equal to the
25	weighted average phase-in proportion deter-



1	mined under this subparagraph for the area
2	for the previous year plus ½, but in no case
3	more than 1.
4	"(C) Medicare advantage area-wide
5	NON-DRUG BENCHMARK AMOUNT.—For purposes
6	of $subparagraph\ (A)(ii)(II),\ the\ term\ `Medicare$
7	Advantage area-wide non-drug benchmark
8	amount' means, for an area and year, the
9	weighted average of the amounts described in sec-
10	tion 1853(j) for Medicare Advantage payment
11	area or areas included in the area (based on the
12	number of traditional fee-for-service enrollees in
13	such payment area or areas) and year.".
14	(2) Application.—Section 1854 (42 U.S.C.
15	1395w-24) is amended—
16	(A) in subsection $(b)(1)(C)(i)$ , as added by
17	section 221(b)(1)(A), by striking "(i) Require-
18	MENT.—The" and inserting "(i) REQUIREMENT
19	FOR NON-COMPETITIVE AREAS.—In the case of a
20	Medicare Advantage payment area that is not a
21	competitive Medicare Advantage area designated
22	under section 1853(k)(1), the";
23	(B) in subsection $(b)(1)(C)$ , as so added, by
24	inserting after clause (i) the following new
25	clause:



1	"(ii) Requirement for competitive
2	MEDICARE ADVANTAGE AREAS.—In the case
3	of a Medicare Advantage payment area that
4	is designated as a competitive Medicare Ad-
5	$vantage \ area \ under \ section \ 1853(k)(1), \ if$
6	there are average per capita monthly sav-
7	ings described in paragraph (6) for a Medi-
8	care Advantage plan and year, the Medicare
9	Advantage plan shall provide to the enrollee
10	a monthly rebate equal to 75 percent of such
11	savings."; and
12	(C) by adding at the end of subsection (b),
13	as amended by sections $221(b)(1)(B)$ and
14	221(b)(2), the following new paragraph:
15	"(6) Computation of average per capita
16	MONTHLY SAVINGS FOR COMPETITIVE MEDICARE AD-
17	VANTAGE AREAS.—For purposes of paragraph
18	(1)(C)(ii), the average per capita monthly savings re-
19	ferred to in such paragraph for a Medicare Advantage
20	plan and year shall be computed in the same manner
21	as the average per capita monthly savings is com-
22	puted under paragraph (3) except that the reference
23	to the Medicare Advantage area-specific non-drug
24	$monthly\ benchmark\ amount\ in\ paragraph\ (3)(B)(i)$
25	(or to the benchmark amount as adjusted under para-



1	graph $(3)(C)(i)$ ) is deemed to be a reference to the
2	competitive Medicare Advantage non-drug monthly
3	benchmark amount (or such amount as adjusted in
4	the manner described in paragraph $(3)(B)(i)$ .".
5	(3) Additional conforming amendments.—
6	(A) PAYMENT OF PLANS.—Section
7	1853(a)(1)(A)(ii), as amended by section
8	221(c)(1), is amended—
9	(i) in subclauses (I) and (II), by in-
10	serting "(or, insofar as such payment area
11	is a competitive Medicare Advantage area,
12	described in section 1854(b)(6))" after "sec-
13	tion $1854(b)(3)(C)$ "; and
14	(ii) in subclause (II), by inserting
15	"(or, insofar as such payment area is a
16	competitive Medicare Advantage area, the
17	competitive Medicare Advantage non-drug
18	monthly benchmark amount)" after "Medi-
19	care Advantage area-specific non-drug
20	monthly benchmark amount"; and
21	(B) Disclosure of information.—Sec-
22	tion $1853(b)(1)(B)$ , as amended by section
23	221(e)(1), is amended to read as follows:
24	"(B) Competition information.—For
25	years beginning with 2006, the following:



1	"(i) Benchmarks.—The Medicare Ad-
2	vantage area-specific non-drug benchmark
3	under section 1853(j) and, if applicable, the
4	competitive Medicare Advantage non-drug
5	benchmark under section $1853(k)(2)$ , for the
6	year and competitive Medicare Advantage
7	area involved and the national fee-for-serv-
8	ice market share percentage for the area
9	and year.
10	"(ii) Adjustment factors.—The ad-
11	justment factors applied under section
12	1853(a)(1)(A)(iv) (relating to demographic
13	adjustment), section 1853(a)(1)(B) (relating
14	to adjustment for end-stage renal disease),
15	and section 1853(a)(3) (relating to health
16	$status\ adjustment).$
17	"(iii) Certain benchmarks and
18	AMOUNTS.—In the case of a competitive
19	Medicare Advantage area, the Medicare Ad-
20	vantage area-wide non-drug benchmark
21	amount (as defined in subsection $(k)(8)(C)$ )
22	and the fee-for-service area-specific non-
23	drug amount (as defined in section

1853(k)(6)) for the area.



1	``(iv) Individuals.—The number of
2	individuals counted under subsection
3	(k)(4)(B) and enrolled in each Medicare Ad-
4	vantage plan in the area.".
5	(C) Definition of monthly basic pre-
6	MIUM.—Section $1854(b)(2)(A)(ii)$ , as amended
7	by section 221(d)(2), is amended by inserting
8	"(or, in the case of a competitive Medicare Ad-
9	vantage area, the competitive Medicare Advan-
10	tage non-drug monthly benchmark amount or, in
11	applying this paragraph under part $E$ in the
12	case of a competitive EFFS region, the competi-
13	tive EFFS non-drug monthly benchmark
14	amount)" after "benchmark amount".
15	(c) Premium Adjustment.—
16	(1) In General.—Section 1839 (42 U.S.C.
17	1395r) is amended by adding at the end the following
18	new subsection:
19	"(h)(1)(A) In the case of an individual who resides in
20	a competitive Medicare Advantage area under section
21	1853(k)(1) (regardless of whether such area is in a competi-
22	tive EFFS region under section 1860E-3(e)) and who is
23	not enrolled in a Medicare Advantage plan under part C
24	or in an EFFS plan under part E, the monthly premium
25	otherwise applied under this part (determined without re-



1	gard to subsections (b) and (f) or any adjustment under
2	this subsection) shall be adjusted as follows: If the fee-for-
3	service area-specific non-drug amount (as defined in section
4	1853(k)(6)) for the competitive Medicare Advantage area in
5	which the individual resides for a month—
6	"(i) does not exceed the competitive Medicare Ad-
7	vantage non-drug benchmark (as determined under
8	paragraph (2) of section 1853(k), without regard to
9	paragraph (8) thereof) for such area, the amount of
10	the premium for the individual for the month shall be
11	reduced by an amount equal to the product of the ad-
12	justment factor under subparagraph (C) and 75 per-
13	cent of the amount by which such competitive bench-
14	mark exceeds such fee-for-service area-specific non-
15	drug amount; or
16	"(ii) exceeds such competitive Medicare Advan-
17	tage non-drug benchmark, the amount of the premium
18	for the individual for the month shall be adjusted to
19	ensure, subject to subparagraph (B), that—
20	"(I) the sum of the amount of the adjusted
21	premium and the competitive Medicare Advan-
22	tage non-drug benchmark for the area, is equal
23	to



1	"(II) the sum of the unadjusted premium
2	plus amount of the fee-for-service area-specific
3	non-drug amount for the area.
4	"(B) In no case shall the actual amount of an adjust-
5	ment under subparagraph (A)(ii) exceed the product of the
6	adjustment factor under subparagraph (C) and the amount
7	of the adjustment otherwise computed under subparagraph
8	(A)(ii) without regard to this subparagraph.
9	"(C) The adjustment factor under this subparagraph
10	for an area for a year is equal to—
11	"(i) the number of consecutive years (in the 5-
12	year period ending with the year involved) in which
13	such area was a competitive Medicare Advantage
14	area; divided by
15	"(ii) 5.
16	"(2)(A) In the case of an individual who resides in
17	an area that is within a competitive EFFS region under
18	section 1860E-3(e) but is not within a competitive Medi-
19	care Advantage area under section 1853(k)(1) and who is
20	not enrolled in a Medicare Advantage plan under part C
21	or in an EFFS plan under part E, the monthly premium
22	otherwise applied under this part (determined without re-
23	gard to subsections (b) and (f) or any adjustment under
24	this subsection) shall be adjusted as follows: If the fee-for-



1	service region-specific non-drug amount (as defined in sec-
2	tion 1860E-3(e)(6)) for a region for a month—
3	"(i) does not exceed the competitive EFFS non-
4	drug monthly benchmark amount (as determined
5	under paragraph (2) of section 1860E-3(e), without
6	regard to paragraph (8) thereof) for such region, the
7	amount of the premium for the individual for the
8	month shall be reduced by an amount equal to the
9	product of the adjustment factor under subparagraph
10	(C) and 75 percent of the amount by which such com-
11	petitive benchmark amount exceeds such fee-for-service
12	region-specific non-drug benchmark amount; or
13	"(ii) exceeds such competitive EFFS non-drug
14	monthly benchmark amount, the amount of the pre-
15	mium for the individual for the month shall be ad-
16	justed to ensure, subject to subparagraph (B), that—
17	"(I) the sum of the amount of the adjusted
18	premium and the competitive EFFS non-drug
19	monthly benchmark amount for the region, is
20	equal to
21	"(II) the sum of the unadjusted premium
22	plus the amount of the EFFS region-specific
23	non-drug monthly bidfor the region.
24	"(B) In no case shall the actual amount of an adjust-
25	ment under subparagraph (A)(ii) exceed the product of the



adjustment factor under subparagraph (C) and the amount of the adjustment otherwise computed under subparagraph 3 (A)(ii) without regard to this subparagraph. 4 "(C) The adjustment factor under this subparagraph for an EFFS region for a year is equal to— 6 "(i) the number of consecutive years (in the 5-7 year period ending with the year involved) in which 8 such region was a competitive EFFS region; divided 9 by10 "(ii) 5. "(3) Nothing in this subsection shall be construed as 11 preventing a reduction under paragraph (1)(A) or paragraph (2)(A) in the premium otherwise applicable under this part to zero or from requiring the provision of a rebate 14 15 to the extent such premium would otherwise be required to be less than zero. 16 17 "(4) The adjustment in the premium under this sub-18 section shall be effected in such manner as the Medicare Benefits Administrator determines appropriate. 19 20 "(5) In order to carry out this subsection (insofar as 21 it is effected through the manner of collection of premiums 22 under 1840(a)), the Medicare Benefits Administrator shall 23 transmit to the Commissioner of Social Security— 24 "(A) at the beginning of each year, the name, so-

cial security account number, and the amount of the



1	adjustment (if any) under this subsection for each in-
2	dividual enrolled under this part for each month dur-
3	ing the year; and
4	"(B) periodically throughout the year, informa-
5	tion to update the information previously transmitted
6	under this paragraph for the year.".
7	(2) Conforming amendment.—Section 1844(c)
8	(42 U.S.C. $1395w(c)$ ) is amended by inserting "and
9	without regard to any premium adjustment effected
10	under section 1839(h)" before the period at the end.
11	(d) Effective Date.—The amendments made by this
12	section shall take effect on January 1, 2010.
13	TITLE III—COMBATTING WASTE,
14	FRAUD, AND ABUSE
15	SEC. 301. MEDICARE SECONDARY PAYOR (MSP) PROVI
16	SIONS.
17	5101151
1 /	(a) Technical Amendment Concerning Sec-
	(a) Technical Amendment Concerning Sec-
18	(a) Technical Amendment Concerning Sec- retary's Authority to Make Conditional Payment
18 19	(a) Technical Amendment Concerning Sec- retary's Authority to Make Conditional Payment When Certain Primary Plans Do Not Pay Prompt-
18 19 20	(a) Technical Amendment Concerning Sec- retary's Authority to Make Conditional Payment When Certain Primary Plans Do Not Pay Prompt- ly.—
18 19 20 21	(a) Technical Amendment Concerning Sec- retary's Authority to Make Conditional Payment When Certain Primary Plans Do Not Pay Prompt- ly.—  (1) In General.—Section 1862(b)(2) (42 U.S.C.
18 19 20 21 22	(a) Technical Amendment Concerning Sec- retary's Authority to Make Conditional Payment When Certain Primary Plans Do Not Pay Prompt- ly.—  (1) In General.—Section 1862(b)(2) (42 U.S.C. 1395y(b)(2)) is amended—



1	$(B)\ in\ subparagraph\ (B)$ —
2	(i) by redesignating clauses (i) through
3	(iii) as clauses (ii) through (iv), respec-
4	tively; and
5	(ii) by inserting before clause (ii), as
6	so redesignated, the following new clause:
7	"(i) Authority to make condi-
8	TIONAL PAYMENT.—The Secretary may
9	make payment under this title with respect
10	to an item or service if a primary plan de-
11	scribed in subparagraph (A)(ii) has not
12	made or cannot reasonably be expected to
13	make payment with respect to such item or
14	service promptly (as determined in accord-
15	ance with regulations). Any such payment
16	by the Secretary shall be conditioned on re-
17	imbursement to the appropriate Trust Fund
18	in accordance with the succeeding provi-
19	sions of this subsection.".
20	(2) Effective date.—The amendments made
21	by paragraph (1) shall be effective as if included in
22	the enactment of title III of the Medicare and Med-
23	icaid Budget Reconciliation Amendments of 1984
24	(Public Law 98-369).



1	(b) Clarifying Amendments to Conditional Pay-
2	MENT PROVISIONS.—Section 1862(b)(2) (42 U.S.C.
3	1395y(b)(2)) is further amended—
4	(1) in subparagraph (A), in the matter following
5	clause (ii), by inserting the following sentence at the
6	end: "An entity that engages in a business, trade, or
7	profession shall be deemed to have a self-insured plan
8	if it carries its own risk (whether by a failure to ob-
9	tain insurance, or otherwise) in whole or in part.";
10	(2) in subparagraph (B)(ii), as redesignated by
11	$subsection \ (a)(2)(B)$ —
12	(A) by striking the first sentence and insert-
13	ing the following: "A primary plan, and an enti-
14	ty that receives payment from a primary plan,
15	shall reimburse the appropriate Trust Fund for
16	any payment made by the Secretary under this
17	title with respect to an item or service if it is
18	demonstrated that such primary plan has or had
19	a responsibility to make payment with respect to
20	such item or service. A primary plan's responsi-
21	bility for such payment may be demonstrated by
22	a judgment, a payment conditioned upon the re-
23	cipient's compromise, waiver, or release (whether
24	or not there is a determination or admission of

liability) of payment for items or services in-



	<b></b>
1	cluded in a claim against the primary plan or
2	the primary plan's insured, or by other means.";
3	and
4	(B) in the final sentence, by striking "on
5	the date such notice or other information is re-
6	ceived" and inserting "on the date notice of, or
7	information related to, a primary plan's respon-
8	sibility for such payment or other information is
9	received"; and
10	(3) in subparagraph (B)(iii), , as redesignated
11	by subsection (a)(2)(B), by striking the first sentence
12	and inserting the following: "In order to recover pay-
13	ment made under this title for an item or service, the
14	United States may bring an action against any or all
15	entities that are or were required or responsible (di-
16	rectly, as an insurer or self-insurer, as a third-party
17	administrator, as an employer that sponsors or con-
18	tributes to a group health plan, or large group health
19	plan, or otherwise) to make payment with respect to
20	the same item or service (or any portion thereof)
21	under a primary plan. The United States may, in ac-
22	cordance with paragraph (3)(A) collect double dam-
23	ages against any such entity. In addition, the United
24	States may recover under this clause from any entity

that has received payment from a primary plan or



1	from the proceeds of a primary plan's payment to
2	any entity.".
3	(c) Clerical Amendments.—Section 1862(b) (42
4	$U.S.C.\ 1395y(b))$ is amended—
5	(1) in paragraph (1)(A), by moving the indenta-
6	tion of clauses (ii) through (v) 2 ems to the left; and
7	(2) in paragraph (3)(A), by striking "such" be-
8	fore "paragraphs".
9	SEC. 302. COMPETITIVE ACQUISITION OF CERTAIN ITEMS
10	AND SERVICES.
11	(a) In General.—Section 1847 (42 U.S.C. 1395w-
12	3) is amended to read as follows:
13	"COMPETITIVE ACQUISITION OF CERTAIN ITEMS AND
14	SERVICES
15	"Sec. 1847. (a) Establishment of Competitive
16	Acquisition Programs.—
17	"(1) Implementation of programs.—
18	"(A) In General.—The Secretary shall es-
19	tablish and implement programs under which
20	competitive acquisition areas are established
21	throughout the United States for contract award
22	purposes for the furnishing under this part of
23	competitively priced items and services (de-
24	scribed in paragraph (2)) for which payment is
25	made under this part. Such areas may differ for
26	different items and services.



1	"(B) Phased-in implementation.—The
2	programs shall be phased-in—
3	"(i) among competitive acquisition
4	areas over a period of not longer than 3
5	years in a manner so that the competition
6	under the programs occurs in—
7	"(I) at least 1/3 of such areas in
8	2005; and
9	"(II) at least 2/3 of such areas in
10	2006; and
11	"(ii) among items and services in a
12	manner such that the programs apply to the
13	highest cost and highest volume items and
14	services first.
15	"(C) Waiver of Certain Provisions.—In
16	carrying out the programs, the Secretary may
17	waive such provisions of the Federal Acquisition
18	Regulation as are necessary for the efficient im-
19	plementation of this section, other than provi-
20	sions relating to confidentiality of information
21	and such other provisions as the Secretary deter-
22	mines appropriate.
23	"(2) Items and services described.—The
24	items and services referred to in paragraph (1) are
25	the following:



1	"(A) Durable medical equipment and
2	MEDICAL SUPPLIES.—Covered items (as defined
3	in section 1834(a)(13)) for which payment is
4	otherwise made under section 1834(a), including
5	items used in infusion and drugs and supplies
6	used in conjunction with durable medical equip-
7	ment, but excluding class III devices under the
8	Federal Food, Drug, and Cosmetic Act.
9	"(B) Other equipment and supplies.—
10	Items, equipment, and supplies (as described in
11	section $1842(s)(2)(D)$ other than enteral nutri-
12	ents).
13	"(C) Off-the-shelf orthotics.—
14	Orthotics (described in section 1861(s)(9)) for
15	which payment is otherwise made under section
16	1834(h) which require minimal self-adjustment
17	for appropriate use and does not require exper-
18	tise in trimming, bending, molding, assembling,
19	or customizing to fit to the patient.
20	"(3) Exception authority.—In carrying out
21	the programs under this section, the Secretary may
22	exempt—
23	"(A) rural areas and areas with low popu-
24	lation density within urban areas that are not
25	competitive, unless there is a significant national



1	market through mail order for a particular item
2	or service; and
3	"(B) items and services for which the appli-
4	cation of competitive acquisition is not likely to
5	result in significant savings.
6	"(4) Special rule for certain rented items
7	OF DURABLE MEDICAL EQUIPMENT.—In the case of a
8	covered item for which payment is made on a rental
9	basis under section 1834(a), the Secretary shall estab-
10	lish a process by which rental agreements for the cov-
11	ered items entered into before the application of the
12	competitive acquisition program under this section
13	for the item may be continued notwithstanding this
14	section. In the case of any such continuation, the sup-
15	plier involved shall provide for appropriate servicing
16	and replacement, as required under section 1834(a).
17	"(5) Physician authorization.—The Sec-
18	retary may establish a process under which a physi-
19	cian may prescribe a particular brand or mode of de-
20	livery of an item or service if the item or service in-
21	volved is clinically more appropriate than other simi-
22	lar items or services.
23	"(6) APPLICATION.—For each competitive acqui-
24	sition area in which the program is implemented

under this subsection with respect to items and serv-



1	ices, the payment basis determined under the competi-
2	tion conducted under subsection (b) shall be sub-
3	stituted for the payment basis otherwise applied
4	$under\ section\ 1834(a).$
5	"(b) Program Requirements.—
6	"(1) In general.—The Secretary shall conduct
7	a competition among entities supplying items and
8	services described in subsection (a)(2) for each com-
9	petitive acquisition area in which the program is im-
10	plemented under subsection (a) with respect to such
11	items and services.
12	"(2) Conditions for awarding contract.—
13	"(A) In general.—The Secretary may not
14	award a contract to any entity under the com-
15	petition conducted in an competitive acquisition
16	area pursuant to paragraph (1) to furnish such
17	items or services unless the Secretary finds all of
18	$the\ following:$
19	"(i) The entity meets quality and fi-
20	nancial standards specified by the Secretary
21	or developed by the Program Advisory and
22	Oversight Committee established under sub-
23	section (c).
24	"(ii) The total amounts to be paid
25	under the contract (including costs associ-



1	ated with the administration of the con-
2	tract) are expected to be less than the total
3	amounts that would otherwise be paid.
4	"(iii) Beneficiary access to a choice of
5	multiple suppliers in the area is main-
6	tained.
7	"(iv) Beneficiary liability is limited to
8	20 percent of the applicable contract award
9	price, except in such cases where a supplier
10	has furnished an upgraded item and has ex-
11	ecuted an advanced beneficiary notice.
12	"(B) Development of quality stand-
13	ARDS FOR DME PRODUCTS.—
14	"(i) In general.—The quality stand-
15	ards $specified$ $under$ $subparagraph$ $(A)(i)$
16	shall not be less than the quality standards
17	that would otherwise apply if this section
18	did not apply and shall include consumer
19	services standards. Not later than July 1,
20	2004, the Secretary shall establish new
21	quality standards for products subject to
22	competitive acquisition under this section.
23	Such standards shall be applied prospec-
24	tively and shall be published on the website



1	of the Department of Health and Human
2	Services.
3	"(ii) Consultation with program
4	ADVISORY AND OVERSIGHT COMMITTEE.—
5	The Secretary shall consult with the Pro-
6	gram Advisory and Oversight Committee
7	(established under subsection (c)) to review
8	(and advise the Secretary concerning) the
9	quality standards referred to in clause (i).
10	"(3) Contents of contract.—
11	"(A) In general.—A contract entered into
12	with an entity under the competition conducted
13	pursuant to paragraph (1) is subject to terms
14	and conditions that the Secretary may specify.
15	"(B) Term of contracts.—The Secretary
16	shall recompete contracts under this section not
17	less often than once every 3 years.
18	"(4) Limit on number of contractors.—
19	"(A) In General.—The Secretary may
20	limit the number of contractors in a competitive
21	acquisition area to the number needed to meet
22	projected demand for items and services covered
23	under the contracts. In awarding contracts, the
24	Secretary shall take into account the ability of

bidding entities to furnish items or services in



1	sufficient quantities to meet the anticipated
2	needs of beneficiaries for such items or services
3	in the geographic area covered under the contract
4	on a timely basis.
5	"(B) Multiple winners.—The Secretary
6	shall award contracts to multiple entities sub-
7	mitting bids in each area for an item or service.
8	"(5) Payment under this part for
9	competitively priced items and services described in
10	subsection (a)(2) shall be based on the bids submitted
11	and accepted under this section for such items and
12	services.
13	"(6) Participating contractors.—Payment
14	shall not be made for items and services described in
15	subsection (a)(2) furnished by a contractor and for
16	which competition is conducted under this section
17	unless—
18	"(A) the contractor has submitted a bid for
19	such items and services under this section; and
20	"(B) the Secretary has awarded a contract
21	to the contractor for such items and services
22	under this section.
23	In this section, the term 'bid' means a request for a
24	proposal for an item or service that includes the cost
25	of the item or service, and where appropriate, any



1	services that are attendant to the provision of the
2	item or service.
3	"(7) Consideration in Determining Cat-
4	EGORIES FOR BIDS.—The Secretary shall consider the
5	similarity of the clinical efficiency and value of spe-
6	cific codes and products, including products that may
7	provide a therapeutic advantage to beneficiaries, be-
8	fore delineating the categories and products that will
9	be subject to bidding.
10	"(8) Authority to contract for education,
11	MONITORING, OUTREACH AND COMPLAINT SERV-
12	ICES.—The Secretary may enter into a contract with
13	an appropriate entity to address complaints from
14	beneficiaries who receive items and services from an
15	entity with a contract under this section and to con-
16	duct appropriate education of and outreach to such
17	beneficiaries and monitoring quality of services with
18	respect to the program.
19	"(c) Program Advisory and Oversight Com-
20	MITTEE.—
21	"(1) Establishment.—There is established a
22	Program Advisory and Oversight Committee (herein-
23	after in this section referred to as the 'Committee').
24	"(2) Membership; terms.—The Committee

shall consist of such members as the Secretary may



1	appoint who shall serve for such term as the Secretary
2	may specify.
3	"(3) Duties.—
4	"(A) TECHNICAL ASSISTANCE.—The Com-
5	mittee shall provide advice and technical assist-
6	ance to the Secretary with respect to the fol-
7	lowing functions:
8	"(i) The implementation of the pro-
9	gram under this section.
10	"(ii) The establishment of requirements
11	for collection of data.
12	"(iii) The development of proposals for
13	efficient interaction among manufacturers
14	and distributors of the items and services
15	and providers and beneficiaries.
16	"(B) Additional duties.—The Committee
17	shall perform such additional functions to assist
18	the Secretary in carrying out this section as the
19	Secretary may specify.
20	"(4) Inapplicability of faca.—The provisions
21	of the Federal Advisory Committee Act (5 U.S.C.
22	App.) shall not apply.
23	"(d) Annual Reports.—The Secretary shall submit
24	to Congress an annual management report on the programs
25	under this section. Each such report shall include informa-



1	tion on savings, reductions in beneficiary cost-sharing, ac-
2	cess to and quality of items and services, and beneficiary
3	satisfaction.
4	"(e) Demonstration Project for Clinical Lab-
5	ORATORY SERVICES.—
6	"(1) In General.—The Secretary shall conduct
7	a demonstration project on the application of com-
8	petitive acquisition under this section to clinical di-
9	agnostic laboratory tests—
10	"(A) for which payment is otherwise made
11	under section 1833(h) or 1834(d)(1) (relating to
12	colorectal cancer screening tests); and
13	"(B) which are furnished by entities that
14	did not have a face-to-face encounter with the in-
15	dividual.
16	"(2) Terms and conditions.—Such project
17	shall be under the same conditions as are applicable
18	to items and services described in subsection (a)(2).
19	"(3) Report.—The Secretary shall submit to
20	Congress—
21	"(A) an initial report on the project not
22	later than December 31, 2005; and
23	"(B) such progress and final reports on the
24	project after such date as the Secretary deter-
25	mines appropriate.".



1	(b) Conforming Amendments.—
2	(1) Durable medical equipment; elimi-
3	NATION OF INHERENT REASONABLENESS AUTHOR-
4	ITY.—Section $1834(a)$ $(42\ U.S.C.\ 1395m(a))$ is
5	amended—
6	(A) in paragraph $(1)(B)$ , by striking "The
7	payment basis" and inserting "Subject to sub-
8	$paragraph\ (E)(i),\ the\ payment\ basis";$
9	(B) in paragraph (1)(C), by striking "This
10	subsection" and inserting "Subject to subpara-
11	$graph\ (E)(ii),\ this\ subsection";$
12	(C) by adding at the end of paragraph (1)
13	the following new subparagraph:
14	"(E) Application of competitive acqui-
15	SITION; ELIMINATION OF INHERENT REASON-
16	ABLENESS AUTHORITY.—In the case of covered
17	items and services that are included in a com-
18	petitive acquisition program in a competitive ac-
19	quisition area under section 1847(a)—
20	"(i) the payment basis under this sub-
21	section for such items and services furnished
22	in such area shall be the payment basis de-
23	termined under such competitive acquisition
24	program; and



1	"(ii) the Secretary may use informa-
2	tion on the payment determined under such
3	competitive acquisition programs to adjust
4	the payment amount otherwise recognized
5	under subparagraph (B)(ii) for an area
6	that is not a competitive acquisition area
7	under section 1847 and in the case of such
8	adjustment, paragraph (10)(B) shall not be
9	applied."; and
10	(D) in paragraph $(10)(B)$ , by inserting "in
11	an area and with respect to covered items and
12	services for which the Secretary does not make a
13	payment amount adjustment under paragraph
14	(1)(E)" after "under this subsection".
15	(2) Off-the-shelf orthotics; elimination
16	of inherent reasonableness authority.—Sec-
17	tion 1834(h) (42 U.S.C. 1395m(h)) is amended—
18	(A) in paragraph (1)(B), by striking "and
19	(E)" and inserting ", (E) , and (H)(i)";
20	(B) in paragraph (1)(D), by striking "This
21	subsection" and inserting "Subject to subpara-
22	$graph\ (H)(ii),\ this\ subsection";$
23	(C) by adding at the end of paragraph (1)
24	the following new subparagraph:



1	"(H) Application of competitive acqui-
2	SITION TO ORTHOTICS; ELIMINATION OF INHER-
3	ENT REASONABLENESS AUTHORITY.—In the case
4	of orthotics described in paragraph $(2)(B)$ of sec-
5	tion 1847(a) that are included in a competitive
6	acquisition program in a competitive acquisition
7	area under such section—
8	"(i) the payment basis under this sub-
9	section for such orthotics furnished in such
10	area shall be the payment basis determined
11	under such competitive acquisition pro-
12	gram; and
13	"(ii) the Secretary may use informa-
14	tion on the payment determined under such
15	competitive acquisition programs to adjust
16	the payment amount otherwise recognized
17	under subparagraph (B)(ii) for an area
18	that is not a competitive acquisition area
19	under section 1847, and in the case of such
20	adjustment, paragraphs (8) and (9) of sec-
21	tion 1842(b) shall not be applied.".
22	(c) Report on Activities of Suppliers.—The Sec-
23	retary shall conduct a study to determine the extent to
24	which (if any) suppliers of covered items of durable medical
25	equipment that are subject to the competitive acquisition



1	program under section 1847 of the Social Security Act, as
2	amended by subsection (a), are soliciting physicians to pre-
3	scribe certain brands or modes of delivery of covered items
4	based on profitability.
5	SEC. 303. COMPETITIVE ACQUISITION OF COVERED OUT-
6	PATIENT DRUGS AND BIOLOGICALS.
7	(a) Adjustment to Physician Fee Schedule.—
8	(1) Adjustment in practice expense rel-
9	ATIVE VALUE UNITS.—Section 1848(c)(2) (42 U.S.C.
10	1395w-4(c)(2)) is amended—
11	(A) in subparagraph (B)—
12	(i) in clause (ii)(II), by striking "The
13	adjustments" and inserting "Subject to
14	clause (iv), the adjustments"; and
15	(ii) by adding at the end of subpara-
16	graph (B), the following new clause:
17	"(iv) Exception to budget neu-
18	TRALITY.—The additional expenditures at-
19	tributable to clauses (ii) and (iii) of sub-
20	paragraph (H) shall not be taken into ac-
21	count in applying clause (ii)(II) for 2005.";
22	and
23	(B) by adding at the end the following new
24	subparagraph:



1	"(H) Adjustments in practice expense
2	RELATIVE VALUE UNITS FOR 2004.—
3	"(i) In general.—As part of the an-
4	nual process of establishing the physician
5	fee schedule under subsection (b) for 2004,
6	the Secretary shall increase the practice ex-
7	pense relative value units for 2004 con-
8	sistent with clauses (ii) and (iii).
9	"(ii) Use of supplemental survey
10	DATA.—For 2004 for any specialty that
11	submitted survey data that included ex-
12	penses for the administration of drugs and
13	biologicals for which payment is made
14	under section 1842(o) (or section 1847A),
15	the Secretary shall use such supplemental
16	survey data in carrying out this subpara-
17	graph insofar as they are collected and pro-
18	vided by entities and organizations con-
19	sistent with the criteria established by the
20	Secretary pursuant to section 212(a) of the
21	Medicare, Medicaid, and SCHIP Balanced
22	Budget Refinement Act of 1999 and insofar
23	as such data are submitted to the Secretary
24	by the date of the enactment of this sub-
25	paragraph.



1	"(iii) Expediting consideration of
2	CPT CODES FOR AFFECTED PHYSICIAN SPE-
3	CIALTIES.—The Secretary shall, in coopera-
4	tion with representatives of physician speci-
5	alities affected by section 1847A, take such
6	actions as are necessary to expedite consid-
7	erations of CPT codes, or expand the ability
8	to appropriately bill for physicians' services
9	under existing CPT codes, for costs associ-
10	ated with the administration of covered out-
11	patient drugs. The Secretary shall consult
12	with representatives of advisory physician
13	groups in expediting such considerations.
14	"(iv) Subsequent, budget neutral
15	ADJUSTMENTS PERMITTED.—Nothing in
16	this subparagraph shall be construed as pre-
17	venting the Secretary from providing for
18	adjustments in practice expense relative
19	value units under (and consistent with)
20	subparagraph (B) for years after 2004.
21	"(v) Consultation.—Before pub-
22	lishing the notice of proposed rulemaking to
23	carry out this subparagraph, the Secretary

shall consult with the Comptroller General



1	of the United States and with groups rep-
2	resenting the physician specialties involved.
3	"(vi) Treatment as change in law
4	AND REGULATION IN SUSTAINABLE GROWTH
5	RATE DETERMINATION.—The enactment of
6	subparagraph (B)(iv) and this subpara-
7	graph shall be treated as a change in law
8	for purposes of applying subsection
9	(f)(2)(D).".
10	(2) Prohibition of administrative and judi-
11	CIAL REVIEW.—Section 1848(i)(1) (42 U.S.C. 1395w-
12	4(i)(1)) is amended—
13	(A) by striking "and" at the end of subpara-
14	graph(D);
15	(B) by striking the period at the end of subpara-
16	graph (E) and inserting ", and"; and
17	(C) by adding at the end the following new sub-
18	paragraph:
19	"(F) adjustments in practice expense rel-
20	ative value units for 2005 under subsection
21	(c)(2)(H).".
22	(3) Treatment of other services cur-
23	RENTLY IN THE NON-PHYSICIAN WORK POOL.—The
24	Secretary shall make adjustments to the non-physi-
25	cian work pool methodology (as such term is used in



1	the regulations promulgated by the Secretary in the
2	Federal Register as of December 31, 2002) for deter-
3	mination of practice expense relative value units
4	under the physician fee schedule described in section
5	1848(c)(2)(C)(ii) of the Social Security Act so that
6	the practice expense relative value units for services
7	determined under such methodology are not dis-
8	proportionately reduced relative to the practice ex-
9	pense relative value units of other services not deter-
10	mined under such non-physician work pool method-
11	ology, as the result of amendments made by para-
12	graph (1).
13	(4) Submission of practice expense survey
14	DATA.—Any physician specialty may submit survey
15	data related to practice expenses to the Secretary
16	through Decmeber 31, 2004. Nothing in this para-
17	graph shall be construed as waiving the application
18	of budget neutrality under section 1848 of the Social
19	Security Act.
20	(b) Payment Based on Competition.—Title XVIII
21	is amended by inserting after section 1847 (42 U.S.C.
22	1395w-3), as amended by section 302, the following new



23 sections:

1	"COMPETITIVE ACQUISITION OF COVERED OUTPATIENT
2	DRUGS AND BIOLOGICALS
3	"Sec. 1847A. (a) Implementation of Competitive
4	Acquisition.—
5	"(1) Implementation of program.—
6	"(A) In General.—The Secretary shall es-
7	tablish and implement a competitive acquisition
8	program under which—
9	"(i) competitive acquisition areas are
10	established throughout the United States for
11	contract award purposes for acquisition of
12	and payment for categories of covered out-
13	patient drugs and biologicals (as defined in
14	paragraph (2)) under this part; and
15	"(ii) each physician who does not elect
16	section 1847B to apply makes an annual
17	selection, under paragraph (5) of the con-
18	tractor through which drugs and biologicals
19	within a category of drugs and biologicals
20	will be acquired and delivered to the physi-
21	cian under this part.
22	"(B) Implementation.—The Secretary
23	shall implement the program so that the pro-
24	gram applies to—



1	"(i) the oncology category beginning in
2	2005; and
3	"(ii) the non-oncology category begin-
4	ning in 2006.
5	This section shall not apply in the case of a phy-
6	sician who elects section 1847B to apply.
7	"(C) Waiver of Certain Provisions.—In
8	order to promote competition, efficient service,
9	and product quality, in carrying out the pro-
10	gram the Secretary may waive such provisions of
11	the Federal Acquisition Regulation as are nec-
12	essary for the efficient implementation of this
13	section, other than provisions relating to con-
14	fidentiality of information and such other provi-
15	sions as the Secretary determines appropriate.
16	"(D) Exclusion authority.—The Sec-
17	retary may exclude covered outpatient drugs and
18	biologicals (including a class of such drugs and
19	biologicals) from the competitive bidding system
20	under this section if the drugs or biologicals (or
21	class) are not appropriate for competitive bid-
22	ding due to low volume of utilization by bene-
23	ficiaries under this part or a unique mode or

method of delivery or similar reasons.



1	"(2) Covered outpatient drugs and
2	BIOLOGICALS, CATEGORIES, PROGRAM DEFINED.—For
3	purposes of this section—
4	"(A) Covered outpatient drugs and
5	BIOLOGICALS DEFINED.—The term 'covered out-
6	patient drugs and biologicals' means drugs and
7	biologicals to which section 1842(o) applies and
8	which are not covered under section 1847 (relat-
9	ing to competitive acquisition for items of dura-
10	ble medical equipment). Such term does not in-
11	clude the following:
12	"(i) Blood clotting factors.
13	"(ii) Drugs and biologicals furnished
14	to individuals in connection with the treat-
15	ment of end stage renal disease.
16	$\it ``(iii) \ Radiopharmac euticals.$
17	"(B) 2 CATEGORIES.—Each of the following
18	shall be a separate category of covered outpatient
19	drugs and biologicals, as identified by the Sec-
20	retary:
21	"(i) Oncology category.—A cat-
22	egory (in this section referred to as the 'on-
23	cology category') consisting of those covered
24	outpatient drugs and biologicals that, as de-
25	termined by the Secretary, are typically



1	primarily billed by oncologists or are other-
2	wise used to treat cancer.
3	"(ii) Non-oncology categories.—
4	Such numbers of categories (in this section
5	referred to as the 'non-oncology categories')
6	consisting of covered outpatient drugs and
7	biologicals not described in clause (i), and
8	appropriate subcategories of such drugs and
9	biologicals as the Secretary may specify.
10	"(C) Program.—The term 'program'
11	means the competitive acquisition program
12	under this section.
13	"(D) Competitive acquisition area;
14	AREA.—The terms 'competitive acquisition area'
15	and 'area' mean an appropriate geographic re-
16	gion established by the Secretary under the pro-
17	gram.
18	"(E) Contractor.—The term 'contractor'
19	means an entity that has entered into a contract
20	with the Secretary under this section.
21	"(3) Application of program payment meth-
22	odology.—With respect to covered outpatient drugs
23	and biologicals which are supplied under the program
24	in an area and which are prescribed by a physician

who has not elected section 1847B to apply—



1	"(A) the claim for such drugs and
2	biologicals shall be submitted by the contractor
3	that supplied the drugs and biologicals;
4	"(B) collection of amounts of any deductible
5	and coinsurance applicable with respect to such
6	drugs and biologicals shall be the responsibility
7	of such contractor and shall not be collected un-
8	less the drug or biological is administered to the
9	beneficiary involved; and
10	"(C) the payment under this section (and
11	related coinsurance amounts) for such drugs and
12	biologicals—
13	"(i) shall be made only to such con-
14	tractor;
15	"(ii) shall be conditioned upon the ad-
16	ministration of such drugs and biologicals;
17	and
18	"(iii) shall be based on the average of
19	the bid prices for such drugs and biologicals
20	in the area, as computed under subsection
21	(d).
22	The Secretary shall provide a process for
23	recoupment in the case in which payment is
24	made for drugs and biologicals which were billed



1	at the time of dispensing but which were not ac-
2	tually administered.
3	"(4) Contract required.—
4	"(A) In general.—Payment may not be
5	made under this part for covered outpatient
6	drugs and biologicals prescribed by a physician
7	who has not elected section 1847B to apply with-
8	in a category and a competitive acquisition area
9	with respect to which the program applies
10	unless—
11	"(i) the drugs or biologicals are sup-
12	plied by a contractor with a contract under
13	this section for such category of drugs and
14	biologicals and area; and
15	"(ii) the physician has elected such
16	contractor under paragraph (5) for such
17	category and area.
18	"(B) Physician choice.—Subparagraph
19	(A) shall not apply for a category of drugs for
20	an area if the physician prescribing the covered
21	outpatient drug in such category and area has
22	elected to apply section 1847B instead of this
23	section.
24	"(5) Contractor selection process.—



1	"(A) In General.—The Secretary shall
2	provide a process for the selection of a con-
3	tractor, on an annual basis and in such exigent
4	circumstances as the Secretary may provide and
5	with respect to each category of covered out-
6	patient drugs and biologicals for an area, by
7	physicians prescribing such drugs and
8	biologicals in the area of the contractor under
9	this section that will supply the drugs and
10	biologicals within that category and area. Such
11	selection shall also include the election described
12	in section $1847B(a)$ .
13	"(B) Information on contractors.—The
14	Secretary shall make available to physicians on
15	an ongoing basis, through a directory posted on
16	the Department's Internet website or otherwise
17	and upon request, a list of the contractors under
18	this section in the different competitive acquisi-
19	tion areas.
20	"(C) Selecting physician defined.—For
21	purposes of this section, the term 'selecting phy-
22	sician' means, with respect to a contractor and
23	category and competitive acquisition area, a

 $physician\ who\ has\ not\ elected\ section\ 1847B\ to$ 



1	apply and has selected to apply under this sec-
2	tion such contractor for such category and area.
3	"(b) Program Requirements.—
4	"(1) Contract for covered outpatient
5	DRUGS AND BIOLOGICALS.—The Secretary shall con-
6	duct a competition among entities for the acquisition
7	of a covered outpatient drug or biological within each
8	HCPCS code within each category for each competi-
9	tive acquisition area.
10	"(2) Conditions for awarding contract.—
11	"(A) In general.—The Secretary may not
12	award a contract to any entity under the com-
13	petition conducted in a competitive acquisition
14	area pursuant to paragraph (1) with respect to
15	the acquisition of covered outpatient drugs and
16	biologicals within a category unless the Secretary
17	finds that the entity meets all of the following
18	with respect to the contract period involved:
19	"(i) Capacity to supply covered
20	OUTPATIENT DRUG OR BIOLOGICAL WITHIN
21	CATEGORY.—
22	"(I) In general.—The entity has
23	sufficient arrangements to acquire and
24	to deliver covered outpatient drugs and
25	biologicals within such category in the



1	area specified in the contract at the bid
2	price specified in the contract for all
3	physicians that may elect such entity.
4	"(II) Shipment method-
5	OLOGY.—The entity has arrangements
6	in effect for the shipment at least 5
7	days each week of covered outpatient
8	drugs and biologicals under the con-
9	tract and for the timely delivery (in-
10	cluding for emergency situations) of
11	such drugs and biologicals in the area
12	under the contract.
13	"(ii) Quality, service, financial
14	PERFORMANCE AND SOLVENCY STAND-
15	ARDS.—The entity meets quality, service, fi-
16	nancial performance, and solvency stand-
17	ards specified by the Secretary, including—
18	``(I) the establishment of proce-
19	dures for the prompt response and res-
20	olution of physician and beneficiary
21	complaints and inquiries regarding the
22	shipment of covered outpatient drugs
23	and biologicals; and
24	"(II) a grievance process for the
25	resolution of disputes.



1	"(B) Additional considerations.—The
2	Secretary may refuse to award a contract under
3	this section, and may terminate such a contract,
4	with an entity based upon—
5	"(i) the suspension or revocation, by
6	the Federal Government or a State govern-
7	ment, of the entity's license for the distribu-
8	tion of drugs or biologicals (including con-
9	$trolled\ substances);\ or$
10	"(ii) the exclusion of the entity under
11	section 1128 from participation under this
12	title.
13	"(C) Application of medicare provider
14	OMBUDSMAN.—For provision providing for a
15	program-wide Medicare Provider Ombudsman to
16	review complaints, see section 1868(b), as added
17	by section 923 of the Medicare Prescription Drug
18	and Modernization Act of 2003.
19	"(3) Awarding multiple contracts for a
20	CATEGORY AND AREA.—In order to provide a choice
21	of at least 2 contractors in each competitive acquisi-
22	tion area for a category of drugs and biologicals, the
23	Secretary may limit (but not below 2) the number of
24	qualified entities that are awarded such contracts for



1	any category and area. The Secretary shall select
2	among qualified entities based on the following:
3	"(A) The bid prices for covered outpatient
4	drugs and biologicals within the category and
5	area.
6	"(B) Bid price for distribution of such
7	drugs and biologicals.
8	"(C) Ability to ensure product integrity.
9	"(D) Customer service.
10	"(E) Past experience in the distribution of
11	drugs and biologicals, including controlled sub-
12	stances.
13	"(F) Such other factors as the Secretary
14	may specify.
15	"(4) Terms of contracts.—
16	"(A) In general.—A contract entered into
17	with an entity under the competition conducted
18	pursuant to paragraph (1) is subject to terms
19	and conditions that the Secretary may specify
20	consistent with this section.
21	"(B) Period of contracts.—A contract
22	under this section shall be for a term of 2 years,
23	but may be terminated by the Secretary or the
24	entity with appropriate, advance notice.



1	"(C) Integrity of drug and biological
2	DISTRIBUTION SYSTEM.—The Secretary—
3	"(i) shall require that for all drug and
4	biological products distributed by a con-
5	tractor under this section be acquired di-
6	rectly from the manufacturer or from a dis-
7	tributor that has acquired the products di-
8	rectly from the manufacturer; and
9	"(ii) may require, in the case of such
10	products that are particularly susceptible to
11	counterfeit or diversion, that the contractor
12	comply with such additional product integ-
13	rity safeguards as may be determined to be
14	necessary.
15	"(D) Implementation of anti-counter-
16	FEITING, QUALITY, SAFETY, AND RECORD KEEP-
17	ing requirements.—The Secretary shall re-
18	quire each contractor to implement (through its
19	officers, agents, representatives, and employees)
20	requirements relating to the storage and han-
21	dling of covered outpatient drugs and biologicals
22	and for the establishment and maintenance of
23	distribution records for such drugs and
24	biologicals. A contract under this section may
25	include requirements relating to the following:



1	"(i) Secure facilities.
2	"(ii) Safe and appropriate storage of
3	drugs and biologicals.
4	"(iii) Examination of drugs and
5	biologicals received and dispensed.
6	"(iv) Disposition of damaged and out-
7	dated drugs and biologicals.
8	"(v) Record keeping and written poli-
9	cies and procedures.
10	"(vi) Compliance personnel.
11	"(E) Compliance with code of conduct
12	AND FRAUD AND ABUSE RULES.—Under the
13	contract—
14	"(i) the contractor shall comply with a
15	code of conduct, specified or recognized by
16	the Secretary, that includes standards relat-
17	ing to conflicts of interest; and
18	"(ii) the contractor shall comply with
19	all applicable provisions relating to preven-
20	tion of fraud and abuse, including compli-
21	ance with applicable guidelines of the De-
22	partment of Justice and the Inspector Gen-
23	eral of the Department of Health and
24	Human Services.



1	"(F) Direct delivery of drugs and
2	BIOLOGICALS TO PHYSICIANS.—Under the con-
3	tract the contractor shall only supply covered
4	outpatient drugs and biologicals directly to the
5	selecting physicians and not directly to bene-
6	ficiaries, except under circumstances and settings
7	where a beneficiary currently receives a drug or
8	biological in the beneficiary's home or other non-
9	physician office setting as the Secretary may
10	provide. The contractor shall not deliver drugs
11	and biologicals to a selecting physician except
12	upon receipt of a prescription for such drugs and
13	biologicals, and such necessary data as may be
14	required by the Secretary to carry out this sec-
15	tion. This section does not require a physician to
16	submit a prescription for each individual treat-
17	ment and does not change the physician's flexi-
18	bility in terms of writing a prescription for
19	drugs for a single treatment or a course of treat-
20	ment.
21	"(5) Permitting access to drugs and
22	BIOLOGICALS.—The Secretary shall establish rules
23	under this section under which drugs and biologicals
24	which are acquired through a contractor under this

section may be used to resupply inventories of such



1	drugs and biologicals which are administered con-
2	sistent with safe drug practices and with adequate
3	safeguards against fraud and abuse. The previous sen-
4	tence shall apply—
5	"(A) in cases in which the drugs or
6	biologicals are immediately required;
7	"(B) in cases in which the physician could
8	not have reasonably anticipated the immediate
9	requirement for the drugs or biologicals;
10	"(C) in cases in which the contractor could
11	not deliver to the physician the drugs or
12	biologicals in a timely manner; and
13	"(D) in emergency situations.
14	"(6) Construction.—Nothing in this section
15	shall be construed as waiving applicable State re-
16	quirements relating to licensing of pharmacies.
17	"(c) Bidding Process.—
18	"(1) In general.—In awarding a contract for
19	a category of drugs and biologicals in an area under
20	the program, the Secretary shall consider with respect
21	to each entity seeking to be awarded a contract the
22	prices bid to acquire and supply the covered out-
23	patient drugs and biologicals for that category and
24	area and the other factors referred to in subsection
25	(b)(3).



1	"(2) Prices bid by an entity
2	under paragraph (1) shall be the prices in effect and
3	available for the supply of contracted drugs and
4	biologicals in the area through the entity for the con-
5	tract period.
6	"(3) Rejection of contract offer.—The Sec-
7	retary shall reject the contract offer of an entity with
8	respect to a category of drugs and biologicals for an
9	area if the Secretary estimates that the prices bid, in
10	the aggregate on average, would exceed 120 percent o
11	the average sales price (as determined under section
12	1847B).
13	"(4) Bidding on a national or regional
14	BASIS.—Nothing in this section shall be construed as
15	precluding a bidder from bidding for contracts in al
16	areas of the United States or as requiring a bidder
17	to submit a bid for all areas of the United States.
18	"(5) Uniformity of bids within area.—The
19	amount of the bid submitted under a contract offer for
20	any covered outpatient drug or biological for an area
21	shall be the same for that drug or biological for al
21 22	shall be the same for that drug or biological for al portions of that area.

to a bid submitted in a contract offer for a covered



1	outpatient drug or biological under this section in the
2	same manner as it applies to information disclosed
3	under such section, except that any reference—
4	"(A) in that subparagraph to a 'manufac-
5	turer or wholesaler' is deemed a reference to a
6	bidder' under this section;
7	"(B) in that section to 'prices charged for
8	drugs' is deemed a reference to a 'bid' submitted
9	under this section; and
10	"(C) in clause (i) of that section to 'this sec-
11	tion', is deemed a reference to 'part B of title
12	XVIII'.
13	"(7) Inclusion of costs.—The bid price sub-
14	mitted in a contract offer for a covered outpatient
15	drug or biological shall—
16	"(A) include all costs related to the delivery
17	of the drug or biological to the selecting physi-
18	cian (or other point of delivery); and
19	"(B) include the costs of dispensing (includ-
20	ing shipping) of such drug or biological and
21	management fees, but shall not include any costs
22	related to the administration of the drug or bio-
23	logical, or wastage, spillage, or spoilage.



1	"(8) Price adjustments during contract
2	PERIOD; DISCLOSURE OF COSTS.—Each contract
3	awarded shall provide for—
4	"(A) disclosure to the Secretary the contrac-
5	tor's reasonable, net acquisition costs for periods
6	specified by the Secretary, not more often than
7	quarterly, of the contract; and
8	"(B) appropriate price adjustments over the
9	period of the contract to reflect significant in-
10	creases or decreases in a contractor's reasonable,
11	net acquisition costs, as so disclosed.
12	"(d) Computation of Average Bid Prices for A
13	Category and Area.—
14	"(1) In general.—For each year or other con-
15	tract period for each covered outpatient drug or bio-
16	logical and area with respect to which a competition
17	is conducted under the program, the Secretary shall
18	compute an area average of the bid prices submitted,
19	in contract offers accepted for the category and area,
20	for that year or other contract period.
21	"(2) Special rules.—The Secretary shall es-
22	tablish rules regarding the use under this section of
23	the alternative payment amount provided under sec-
24	tion 1847B to the use of a price for specific covered



1	outpatient drugs and biologicals in the following
2	cases:
3	"(A) New drugs and biologicals.—A
4	covered outpatient drug or biological for which
5	an average bid price has not been previously de-
6	termined.
7	"(B) Other cases.—Such other excep-
8	tional cases as the Secretary may specify in reg-
9	ulations.
10	Such alternative payment amount shall be based
11	upon actual market price information and in no case
12	shall it exceed the average sales price (as determined
13	under section 1847B).
14	"(e) Coinsurance.—
15	"(1) In general.—Coinsurance under this part
16	with respect to a covered outpatient drug or biological
17	for which payment is payable under this section shall
18	be based on 20 percent of the payment basis under
19	this section.
20	"(2) Collection.—Such coinsurance shall be
21	collected by the contractor that supplies the drug or
22	biological involved and, subject to subsection
23	(a)(3)(B), in the same manner as coinsurance is col-
24	lected for durable medical equipment under this part.
25	"(f) Special Payment Rules.—



1	"(1) In general.—The Secretary may not pro-
2	vide for an adjustment to reimbursement for covered
3	outpatient drugs and biologicals unless adjustments to
4	the practice expense payment adjustment are made on
5	the basis of supplemental surveys under section
6	1848(c)(2)(H)(ii) of the Social Security Act, as added
7	by subsection $(a)(1)(B)$ .
8	"(B) Use in exclusion cases.—If the
9	Secretary excludes a drug or biological (or class
10	of drugs or biologicals) under subsection
11	(a)(1)(D), the Secretary may provide for reim-
12	bursement to be made under this part for such
13	drugs and biologicals (or class) using the pay-
14	ment methodology under section 1847B or other
15	market based pricing system.
16	"(2) Coordination rules.—The provisions of
17	section 1842(h)(3) shall apply to a contractor with re-
18	spect to covered outpatients drugs and biologicals sup-
19	plied by that contractor in the same manner as they
20	apply to a participating supplier. In order to admin-
21	ister this section, the Secretary may condition pay-
22	ment under this part to a person for the administra-
23	tion of a drug or biological supplied under this sec-
24	tion upon person's provision of information on such



25

administration.

1	"(3) Application of requirement for as-
2	SIGNMENT.—For provision requiring assignment of
3	claims for covered outpatient drugs and biologicals,
4	see section $1842(0)(3)$ .
5	"(4) Protection for beneficiary in case of
6	MEDICAL NECESSITY DENIAL.—For protection of bene-
7	ficiaries against liability in the case of medical neces-
8	sity determinations, see section $1842(b)(3)(B)(ii)(III)$ .
9	"(5) Physician role in appeals process.—
10	The Secretary shall establish a procedure under which
11	a physician who prescribes a drug or biological for
12	which payment is made under this section has appeal
13	rights that are similar to those provided to a physi-
14	cian who prescribes durable medical equipment or a
15	laboratory test.
16	"(g) Advisory Committee.—The Secretary shall es-
17	tablish an advisory committee that includes representatives
18	of parties affected by the program under this section, in-
19	cluding physicians, specialty pharmacies, distributors,
20	manufacturers, and beneficiaries. The committee shall ad-
21	vise the Secretary on issues relating to the effective imple-
22	mentation of this section.
23	"(h) Annual Reports.—The Secretary shall submit
24	to Congress an annual report in each of 2004, 2005, and
25	2006, on the program. Each such report shall include infor-



1	mation on savings, reductions in cost-sharing, access to cov-
2	ered outpatient drugs and biologicals, the range of choices
3	of contractors available to providers, and beneficiary and
4	provider satisfaction.
5	"OPTIONAL USE OF AVERAGE SALES PRICE PAYMENT
6	METHODOLOGY
7	"Sec. 1847B. (a) In General.—In connection with
8	the election made by a physician under section
9	1847A(a)(5), the physician may elect to apply this section
10	to the payment for covered outpatient drugs instead of the
11	payment methodology under section 1847A. For purposes
12	of this section, the term 'covered outpatient drug' has the
13	meaning given such term in section $1847A(a)(2)(A)$ .
14	"(b) Computation of Payment Amount.—
15	"(1) In general.—If this section applies with
16	respect to a covered outpatient drug, the amount pay-
17	able for the drug (based on a minimum dosage unit)
18	is, subject to applicable deductible and coinsurance—
19	"(A) in the case of a multiple source drug
20	(as defined in subsection $(c)(6)(C)$ ), the amount
21	determined under paragraph (3); or
22	"(B) in the case of a single source drug (as
23	defined in subsection $(c)(6)(D)$ , the amount de-
24	termined under paragraph (4).
25	"(2) Specification of unit



1	"(A) Specification by manufacturer.—
2	The manufacturer of a covered outpatient drug
3	shall specify the unit associated with each Na-
4	tional Drug Code as part of the submission of
5	$data\ under\ section\ 1927(b)(3)(A)(iii).$
6	"(B) Unit defined.—In this section, the
7	term 'unit' means, with respect to a covered out-
8	patient drug, the lowest identifiable quantity
9	(such as a capsule or tablet, milligram of mol-
10	ecules, or grams) of the drug that is dispensed,
11	exclusive of any diluent without reference to vol-
12	ume measures pertaining to liquids.
13	"(3) Multiple source drug.—For all drug
14	products included within the same multiple source
15	drug, the amount specified in this paragraph is the
16	volume-weighted average of the average sales prices
17	reported under section 1927(b)(3)(A)(iii) computed as
18	follows:
19	"(A) Compute the sum of the products (for
20	each national drug code assigned to such drug
21	products) of—
22	"(i) the manufacturer's average sales
23	price (as defined in subsection (c)); and



1	"(ii) the total number of units specified
2	under paragraph (2) sold, as reported under
3	section $1927(b)(3)(A)(iii)$ .
4	"(B) Divide the sum computed under sub-
5	paragraph (A) by the sum of the total number
6	of units under subparagraph (A)(ii) for all na-
7	tional drug codes assigned to such drug products.
8	"(4) Single source drug.—The amount speci-
9	fied in this paragraph for a single source drug is the
10	lesser of the following:
11	"(A) Manufacturer's average sales
12	PRICE.—The manufacturer's average sales price
13	for a national drug code, as computed using the
14	methodology applied under paragraph (3).
15	"(B) Wholesale acquisition cost
16	(WAC).—The wholesale acquisition cost (as de-
17	fined in subsection $(c)(6)(B)$ ) reported for the
18	single source drug.
19	"(5) Basis for determination.—The payment
20	amount shall be determined under this subsection
21	based on information reported under subsection (e)
22	and without regard to any special packaging, label-
23	ing, or identifiers on the dosage form or product or
24	package.
25	"(c) Manufacturer's Average Sales Price.—



1	"(1) In general.—For purposes of this sub-
2	section, subject to paragraphs (2) and (3), the manu-
3	facturer's 'average sales price' means, of a covered
4	outpatient drug for a NDC code for a calendar quar-
5	ter for a manufacturer for a unit—
6	"(A) the manufacturer's total sales (as de-
7	fined by the Secretary in regulations for pur-
8	poses of section 1927(c)(1)) in the United States
9	for such drug in the calendar quarter; divided by
10	"(B) the total number of such units of such
11	drug sold by the manufacturer in such quarter.
12	"(2) Certain sales exempted from computa-
13	TION.—In calculating the manufacturer's average
14	sales price under this subsection, the following sales
15	shall be excluded:
16	"(A) Sales exempt from best price.—
17	Sales exempt from the inclusion in the deter-
18	mination of best price' under section
19	1927(c)(1)(C)(i).
20	"(B) Sales at nominal charge.—Such
21	other sales as the Secretary identifies by regula-
22	tion as sales to an entity that are nominal in
23	price or do not reflect a market price paid by an
24	entity to which payment is made under this sec-
25	tion.



"(3) Sale price net of discounts.—In calcu-
lating the manufacturer's average sales price under
this subsection, such price shall be determined taking
into account volume discounts, prompt pay discounts,
cash discounts, the free goods that are contingent on
any purchase requirement, chargebacks, and rebates
(other than rebates under section 1927), that result in
a reduction of the cost to the purchaser. A rebate to
a payor or other entity that does not take title to a
covered outpatient drug shall not be taken into ac-
count in determining such price unless the manufac-
turer has an agreement with the payor or other entity
under which the purchaser's price for the drug is re-
duced as a consequence of such rebate.

"(4) AUTHORITY TO DISREGARD AVERAGE SALES
PRICE DURING FIRST QUARTER OF SALES.—In the
case of a covered outpatient drug during an initial
period (not to exceed a full calendar quarter) in
which data on the prices for sales for the drug is not
sufficiently available from the manufacturer to compute an average sales price for the drug, the Secretary
may determine the amount payable under this section
for the drug without considering the manufacturer's
average sales price of that manufacturer for that
drug.



1	"(5) Frequency of Determinations.—
2	"(A) In General on a quarterly
3	BASIS.—The manufacturer's average sales price,
4	for a covered outpatient drug of a manufacturer,
5	shall be determined by such manufacturer under
6	this subsection on a quarterly basis. In making
7	such determination insofar as there is a lag in
8	the reporting of the information on rebates and
9	chargebacks under paragraph (3) so that ade-
10	quate data are not available on a timely basis,
11	the manufacturer shall apply a methodology es-
12	tablished by the Secretary based on a 12-month
13	rolling average for the manufacturer to estimate
14	costs attributable to rebates and chargebacks.
15	"(B) UPDATES IN RATES.—The payment
16	rates under subsection (b)(1) and (b)(2)(A) shall
17	be updated by the Secretary on a quarterly basis
18	and shall be applied based upon the manufactur-
19	er's average sales price determined for the most
20	recent calendar quarter.
21	"(C) Use of contractors; implementa-
22	TION.—The Secretary may use a carrier, fiscal
23	intermediary, or other contractor to determine
24	the payment amount under subsection (b). Not-

withstanding any other provision of law, the



1	Secretary may implement, by program memo-
2	randum or otherwise, any of the provisions of
3	this section.
4	"(6) Definitions and other rules.—In this
5	section:
6	"(A) Manufacturer.—The term 'manufac-
7	turer' means, with respect to a covered out-
8	patient drug, the manufacturer (as defined in
9	section 1927(k)(5)) whose national drug code ap-
10	pears on such drug.
11	"(ii) Wholesale acquisition cost.—The
12	term 'wholesale acquisition cost' means, with re-
13	spect to a covered outpatient drug, the manufac-
14	turer's list price for the drug to wholesalers or
15	direct purchasers in the United States, not in-
16	cluding prompt pay or other discounts, rebates
17	or reductions in price, for the most recent month
18	for which the information is available, as re-
19	ported in wholesale price guides or other publica-
20	tions of drug pricing data.
21	"(C) Multiple source drug.—The term
22	'multiple source drug' means, for a calendar
23	quarter, a covered outpatient drug for which

there are 2 or more drug products which—



1	"(i) are rated as therapeutically equiv-
2	alent (under the Food and Drug Adminis-
3	tration's most recent publication of 'Ap-
4	proved Drug Products with Therapeutic
5	$Equivalence\ Evaluations'),$
6	"(ii) except as provided in subpara-
7	graph (E), are pharmaceutically equivalent
8	and bioequivalent, as determined under sub-
9	paragraph (F) and as determined by the
10	Food and Drug Administration, and
11	"(iii) are sold or marketed in the
12	United States during the quarter.
13	"(D) Single source drug.—The term
14	'single source drug' means a covered outpatient
15	drug which is not a multiple source drug and
16	which is produced or distributed under an origi-
17	nal new drug application approved by the Food
18	and Drug Administration, including a drug
19	product marketed by any cross-licensed pro-
20	ducers or distributors operating under the new
21	drug application, or which is a biological.
22	"(E) Exception from pharmaceutical
23	EQUIVALENCE AND BIOEQUIVALENCE REQUIRE-
24	MENT.—Subparagraph (C)(ii) shall not apply if

the Food and Drug Administration changes by



1	regulation the requirement that, for purposes of
2	the publication described in subparagraph $(C)(i)$ ,
3	in order for drug products to be rated as thera-
4	peutically equivalent, they must be pharmaceuti-
5	cally equivalent and bioequivalent, as defined in
6	subparagraph (F).
7	"(F) Determination of pharmaceutical
8	EQUIVALENCE AND BIOEQUIVALENCE.—For pur-
9	poses of this paragraph—
10	"(i) drug products are pharmaceuti-
11	cally equivalent if the products contain
12	identical amounts of the same active drug
13	ingredient in the same dosage form and
14	meet compendial or other applicable stand-
15	ards of strength, quality, purity, and iden-
16	tity; and
17	"(ii) drugs are bioequivalent if they do
18	not present a known or potential bioequiva-
19	lence problem, or, if they do present such a
20	problem, they are shown to meet an appro-
21	priate standard of bioequivalence.
22	"(G) Inclusion of vaccines.—In apply-
23	ing provisions of section 1927 under this section,
24	'other than a vaccine' is deemed deleted from sec-
25	$tion \ 1927(k)(2)(B).$



1	"(d) Monitoring price information.—
2	"(1) In general.—The Secretary shall monitor
3	available pricing information, including information
4	on average sales price and average manufacturer
5	price.
6	"(2) Response to significant discrep-
7	ANCIES.—
8	"(A) Report to congress.—If the Sec-
9	retary finds that there are significant discrep-
10	ancies among such prices and that the manufac-
11	turer's average sales price does not reflect a
12	broad-based market price or a reasonable ap-
13	proximation of the acquisition cost of the covered
14	outpatient drug involved to purchasers reim-
15	bursed under this section, the Secretary shall
16	submit to Congress a report.
17	"(B) Confidentiality of information
18	Reported.—Consistent with requirements relat-
19	ing to maintaining the confidentiality of infor-
20	mation reported on manufacturer's average
21	prices under section 1927(b)(3)(D), such report
22	shall include details regarding such discrepancies
23	and recommendations on how to best address

such discrepancies. Such report shall not disclose



1	average manufacturer prices or average sales
2	prices.
3	"(C) RECOMMENDATIONS.—Such rec-
4	ommendations may include other changes in
5	$payment\ methodology.$
6	"(D) AUTHORITY TO MODIFY PAYMENT
7	METHODOLOGY BY RULE.—Upon submission of
8	such report, the Secretary may commence a rule-
9	making to change such percent or payment meth-
10	odologies under paragraph (1)(D) and (2) as ap-
11	plied to the covered outpatient drug involved
12	under this section.
13	"(3) Response to public health emer-
14	GENCY.—In the case of a public health emergency
15	under section 319 of the Public Health Service Act in
16	which there is a documented inability to access cov-
17	ered outpatient drugs, and a concomitant increase in
18	the price, of a drug which is not reflected in the man-
19	ufacturer's average sales price for one or more quar-
20	ters, the Secretary may use the wholesale acquisition
21	cost (or other reasonable measure of drug price) in-
22	stead of the manufacturer's average sales price for
23	such quarters and for subsequent quarters until the

price and availability of the drug has stabilized and



1	is substantially reflected in the applicable manufac-
2	turer's average sales price.
3	"(4) Annual report to congress.—The Sec-
4	retary shall submit to the Committees on Energy and
5	Commerce and Ways and Means of the House of Rep-
6	resentatives and the Committee on Finance of the
7	Senate an annual report on the operation of this sec-
8	tion. Such report shall be submitted in coordination
9	with the submission of reports under section 1927(i).
10	Such report shall include information on the fol-
11	lowing:
12	"(A) Trends in average sales price under
13	subsection (b).
14	"(B) Administrative costs associated with
15	compliance with this section.
16	"(C) Total value of payments made under
17	this section.
18	"(D) Comparison of the average manufac-
19	turer price as applied under section 1927 for a
20	covered outpatient drug with the manufacturer's
21	average sales price for the drug under this sec-
22	tion.
23	"(e) Reports on pricing information.—
24	"(1) Reference to reporting requirement
25	ON AVERAGE SALES PRICE.—For requirements for re-



1	porting the manufacturer's average sales price (and,
2	if required to make payment, the manufacturer's
3	wholesale acquisition cost) for the covered outpatient
4	drug, see section 1927(b)(3).
5	"(2) MedPAC review.—The Medicare Payment
6	Advisory Commission shall periodically review the
7	payment methodology established under this section
8	and submit to Congress such recommendations on
9	such methodology as it deems appropriate as part of
10	its annual reports to Congress.
11	"(3) Construction.—Nothing in this subsection
12	shall be construed as authorizing the Secretary to re-
13	view for purposes of this section information reported
14	only under section $1927(b)(3)$ .
15	"(f) Restriction on administrative and judicial
16	REVIEW.—There shall be no administrative or judicial re-
17	view under section 1869, section 1878, or otherwise, of de-
18	terminations of manufacturer's average sales price under
19	subsection (c).".
20	(c) Continuation of Payment Methodology for
21	$Radiopharmaceuticals. \color{red} -Nothing \ in \ the \ amendments$
22	made by this section shall be construed as changing the pay-
23	$ment\ methodology\ under\ part\ B\ of\ title\ XVIII\ of\ the\ Social$
24	Security Act for radiopharmaceuticals, including the use by

25 carriers of invoice pricing methodology.



1	(d) Conforming Amendments.—
2	(1) In General.—Section 1842(o) (42 U.S.C.
3	1395u(o)) is amended—
4	(A) in paragraph (1), by inserting ", sub-
5	ject to section 1847A and 1847B," before "the
6	amount payable for the drug or biological"; and
7	(B) by adding at the end of paragraph (2)
8	the following: "This paragraph shall not apply
9	in the case of payment under section 1847A or
10	1847B.".
11	(2) No change in coverage basis.—Section
12	1861(s)(2)(A) (42 U.S.C. $1395x(s)(2)(A)$ ) is amended
13	by inserting "(or would have been so included but for
14	the application of section 1847A or 1847B)" after
15	"included in the physicians' bills".
16	(3) Payment.—Section $1833(a)(1)(S)$ (42)
17	$U.S.C.\ 1395l(a)(1)(S))$ is amended by inserting "(or,
18	if applicable, under section 1847A or 1847B)" after
19	"1842(o)".
20	(4) Consolidated reporting of pricing in-
21	FORMATION.—Section 1927 (42 U.S.C. 1396r-8) is
22	amended—
23	(A) in subsection (a)(1), by inserting "or
24	under part B of title XVIII' after "section
25	1903(a)";



1	(B) in subsection $(b)(3)(A)$ —
2	(i) in clause (i), by striking "and" at
3	$the\ end;$
4	(ii) in clause (ii), by striking the pe-
5	riod and inserting "; and"; and
6	(iii) by adding at the end the following
7	new clause:
8	"(iii) for calendar quarters beginning
9	on or after April 1, 2004, in conjunction
10	with reporting required under clause (i)
11	and by national drug code (NDC)—
12	"(I) the manufacturer's average
13	sales price (as defined in section
14	1847B(c)) and the total number of
15	units specified under section
16	1847B(b)(2)(A);
17	"(II) if required to make payment
18	under section 1847B, the manufactur-
19	er's wholesale acquisition cost, as de-
20	fined in subsection $(c)(6)$ of such sec-
21	$tion; \ and$
22	"(III) information on those sales
23	that were made at a nominal price or
24	otherwise described in section
25	1847B(c)(2)(B), which information is



1	subject to audit by the Inspector Gen-
2	eral of the Department of Health and
3	Human Services;
4	for a covered outpatient drug for which
5	payment is made under section 1847B.";
6	(C) in subsection $(b)(3)(B)$ —
7	(i) in the heading, by inserting "AND
8	MANUFACTURER'S AVERAGE SALES PRICE"
9	after "PRICE"; and
10	(ii) by inserting "and manufacturer's
11	average sales prices (including wholesale ac-
12	quisition cost) if required to make pay-
13	ment" after "manufacturer prices"; and
14	(D) in subsection $(b)(3)(D)(i)$ , by inserting
15	"and section 1847B" after "this section".
16	(e) GAO STUDY.—
17	(1) Study.—The Comptroller General of the
18	United States shall conduct a study to assess the im-
19	pact of the amendments made by this section on the
20	delivery of services, including their impact on—
21	(A) beneficiary access to drugs and
22	biologicals for which payment is made under
23	part B of title XVIII of the Social Security Act;
24	and
25	(B) the site of delivery of such services.



1	(2) Report.—Not later than 2 years after the
2	year in which the amendment made by subsection
3	(a)(1) first takes effect, the Comptroller General shall
4	submit to Congress a report on the study conducted
5	under paragraph (1).
6	(f) MedPAC Recommendations on Blood Clot-
7	TING FACTORS.—The Medicare Payment Advisory Commis-
8	sion shall submit to Congress, in its annual report in 2004,
9	specific recommendations regarding a payment amount (or
10	amounts) for blood clotting factors and its administration
11	under the medicare program.
12	(g) Establishment of Pharmaceutical Manage-
13	MENT FEE WHERE DRUGS PROVIDED THROUGH A CON-
14	TRACTOR.—Section 1848(a) (42 U.S.C. 1395w-4(a)) is
15	amended by adding at the end the following new paragraph:
16	"(5) Recognition of pharmaceutical man-
17	AGEMENT FEE IN CERTAIN CASES.—In establishing
18	the fee schedule under this section, the Secretary shall
19	provide for a separate payment with respect to physi-
20	cians' services consisting of the unique administrative
21	and management costs associated with covered drugs
22	and biologicals which are furnished to physicians
23	through a contractor under section 1847A (compared
24	with such costs if such drugs and biologicals were ac-
25	quired directly by such physicians).".



1	SEC. 304. DEMONSTRATION PROJECT FOR USE OF RECOV-
2	ERY AUDIT CONTRACTORS.
3	(a) In General.—The Secretary of Health and
4	Human Services shall conduct a demonstration project
5	under this section (in this section referred to as the
6	"project") to demonstrate the use of recovery audit contrac-
7	tors under the Medicare Integrity Program in identifying
8	underpayments and overpayments and recouping overpay-
9	ments under the medicare program for services for which
10	payment is made under part A or part B of title XVIII
11	of the Social Security Act. Under the project—
12	(1) payment may be made to such a contractor
13	on a contingent basis;
14	(2) a percentage of the amount recovered may be
15	retained by the Secretary and shall be available to the
16	program management account of the Centers for
17	Medicare & Medicaid Services; and
18	(3) the Secretary shall examine the efficacy of
19	such use with respect to duplicative payments, accu-
20	racy of coding, and other payment policies in which
21	inaccurate payments arise.
22	(b) Scope and Duration.—
23	(1) Scope.—The project shall cover at least 2
24	States that are among the States with—
25	(A) the highest per capita utilization rates
26	of medicare services, and



1	(B) at least 3 contractors.
2	(2) Duration.—The project shall last for not
3	longer than 3 years.
4	(c) Waiver.—The Secretary of Health and Human
5	Services shall waive such provisions of title XVIII of the
6	Social Security Act as may be necessary to provide for pay-
7	ment for services under the project in accordance with sub-
8	section (a).
9	(d) Qualifications of Contractors.—
10	(1) In general.—The Secretary shall enter into
11	a recovery audit contract under this section with an
12	entity only if the entity has staff that has the appro-
13	priate clinical knowledge of and experience with the
14	payment rules and regulations under the medicare
15	program or the entity has or will contract with an-
16	other entity that has such knowledgeable and experi-
17	enced staff.
18	(2) Ineligibility of certain contractors.—
19	The Secretary may not enter into a recovery audit
20	contract under this section with an entity to the ex-
21	tent that the entity is a fiscal intermediary under sec-
22	tion 1816 of the Social Security Act (42 U.S.C.
23	1395h), a carrier under section 1842 of such Act (42
24	U.S.C. 1395u), or a Medicare Administrative Con-
25	tractor under section 1874A of such Act.



1	(3) Preference for entities with dem-
2	ONSTRATED PROFICIENCY WITH PRIVATE INSURERS.—
3	In awarding contracts to recovery audit contractors
4	under this section, the Secretary shall give preference
5	to those risk entities that the Secretary determines
6	have demonstrated more than 3 years direct manage-
7	ment experience and a proficiency in recovery audits
8	with private insurers or under the medicaid program
9	under title XIX of such Act.
10	(e) Construction Relating to Conduct of Inves-
11	TIGATION OF FRAUD.—A recovery of an overpayment to a
12	provider by a recovery audit contractor shall not be con-
13	strued to prohibit the Secretary or the Attorney General
14	from investigating and prosecuting, if appropriate, allega-
15	tions of fraud or abuse arising from such overpayment.
16	(f) Report.—The Secretary of Health and Human
17	Services shall submit to Congress a report on the project
18	not later than 6 months after the date of its completion.
19	Such reports shall include information on the impact of the
20	project on savings to the medicare program and rec-
21	ommendations on the cost-effectiveness of extending or ex-
22	panding the project.



1	TITLE IV—RURAL HEALTH CARE
2	<b>IMPROVEMENTS</b>
3	SEC. 401. ENHANCED DISPROPORTIONATE SHARE HOS-
4	PITAL (DSH) TREATMENT FOR RURAL HOS-
5	PITALS AND URBAN HOSPITALS WITH FEWER
6	THAN 100 BEDS.
7	(a) Doubling the Cap.—
8	(1) In General.—Section $1886(d)(5)(F)$ (42)
9	$U.S.C.\ 1395ww(d)(5)(F))$ is amended by adding at
10	the end the following new clause:
11	"(xiv)(I) In the case of discharges in a fiscal year be-
12	ginning on or after October 1, 2003, subject to subclause
13	(II), there shall be substituted for the disproportionate share
14	adjustment percentage otherwise determined under clause
15	(iv) (other than subclause (I)) or under clause (viii), (x),
16	(xi), (xii), or (xiii), the disproportionate share adjustment
17	percentage determined under clause (vii) (relating to large,
18	urban hospitals).
19	"(II) Under subclause (I), the disproportionate share
20	adjustment percentage shall not exceed 10 percent for a hos-
21	pital that is not classified as a rural referral center under
22	subparagraph (C).".
23	(2) Conforming amendments.—Section
24	1886(d)(5)(F) (42 U.S.C. $1395ww(d)(5)(F)$ ) is



25

amended—

1	(A) in each of subclauses (II), (III), (IV),
2	(V), and (VI) of clause (iv), by inserting "subject
3	to clause (xiv) and" before "for discharges occur-
4	ring";
5	(B) in clause (viii), by striking "The for-
6	mula" and inserting "Subject to clause (xiv), the
7	formula"; and
8	(C) in each of clauses (x), (xi), (xii), and
9	(xiii), by striking "For purposes" and inserting
10	"Subject to clause (xiv), for purposes".
11	(b) Effective Date.—The amendments made by this
12	section shall apply with respect to discharges occurring on
13	or after October 1, 2003.
14	SEC. 402. IMMEDIATE ESTABLISHMENT OF UNIFORM
15	STANDARDIZED AMOUNT IN RURAL AND
16	SMALL URBAN AREAS.
17	(a) In General.—Section 1886(d)(3)(A) (42 U.S.C.
18	1395ww(d)(3)(A)) is amended—
19	(1) in clause (iv), by inserting "and ending on
20	or before September 30, 2003," after "October 1,
21	1995,"; and
22	(2) by redesignating clauses (v) and (vi) as
23	clauses (vii) and (viii), respectively, and inserting
24	after clause (iv) the following new clauses:



1	"(v) For discharges occurring in the fiscal year
2	beginning on October 1, 2003, the average standard-
3	ized amount for hospitals located in areas other than
4	a large urban area shall be equal to the average
5	standardized amount for hospitals located in a large
6	urban area.".
7	(b) Conforming Amendments.—
8	(1) Computing drg-specific rates.—Section
9	1886(d)(3)(D) (42 U.S.C. $1395ww(d)(3)(D)$ ) is
10	amended—
11	(A) in the heading, by striking "IN DIF-
12	FERENT AREAS";
13	(B) in the matter preceding clause (i), by
14	striking ", each of";
15	(C) in clause (i)—
16	(i) in the matter preceding subclause
17	(I), by inserting "for fiscal years before fis-
18	cal year 2004," before "for hospitals"; and
19	(ii) in subclause (II), by striking
20	"and" after the semicolon at the end;
21	(D) in clause (ii)—
22	(i) in the matter preceding subclause
23	(I), by inserting "for fiscal years before fis-
24	cal year 2004," before "for hospitals"; and



1	(ii) in subclause (II), by striking the
2	period at the end and inserting "; and";
3	and
4	(E) by adding at the end the following new
5	clause:
6	"(iii) for a fiscal year beginning after fiscal
7	year 2003, for hospitals located in all areas, to
8	the product of—
9	``(I) the applicable standardized
10	amount (computed under subparagraph
11	(A)), reduced under subparagraph (B), and
12	adjusted or reduced under subparagraph (C)
13	for the fiscal year; and
14	"(II) the weighting factor (determined
15	under paragraph (4)(B)) for that diagnosis-
16	related group.".
17	(2) Technical conforming sunset.—Section
18	1886(d)(3) (42 U.S.C. 1395ww(d)(3)) is amended—
19	(A) in the matter preceding subparagraph
20	(A), by inserting ", for fiscal years before fiscal
21	year 1997," before "a regional adjusted DRG
22	prospective payment rate"; and
23	(B) in subparagraph (D), in the matter
24	preceding clause (i), by inserting ", for fiscal



1	years before fiscal year 1997," before "a regional
2	DRG prospective payment rate for each region,".
3	SEC. 403. ESTABLISHMENT OF ESSENTIAL RURAL HOSPITAL
4	CLASSIFICATION.
5	(a) Classification.—Section 1861(mm) (42 U.S.C.
6	1395x(mm)) is amended—
7	(1) in the heading by adding "Essential
8	Rural Hospitals" at the end; and
9	(2) by adding at the end the following new para-
10	graphs:
11	"(4)(A) The term 'essential rural hospital' means a
12	subsection (d) hospital (as defined in section $1886(d)(1)(B)$ )
13	that is located in a rural area (as defined for purposes of
14	section 1886(d)), has more than 25 licensed acute care inpa-
15	tient beds, has applied to the Secretary for classification
16	as such a hospital, and with respect to which the Secretary
17	has determined that the closure of the hospital would sig-
18	nificantly diminish the ability of medicare beneficiaries to
19	obtain essential health care services.
20	"(B) The determination under subparagraph (A) shall
21	be based on the following criteria:
22	"(i) High proportion of medicare bene-
23	FICIARIES RECEIVING CARE FROM HOSPITAL.—(I) A
24	high percentage of such beneficiaries residing in the
25	area of the hospital who are hospitalized (during the



1	most recent year for which complete data are avail-
2	able) receive basic inpatient medical care at the hos-
3	pital.
4	"(II) For a hospital with more than 200 licensed
5	beds, a high percentage of such beneficiaries residing
6	in such area who are hospitalized (during such recent
7	year) receive specialized surgical inpatient care at the
8	hospital.
9	"(III) Almost all physicians described in section
10	1861(r)(1) in such area have privileges at the hospital
11	and provide their inpatient services primarily at the
12	hospital.
13	"(ii) Significant adverse impact in absence
14	OF HOSPITAL.—If the hospital were to close—
15	"(I) there would be a significant amount of
16	time needed for residents to reach emergency
17	treatment, resulting in a potential significant
18	harm to beneficiaries with critical illnesses or
19	injuries;
20	"(II) there would be an inability in the
21	community to stablize emergency cases for trans-
22	fers to another acute care setting, resulting in a
23	potential for significant harm to medicare bene-
24	ficiaries; and



1	"(III) any other nearby hospital lacks the
2	physical and clinical capacity to take over the
3	hospital's typical admissions.
4	"(C) In making such determination, the Secretary
5	may also consider the following:
6	"(i) Free-standing ambulatory surgery centers,
7	office-based oncology care, and imaging center services
8	are insufficient in the hospital's area to handle the
9	outpatient care of the hospital.
10	"(ii) Beneficiaries in nearby areas would be ad-
11	versely affected if the hospital were to close as the hos-
12	pital provides specialized knowledge and services to a
13	network of smaller hospitals and critical access hos-
14	pitals.
15	"(iii) Medicare beneficiaries would have dif-
16	ficulty in accessing care if the hospital were to close
17	as the hospital provides significant subsidies to sup-
18	port ambulatory care in local clinics, including men-
19	tal health clinics and to support post acute care.
20	"(iv) The hospital has a committment to provide
21	graduate medical education in a rural area.
22	"(C) Quality care.—The hospital inpatient
23	score for quality of care is not less than the median
24	hospital score for qualify of care for hospitals in the

State, as established under standards of the utiliza-



1	tion and quality control peer review organization
2	under part B of title XI or other quality standards
3	recognized by the Secretary.
4	A hospital classified as an essential rural hospital may not
5	change such classification and a hospital so classified shall
6	not be treated as a sole community hospital, medicare de-
7	pendent hospital, or rural referral center for purposes of
8	section 1886.".
9	(b) Payment Based on 102 Percent of Allowed
10	Costs.—
11	(1) Inpatient Hospital Services.—Section
12	1886(d) (42 U.S.C. 1395ww(d)) is amended by add-
13	ing at the end the following:
14	"(11) In the case of a hospital classified as an essential
15	rural hospital under section 1861(mm)(4) for a cost report-
16	ing period, the payment under this subsection for inpatient
17	hospital services for discharges occurring during the period
18	shall be based on 102 percent of the reasonable costs for such
19	services. Nothing in this paragraph shall be construed as
20	affecting the application or amount of deductibles or copay-
21	ments otherwise applicable to such services under part A
22	or as waiving any requirement for billing for such serv-



23 ices.".

1	(2) Hospital outpatient services.—Section
2	1833(t)(13) (42 U.S.C. $1395l(t)(13)$ ) is amended by
3	adding at the end the following new subparagraph:
4	"(B) Special rule for essential rural
5	HOSPITALS.—In the case of a hospital classified
6	as an essential rural hospital under section
7	1861(mm)(4) for a cost reporting period, the
8	payment under this subsection for covered OPD
9	services during the period shall be based on 102
10	percent of the reasonable costs for such services.
11	Nothing in this subparagraph shall be construed
12	as affecting the application or amount of
13	deductibles or copayments otherwise applicable
14	to such services under this part or as waiving
15	any requirement for billing for such services.".
16	(c) Effective Date.—The amendments made by this
17	section shall apply to cost reporting periods beginning on
18	or after October 1, 2004.
19	SEC. 404. MORE FREQUENT UPDATE IN WEIGHTS USED IN
20	HOSPITAL MARKET BASKET.
21	(a) More Frequent Updates in Weights.—After
22	revising the weights used in the hospital market basket
23	under section 1886(b)(3)(B)(iii) of the Social Security Act
24	$(42\ U.S.C.\ 1395ww(b)(3)(B)(iii))$ to reflect the most current
25	data available, the Secretary shall establish a frequency for



1	revising such weights, including the labor share, in such
2	market basket to reflect the most current data available
3	more frequently than once every 5 years.
4	(b) Report.—Not later than October 1, 2004, the Sec-
5	retary shall submit a report to Congress on the frequency
6	established under subsection (a), including an explanation
7	of the reasons for, and options considered, in determining
8	such frequency.
9	SEC. 405. IMPROVEMENTS TO CRITICAL ACCESS HOSPITAL
10	PROGRAM.
11	(a) Increase in Payment Amounts.—
12	(1) In General.—Sections 1814(l), 1834(g)(1),
13	and $1883(a)(3)$ (42 U.S.C. $1395f(l)$ ; $1395m(g)(1)$ ; 42
14	$U.S.C.\ 1395tt(a)(3))$ are each amended by inserting
15	"equal to 102 percent of" before "the reasonable
16	costs".
17	(2) Effective date.—The amendments made
18	by paragraph (1) shall apply to payments for services
19	furnished during cost reporting periods beginning on
20	or after October 1, 2003.
21	(b) Coverage of Costs for Certain Emergency
22	Room On-Call Providers.—
23	(1) In general.—Section 1834(g)(5) (42 U.S.C.
24	1395m(g)(5)) is amended—
25	(A) in the heading—



1	(i) by inserting "CERTAIN" before
2	"EMERGENCY"; and
3	(ii) by striking "PHYSICIANS" and in-
4	serting "PROVIDERS";
5	(B) by striking "emergency room physicians
6	who are on-call (as defined by the Secretary)"
7	and inserting "physicians, physician assistants,
8	nurse practitioners, and clinical nurse specialists
9	who are on-call (as defined by the Secretary) to
10	provide emergency services"; and
11	(C) by striking "physicians' services" and
12	inserting "services covered under this title".
13	(2) Effective date.—The amendment made by
14	paragraph (1) shall apply with respect to costs in-
15	curred for services provided on or after January 1,
16	2004.
17	(c) Modification of the Isolation Test for
18	Cost-Based CAH Ambulance Services.—
19	(1) In General.—Section 1834(l)(8) (42 U.S.C.
20	1395m(l)), as added by section $205(a)$ of BIPA (114
21	Stat. 2763A-482), is amended by adding at the end
22	the following: "The limitation described in the matter
23	following subparagraph (B) in the previous sentence
24	shall not apply if the ambulance services are fur-
25	nished by such a provider or supplier of ambulance



1	services who is a first responder to emergencies (as de-
2	termined by the Secretary).".
3	(2) Effective date.—The amendment made by
4	paragraph (1) shall apply to ambulances services fur-
5	nished on or after the first cost reporting period that
6	begins after the date of the enactment of this Act.
7	(d) Reinstatement of Periodic Interim Payment
8	(PIP).—
9	(1) In General.—Section 1815(e)(2) (42 U.S.C.
10	1395g(e)(2)) is amended—
11	(A) in the matter before subparagraph (A),
12	by inserting ", in the cases described in subpara-
13	graphs (A) through (D)" after "1986"; and
14	(B) by striking "and" at the end of sub-
15	paragraph (C);
16	(C) by adding "and" at the end of subpara-
17	graph (D); and
18	(D) by inserting after subparagraph (D) the
19	following new subparagraph:
20	$\lq\lq(E)$ inpatient critical access hospital services; $\lq\lq$ .
21	(2) Development of alternative methods
22	OF PERIODIC INTERIM PAYMENTS.—With respect to
23	periodic interim payments to critical access hospitals
24	for inpatient critical access hospital services under
25	section $1815(e)(2)(E)$ of the Social Security Act, as



1	added by paragraph (1), the Secretary shall develop
2	alternative methods for such payments that are based
3	on expenditures of the hospital.
4	(3) Reinstatement of PIP.—The amendments
5	made by paragraph (1) shall apply to payments
6	made on or after January 1, 2004.
7	(e) Condition for Application of Special Physi-
8	CIAN PAYMENT ADJUSTMENT.—
9	(1) In General.—Section 1834(g)(2) (42 U.S.C.
10	1395m(g)(2)) is amended by adding after and below
11	subparagraph (B) the following:
12	"The Secretary may not require, as a condition for
13	applying subparagraph (B) with respect to a critical
14	access hospital, that each physician providing profes-
15	sional services in the hospital must assign billing
16	rights with respect to such services, except that such
17	subparagraph shall not apply to those physicians who
18	have not assigned such billing rights.".
19	(2) Effective date.—The amendment made by
20	paragraph (1) shall be effective as if included in the
21	enactment of section 403(d) of the Medicare, Med-
22	icaid, and SCHIP Balanced Budget Refinement Act
23	of 1999 (113 Stat. 1501A–371).
24	(f) Flexibility in Bed Limitation for Hos-
25	PITALS.—Section 1820 (42 U.S.C. 1395i-4) is amended—



1	(1) in subsection $(c)(2)(B)(iii)$ , by inserting
2	"subject to paragraph (3)" after "(iii) provides";
3	(2) by adding at the end of subsection (c) the fol-
4	lowing new paragraph:
5	"(3) Increase in maximum number of beds
6	FOR HOSPITALS WITH STRONG SEASONAL CENSUS
7	FLUCTUATIONS.—
8	"(A) In general.—Subject to subpara-
9	graph (C), in the case of a hospital that dem-
10	onstrates that it meets the standards established
11	under subparagraph (B) and has not made the
12	election described in subsection $(f)(2)(A)$ , the bed
13	limitations otherwise applicable under para-
14	$graph\ (2)(B)(iii)\ and\ subsection\ (f)\ shall\ be\ in-$
15	creased by 5 beds.
16	"(B) Standards.—The Secretary shall
17	specify standards for determining whether a crit-
18	ical access hospital has sufficiently strong sea-
19	sonal variations in patient admissions to justify
20	the increase in bed limitation provided under
21	subparagraph (A)."; and
22	(3) in subsection (f)—
23	(A) by inserting "(1)" after "(f)"; and
24	(B) by adding at the end the following new
25	paragraph:



1	"(2)(A) A hospital may elect to treat the reference in
2	paragraph (1) to '15 beds' as a reference to '25 beds', but
3	only if no more than 10 beds in the hospital are at any
4	time used for non-acute care services. A hospital that makes
5	such an election is not eligible for the increase provided
6	under subsection $(c)(3)(A)$ .
7	"(B) The limitations in numbers of beds under the first
8	sentence of paragraph (1) are subject to adjustment under
9	subsection $(c)(3)$ .".
10	(4) Effective date.—The amendments made
11	by this subsection shall apply to designations made
12	before, on, or after January 1, 2004.
13	(g) Additional 5-Year Period of Funding for
14	Grant Program.—
15	(1) In General.—Section 1820(g) (42 U.S.C.
16	1395i-4(g)) is amended by adding at the end the fol-
17	lowing new paragraph:
18	"(4) Funding.—
19	"(A) In general.—Subject to subpara-
20	graph (B), payment for grants made under this
21	subsection during fiscal years 2004 through 2008
22	shall be made from the Federal Hospital Insur-
23	ance Trust Fund.
24	"(B) Annual aggregate limitation.—In
25	no case may the amount of payment provided for



1	under subparagraph (A) for a fiscal year exceed
2	\$25,000,000.".
3	(2) Conforming amendment.—Section 1820
4	(42 U.S.C. 1395i-4) is amended by striking sub-
5	section $(j)$ .
6	SEC. 406. REDISTRIBUTION OF UNUSED RESIDENT POSI-
7	TIONS.
8	(a) In General.—Section 1886(h)(4) (42 U.S.C.
9	1395ww(h)(4)) is amended—
10	(1) in subparagraph $(F)(i)$ , by inserting "subject
11	to subparagraph (I)," after "October 1, 1997,";
12	(2) in subparagraph $(H)(i)$ , by inserting "subject
13	to subparagraph (I)," after "subparagraphs (F) and
14	(G),"; and
15	(3) by adding at the end the following new sub-
16	paragraph:
17	"(I) Redistribution of unused resi-
18	DENT POSITIONS.—
19	"(i) Reduction in limit based on
20	UNUSED POSITIONS.—
21	"(I) In general.—If a hospital's
22	resident level (as defined in clause
23	(iii)(I)) is less than the otherwise ap-
24	plicable resident limit (as defined in
25	clause $(iii)(II)$ ) for each of the ref-



1	erence periods (as defined in subclause
2	(II)), effective for cost reporting peri-
3	ods beginning on or after January 1,
4	2004, the otherwise applicable resident
5	limit shall be reduced by 75 percent of
6	the difference between such limit and
7	the reference resident level specified in
8	subclause (III) (or subclause (IV) if
9	applicable).
10	"(II) Reference periods de-
11	FINED.—In this clause, the term 'ref-
12	erence periods' means, for a hospital,
13	the 3 most recent consecutive cost re-
14	porting periods of the hospital for
15	which cost reports have been settled (or,
16	if not, submitted) on or before Sep-
17	tember 30, 2002.
18	"(III) REFERENCE RESIDENT
19	LEVEL.—Subject to subclause (IV), the
20	reference resident level specified in this
21	subclause for a hospital is the highest
22	resident level for the hospital during
23	any of the reference periods.
24	"(IV) Adjustment process.—
25	Upon the timely request of a hospital,



1	the Secretary may adjust the reference
2	resident level for a hospital to be the
3	resident level for the hospital for the
4	cost reporting period that includes
5	July 1, 2003.
6	"(V) Affiliation.—With respect
7	to hospitals which are members of the
8	same affiliated group (as defined by
9	the Secretary under subparagraph
10	(H)(ii)), the provisions of this section
11	shall be applied with respect to such an
12	affiliated group by deeming the affili-
13	ated group to be a single hospital.
14	"(ii) Redistribution.—
15	"(I) In general.—The Secretary
16	is authorized to increase the otherwise
17	applicable resident limits for hospitals
18	by an aggregate number estimated by
19	the Secretary that does not exceed the
20	aggregate reduction in such limits at-
21	tributable to clause (i) (without taking
22	into account any adjustment under
23	subclause (IV) of such clause).
24	"(II) Effective date.—No in-

crease under subclause (I) shall be per-



1	mitted or taken into account for a hos-
2	pital for any portion of a cost report-
3	ing period that occurs before July 1,
4	2004, or before the date of the hos-
5	pital's application for an increase
6	under this clause. No such increase
7	shall be permitted for a hospital unless
8	the hospital has applied to the Sec-
9	retary for such increase by December
10	31, 2005.
11	"(III) Considerations in redis-
12	TRIBUTION.—In determining for which
13	hospitals the increase in the otherwise
14	applicable resident limit is provided
15	under subclause (I), the Secretary shall
16	take into account the need for such an
17	increase by specialty and location in-
18	volved, consistent with subclause (IV).
19	"(IV) Priority for rural and
20	SMALL URBAN AREAS.—In determining
21	for which hospitals and residency
22	training programs an increase in the
23	otherwise applicable resident limit is
24	provided under subclause (I), the Sec-

 $retary\ shall\ first\ distribute\ the\ increase$ 



1	to programs of hospitals located in
2	rural areas or in urban areas that are
3	not large urban areas (as defined for
4	purposes of subsection (d)) on a first-
5	come-first-served basis (as determined
6	by the Secretary) based on a dem-
7	onstration that the hospital will fill the
8	positions made available under this
9	clause and not to exceed an increase of
10	25 full-time equivalent positions with
11	respect to any hospital.
12	"(V) Application of locality
13	ADJUSTED NATIONAL AVERAGE PER
14	RESIDENT AMOUNT.—With respect to
15	additional residency positions in a
16	hospital attributable to the increase
17	provided under this clause, notwith-
18	standing any other provision of this
19	subsection, the approved FTE resident
20	amount is deemed to be equal to the lo-
21	cality adjusted national average per
22	resident amount computed under sub-
23	paragraph (E) for that hospital.
24	"(VI) Construction.—Nothing
25	in this clause shall be construed as per-



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1	mitting the redistribution of reductions
2	in residency positions attributable to
3	voluntary reduction programs under
4	paragraph (6) or as affecting the abil-
5	ity of a hospital to establish new med-
6	ical residency training programs
7	$under\ subparagraph\ (H).$
8	"(iii) Resident Level and limit de-
9	FINED.—In this subparagraph:
10	"(I) Resident Level.—The term
11	'resident level' means, with respect to a
12	hospital, the total number of full-time
13	equivalent residents, before the applica-
14	tion of weighting factors (as deter-
15	mined under this paragraph), in the
16	fields of allopathic and osteopathic
17	medicine for the hospital.
18	"(II) Otherwise applicable
19	RESIDENT LIMIT.—The term 'otherwise
20	applicable resident limit' means, with
21	respect to a hospital, the limit other-
22	wise applicable under subparagraphs
23	(F)(i) and $(H)$ on the resident level for
24	the hospital determined without regard
25	to this subparagraph.".



1	(b) Conforming Amendment to IME.—Section
2	1886(d)(5)(B)(v) (42 U.S.C. $1395ww(d)(5)(B)(v)$ ) is
3	amended by adding at the end the following: "The provi-
4	sions of subparagraph (I) of subsection (h)(4) shall apply
5	with respect to the first sentece of this clause in the same
6	manner as it applies with respect to subparagraph (F) of
7	such subsection.".
8	(c) Report on Extension of Applications Under
9	Redistribution Program.—Not later than July 1, 2005,
10	the Secretary shall submit to Congress a report containing
11	recommendations regarding whether to extend the deadline
12	for applications for an increase in resident limits under
13	section 1886(h)(4)(I)(ii)(II) of the Social Security Act (as
14	added by subsection (a)).
15	SEC. 407. TWO-YEAR EXTENSION OF HOLD HARMLESS PRO-
16	VISIONS FOR SMALL RURAL HOSPITALS AND
17	SOLE COMMUNITY HOSPITALS UNDER PRO-
18	SPECTIVE PAYMENT SYSTEM FOR HOSPITAL
19	OUTPATIENT DEPARTMENT SERVICES.
20	(a) Hold Harmless Provisions.—
21	(1) In General.—Section $1833(t)(7)(D)(i)$ (42)
22	$U.S.C.\ 1395l(t)(7)(D)(i))$ is amended—
23	(A) in the heading, by striking "SMALL"
24	and inserting "CERTAIN":



1	(B) by inserting "or a sole community hos-
2	pital (as defined in section 1886(d)(5)(D)(iii))
3	located in a rural area" after "100 beds"; and
4	(C) by striking "2004" and inserting
5	"2006".
6	(2) Effective date.—The amendment made by
7	subsection (a)(2) shall apply with respect to payment
8	for OPD services furnished on and after January 1,
9	2004.
10	(b) Study; Adjustment.—
11	(1) Study.—The Secretary shall conduct a study
12	to determine if, under the prospective payment system
13	for hospital outpatient department services under sec-
14	tion 1833(t) of the Social Security Act (42 U.S.C.
15	1395l(t)), costs incurred by rural providers of services
16	by ambulatory payment classification groups (APCs)
17	exceed those costs incurred by urban providers of serv-
18	ices.
19	(2) Adjustment.—Insofar as the Secretary de-
20	termines under paragraph (1) that costs incurred by
21	rural providers exceed those costs incurred by urban
22	providers of services, the Secretary shall provide for
23	an appropriate adjustment under such section $1833(t)$
24	to reflect those higher costs by January 1, 2005.



1	SEC. 408. EXCLUSION OF CERTAIN RURAL HEALTH CLINIC
2	AND FEDERALLY QUALIFIED HEALTH CENTER
3	SERVICES FROM THE PROSPECTIVE PAYMENT
4	SYSTEM FOR SKILLED NURSING FACILITIES.
5	(a) In General.—Section 1888(e)(2)(A) (42 U.S.C.
6	1395yy(e)(2)(A)) is amended—
7	(1) in clause (i)(II), by striking "clauses (ii) and
8	(iii)" and inserting "clauses (ii), (iii), and (iv)"; and
9	(2) by adding at the end the following new
10	clause:
11	"(iv) Exclusion of certain rural
12	HEALTH CLINIC AND FEDERALLY QUALIFIED
13	HEALTH CENTER SERVICES.—Services de-
14	scribed in this clause are—
15	"(I) rural health clinic services
16	(as defined in paragraph (1) of section
17	1861(aa)); and
18	"(II) Federally qualified health
19	center services (as defined in para-
20	graph (3) of such section);
21	that would be described in clause (ii) if such
22	services were not furnished by an individual
23	affiliated with a rural health clinic or a
24	Federally qualified health center.".



1	(b) Effective Date.—The amendments made by sub-
2	section (a) shall apply to services furnished on or after Jan-
3	uary 1, 2004.
4	SEC. 409. RECOGNITION OF ATTENDING NURSE PRACTI-
5	TIONERS AS ATTENDING PHYSICIANS TO
6	SERVE HOSPICE PATIENTS.
7	(a) In General.—Section 1861(dd)(3)(B) (42 U.S.C.
8	1395x(dd)(3)(B)) is amended by inserting "or nurse practi-
9	tioner (as defined in subsection (aa)(5))" after "the physi-
10	cian (as defined in subsection (r)(1))".
11	(b) Prohibition on Nurse Practitioner Certi-
12	Fying Need for Hospice.—Section $1814(a)(7)(A)(i)(I)$
13	(42 U.S.C. $1395f(a)(7)(A)(i)(I)$ ) is amended by inserting
14	"(which for purposes of this subparagraph does not include
15	a nurse practitioner)" after "attending physician (as de-
16	fined in section $1861(dd)(3)(B)$ )".
17	SEC. 410. IMPROVEMENT IN PAYMENTS TO RETAIN EMER-
18	GENCY CAPACITY FOR AMBULANCE SERVICES
19	IN RURAL AREAS.
20	Section 1834(l) (42 U.S.C. 1395m(l)) is amended—
21	(1) by redesignating paragraph (8), as added by
22	section 221(a) of BIPA (114 Stat. 2763A-486), as
23	paragraph (9); and
24	(2) by adding at the end the following new para-
25	graph:



1	"(10) Assistance for rural providers fur-
2	NISHING SERVICES IN LOW MEDICARE POPULATION
3	DENSITY AREAS.—
4	"(A) In general.—In the case of ground
5	ambulance services furnished on or after Janu-
6	ary 1, 2004, for which the transportation origi-
7	nates in a qualified rural area (as defined in
8	subparagraph (B)), the Secretary shall provide
9	for an increase in the base rate of the fee sched-
10	ule for mileage for a trip established under this
11	subsection. In establishing such increase, the Sec-
12	retary shall, based on the relationship of cost
13	and volume, estimate the average increase in cost
14	per trip for such services as compared with the
15	cost per trip for the average ambulance service.
16	"(B) Qualified rural area defined.—
17	For purposes of subparagraph (A), the term
18	'qualified rural area' is a rural area (as defined
19	in section $1886(d)(2)(D)$ ) with a population den-
20	sity of medicare beneficiaries residing in the
21	area that is in the lowest quartile of all rural
22	county populations.".



1	SEC. 411. TWO-YEAR INCREASE FOR HOME HEALTH SERV-
2	ICES FURNISHED IN A RURAL AREA.
3	(a) In General.—In the case of home health services
4	furnished in a rural area (as defined in section
5	1886(d)(2)(D) of the Social Security Act (42 U.S.C.
6	1395ww(d)(2)(D))) during 2004 and 2005, the Secretary
7	shall increase the payment amount otherwise made under
8	section 1895 of such Act (42 U.S.C. 1395fff) for such serv-
9	ices by 5 percent.
10	(b) Waiving Budget Neutrality.—The Secretary
11	shall not reduce the standard prospective payment amount
12	(or amounts) under section 1895 of the Social Security Act
13	(42 U.S.C. 1395fff) applicable to home health services fur-
14	nished during a period to offset the increase in payments
15	resulting from the application of subsection (a).
16	SEC. 412. PROVIDING SAFE HARBOR FOR CERTAIN COL-
17	LABORATIVE EFFORTS THAT BENEFIT MEDI-
18	CALLY UNDERSERVED POPULATIONS.
19	(a) In General.—Section 1128B(b)(3) (42 U.S.C.
20	1320a-7(b)(3)), as amended by section $101(b)(2)$ , is
21	amended—
22	(1) in subparagraph (F), by striking "and" after
23	the semicolon at the end;
24	(2) in subparagraph (G), by striking the period
25	at the end and inserting "; and"; and



1	(3) by adding at the end the following new sub-
2	paragraph:
3	"(H) any remuneration between a public or
4	nonprofit private health center entity described
5	under clause (i) or (ii) of section $1905(l)(2)(B)$
6	and any individual or entity providing goods,
7	items, services, donations or loans, or a combina-
8	tion thereof, to such health center entity pursu-
9	ant to a contract, lease, grant, loan, or other
10	agreement, if such agreement contributes to the
11	ability of the health center entity to maintain or
12	increase the availability, or enhance the quality,
13	of services provided to a medically underserved
14	population served by the health center entity.".
15	(b) Rulemaking for Exception for Health Cen-
16	TER ENTITY ARRANGEMENTS.—
17	(1) Establishment.—
18	(A) In general.—The Secretary of Health
19	and Human Services (in this subsection referred
20	to as the "Secretary") shall establish, on an ex-
21	pedited basis, standards relating to the exception
22	described in section $1128B(b)(3)(H)$ of the Social
23	Security Act, as added by subsection (a), for
24	health center entity arrangements to the
25	antikickback penalties.



1	(B) Factors to consider.—The Secretary
2	shall consider the following factors, among oth-
3	ers, in establishing standards relating to the ex-
4	ception for health center entity arrangements
5	under subparagraph (A):
6	(i) Whether the arrangement between
7	the health center entity and the other party
8	results in savings of Federal grant funds or
9	increased revenues to the health center enti-
10	ty.
11	(ii) Whether the arrangement between
12	the health center entity and the other party
13	restricts or limits a patient's freedom of
14	choice.
15	(iii) Whether the arrangement between
16	the health center entity and the other party
17	protects a health care professional's inde-
18	pendent medical judgment regarding medi-
19	cally appropriate treatment.
20	The Secretary may also include other standards
21	and criteria that are consistent with the intent
22	of Congress in enacting the exception established
23	under this section.
24	(2) Interim final effect.—No later than 180
25	days after the date of enactment of this Act, the Sec-



1	retary shall publish a rule in the Federal Register
2	consistent with the factors under paragraph $(1)(B)$ .
3	Such rule shall be effective and final immediately on
4	an interim basis, subject to such change and revision,
5	after public notice and opportunity (for a period of
6	not more than 60 days) for public comment, as is
7	consistent with this subsection.
8	SEC. 413. GAO STUDY OF GEOGRAPHIC DIFFERENCES IN
9	PAYMENTS FOR PHYSICIANS' SERVICES.
10	(a) Study.—The Comptroller General of the United
11	States shall conduct a study of differences in payment
12	amounts under the physician fee schedule under section
13	1848 of the Social Security Act (42 U.S.C. 1395w-4) for
14	physicians' services in different geographic areas. Such
15	study shall include—
16	(1) an assessment of the validity of the geo-
17	graphic adjustment factors used for each component of
18	the fee schedule;
19	(2) an evaluation of the measures used for such
20	adjustment, including the frequency of revisions; and
21	(3) an evaluation of the methods used to deter-
22	mine professional liability insurance costs used in
23	computing the malpractice component, including a
24	review of increases in professional liability insurance
25	premiums and variation in such increases by State



1 and physician specialty and methods used to update 2 the geographic cost of practice index and relative weights for the malpractice component. 3 (b) Report.—Not later than 1 year after the date of 4 the enactment of this Act, the Comptroller General shall 5 submit to Congress a report on the study conducted under subsection (a). The report shall include recommendations 8 regarding the use of more current data in computing geographic cost of practice indices as well as the use of data 10 directly representative of physicians' costs (rather than proxy measures of such costs). 11 SEC. 414. TREATMENT OF MISSING COST REPORTING PERI-12 13 ODS FOR SOLE COMMUNITY HOSPITALS. 14 (a) IN GENERAL.—Section 1886(b)(3)(I) (42 U.S.C. 15 1395ww(b)(3)(I)) is amended by adding at the end the following new clause: 16 17 "(iii) In no case shall a hospital be denied treatment as a sole community hospital or payment (on the basis of 18 19 a target rate as such as a hospital) because data are unavailable for any cost reporting period due to changes in 20 21 ownership, changes in fiscal intermediaries, or other extraordinary circumstances, so long as data for at least one

applicable base cost reporting period is available.".



1	(b) Effective Date.—The amendment made by sub-
2	section (a) shall apply to cost reporting periods beginning
3	on or after January 1, 2004.
4	SEC. 415. EXTENSION OF TELEMEDICINE DEMONSTRATION
5	PROJECT.
6	Section 4207 of Balanced Budget Act of 1997 (Public
7	Law 105–33) is amended—
8	(1) in subsection (a)(4), by striking "4-year"
9	and inserting "8-year"; and
10	(2) in subsection $(d)(3)$ , by striking
11	"\$30,000,000" and inserting "\$60,000,000".
12	SEC. 416. ADJUSTMENT TO THE MEDICARE INPATIENT HOS-
13	PITAL PPS WAGE INDEX TO REVISE THE
<ul><li>13</li><li>14</li></ul>	PITAL PPS WAGE INDEX TO REVISE THE
14	LABOR-RELATED SHARE OF SUCH INDEX.
14 15	LABOR-RELATED SHARE OF SUCH INDEX.  (a) IN GENERAL.—Section 1886(d)(3)(E) (42 U.S.C.
<ul><li>14</li><li>15</li><li>16</li></ul>	LABOR-RELATED SHARE OF SUCH INDEX.  (a) IN GENERAL.—Section $1886(d)(3)(E)$ (42 U.S.C. $1395ww(d)(3)(E)$ ) is amended—
14 15 16 17	LABOR-RELATED SHARE OF SUCH INDEX.  (a) IN GENERAL.—Section 1886(d)(3)(E) (42 U.S.C. 1395ww(d)(3)(E)) is amended—  (1) by striking "WAGE LEVELS.—The Secretary"
14 15 16 17 18	LABOR-RELATED SHARE OF SUCH INDEX.  (a) IN GENERAL.—Section 1886(d)(3)(E) (42 U.S.C. 1395ww(d)(3)(E)) is amended—  (1) by striking "WAGE LEVELS.—The Secretary" and inserting "WAGE LEVELS.—
14 15 16 17 18	LABOR-RELATED SHARE OF SUCH INDEX.  (a) IN GENERAL.—Section 1886(d)(3)(E) (42 U.S.C. 1395ww(d)(3)(E)) is amended—  (1) by striking "WAGE LEVELS.—The Secretary" and inserting "WAGE LEVELS.—  "(i) IN GENERAL.—Except as provided in
14 15 16 17 18 19 20	LABOR-RELATED SHARE OF SUCH INDEX.  (a) IN GENERAL.—Section 1886(d)(3)(E) (42 U.S.C. 1395ww(d)(3)(E)) is amended—  (1) by striking "WAGE LEVELS.—The Secretary" and inserting "WAGE LEVELS.—  "(i) IN GENERAL.—Except as provided in clause (ii), the Secretary"; and
14 15 16 17 18 19 20 21	LABOR-RELATED SHARE OF SUCH INDEX.  (a) IN GENERAL.—Section 1886(d)(3)(E) (42 U.S.C. 1395ww(d)(3)(E)) is amended—  (1) by striking "WAGE LEVELS.—The Secretary" and inserting "WAGE LEVELS.—  "(i) IN GENERAL.—Except as provided in clause (ii), the Secretary"; and  (2) by adding at the end the following new



1	"(I) In general.—Except as provided
2	in subclause (II), for discharges occurring
3	on or after October 1, 2003, the Secretary
4	shall substitute the '62 percent' for the pro-
5	portion described in the first sentence of
6	clause $(i)$ .
7	"(II) Hold harmless for certain
8	HOSPITALS.—If the application of subclause
9	(I) would result in lower payments to a hos-
10	pital than would otherwise be made, then
11	this subparagraph shall be applied as if this
12	clause had not been enacted.".
13	(b) Waiving Budget Neutrality.—Section
14	1886(d)(3)(E) (42 U.S.C. $1395ww(d)(3)(E)$ ), as amended
15	by subsection (a), is amended by adding at the end of clause
16	(i) the following new sentence: "The Secretary shall apply
17	the previous sentence for any period as if the amendments
18	made by section 402(a) of the Medicare Prescription Drug
19	and Modernization Act of 2003 had not been enacted.".
20	SEC. 417. MEDICARE INCENTIVE PAYMENT PROGRAM IM-
21	PROVEMENTS FOR PHYSICIAN SCARCITY.
22	(a) Additional Bonus Payment for Certain Phy-
23	SICIAN SCARCITY AREAS.—



1	(1) In General.—Section 1833 (42 U.S.C.
2	1395l) is amended by adding at the end the following
3	new subsection:
4	"(u) Incentive Payments for Physician Scarcity
5	Areas.—
6	"(1) In general.—In the case of physicians'
7	services furnished in a year—
8	"(A) by a primary care physician in a pri-
9	mary care scarcity county (identified under
10	paragraph (4)); or
11	"(B) by a physician who is not a primary
12	care physician in a specialist care scarcity coun-
13	ty (as so identified),
14	in addition to the amount of payment that would oth-
15	erwise be made for such services under this part, there
16	also shall be paid an amount equal to 5 percent of
17	the payment amount for the service under this part.
18	"(2) Determination of ratios of physicians
19	to medicare beneficiaries in area.—Based upon
20	available data, the Secretary shall periodically deter-
21	mine, for each county or equivalent area in the
22	United States, the following:
23	"(A) Number of physicians practicing
24	IN THE AREA.—The number of physicians who
25	furnish physicians' services in the active practice



1	of medicine or osteopathy in that county or area,
2	other than physicians whose practice is exclu-
3	sively for the Federal Government, physicians
4	who are retired, or physicians who only provide
5	administrative services. Of such number, the
6	number of such physicians who are—
7	"(i) primary care physicians; or
8	"(ii) physicians who are not primary
9	care physicians.
10	"(B) Number of medicare beneficiaries
11	RESIDING IN THE AREA.—The number of indi-
12	viduals who are residing in the county and are
13	entitled to benefits under part A or enrolled
14	under this part, or both.
15	"(C) Determination of ratios.—
16	"(i) Primary care ratio.—The ratio
17	(in this paragraph referred to as the 'pri-
18	mary care ratio') of the number of primary
19	care physicians (determined under subpara-
20	$graph \ (A)(i)), \ to \ number \ of \ medicare \ bene-$
21	ficiaries determined under subparagraph
22	(B).
23	"(ii) Specialist care ratio.—The
24	ratio (in this paragraph referred to as the
25	'specialist care ratio') of the number of



1	other physicians (determined under sub-
2	paragraph (A)(ii)), to number of medicare
3	beneficiaries determined under subpara-
4	graph(B).
5	"(3) Ranking of counties.—The Secretary
6	shall rank each such county or area based separately
7	on its primary care ratio and its specialist care ratio.
8	"(4) Identification of counties.—The Sec-
9	retary shall identify—
10	"(A) those counties and areas (in this para-
11	graph referred to as 'primary care scarcity coun-
12	ties') with the lowest primary care ratios that
13	represent, if each such county or area were
14	weighted by the number of medicare beneficiaries
15	determined under paragraph $(2)(B)$ , an aggre-
16	gate total of 20 percent of the total of the medi-
17	care beneficiaries determined under such para-
18	graph; and
19	"(B) those counties and areas (in this sub-
20	section referred to as 'specialist care scarcity
21	counties') with the lowest specialist care ratios
22	that represent, if each such county or area were
23	weighted by the number of medicare beneficiaries
24	determined under paragraph (2)(B), an aggre-

gate total of 20 percent of the total of the medi-



1	care beneficiaries determined under such para-
2	graph.
3	There is no administrative or judicial review respect-
4	ing the identification of a county or area or the as-
5	signment of a specialty of any physician under this
6	paragraph.
7	"(5) Rural census tracks.—To the extent fea-
8	sible, the Secretary shall treat a rural census tract of
9	a metropolitan statistical area (as determined under
10	the most recent modification of the Goldsmith Modi-
11	fication, originally published in the Federal Register
12	on February 27, 1992 (57 Fed. Reg. 6725) as an
13	equivalent area for purposes of qualifying as a pri-
14	mary care scarcity county or specialist care scarcity
15	county under this subsection.
16	"(6) Physician Defined.—For purposes of this
17	paragraph, the term 'physician' means a physician
18	described in section $1861(r)(1)$ and the term 'primary
19	care physician' means a physician who is identified
20	in the available data as a general practitioner, family
21	practice practitioner, general internist, or obstetrician
22	or gynecologist.
23	"(7) Publication of list of counties.—In
24	carrying out this subsection for a year, the Secretary

shall include, as part of the proposed and final rule



1	to implement the physician fee schedule under section
2	1848 for the year, a list of all areas which will qual-
3	ify as a primary care scarcity county or specialist
4	care scarcity county under this subsection for the year
5	involved.".
6	(2) Effective date.—The amendments made
7	by subsection (a) shall apply to physicians' services
8	furnished or after January 1, 2004.
9	(b) Improvement to Medicare Incentive Payment
10	Program.—
11	(1) In General.—Section 1833(m) (42 U.S.C.
12	1395l(m)) is amended—
13	(A) by inserting "(1)" after "(m)"; and
14	(B) by adding at the end the following new
15	paragraphs:
16	"(2) The Secretary shall establish procedures under
17	which the Secretary, and not the physician furnishing the
18	service, is responsible for determining when a payment is
19	required to be made under paragraph (1).
20	"(3) In carrying out paragraph (1) for a year, the Sec-
21	retary shall include, as part of the proposed and final rule
22	to implement the physician fee schedule under section 1848
23	for the year, a list of all areas which will qualify as a health
24	professional shortage area under paragraph (1) for the year
25	involved.".



1	(2) Effective date.—The amendments made
2	by paragraph (1) shall apply to physicians' services
3	furnished or after January 1, 2004.
4	TITLE V—PROVISIONS RELATING
5	TO PART A
6	$Subtitle\ A-Inpatient\ Hospital$
7	Services
8	SEC. 501. REVISION OF ACUTE CARE HOSPITAL PAYMENT
9	UPDATES.
10	Section $1886(b)(3)(B)(i)$ (42 U.S.C.
11	1395ww(b)(3)(B)(i)) is amended—
12	(1) by striking "and" at the end of subclause
13	(XVIII);
14	(2) by striking subclause (XIX); and
15	(3) by inserting after subclause (XVIII) the fol-
16	lowing new subclauses:
17	"(XIX) for each of fiscal years 2004 through
18	2006, the market basket percentage increase minus 0.4
19	percentage points for hospitals in all areas; and
20	"(XX) for fiscal year 2007 and each subsequent
21	fiscal year, the market basket percentage increase for
22.	hospitals in all areas "



1	SEC. 502. RECOGNITION OF NEW MEDICAL TECHNOLOGIES
2	UNDER INPATIENT HOSPITAL PPS.
3	(a) Improving Timeliness of Data Collection.—
4	Section $1886(d)(5)(K)$ (42 U.S.C. $1395ww(d)(5)(K)$ ) is
5	amended by adding at the end the following new clause:
6	"(vii) Under the mechanism under this subparagraph,
7	the Secretary shall provide for the addition of new diagnosis
8	and procedure codes in April 1 of each year, but the addi-
9	tion of such codes shall not require the Secretary to adjust
10	the payment (or diagnosis-related group classification)
11	under this subsection until the fiscal year that begins after
12	such date.".
13	(b) Eligibility Standard for Technology
14	Outliers.—
15	(1) Minimum period for recognition of new
16	TECHNOLOGIES.—Section $1886(d)(5)(K)(vi)$ (42)
17	U.S.C. $1395ww(d)(5)(K)(vi)$ is amended—
18	(A) by inserting "(I)" after "(vi)"; and
19	(B) by adding at the end the following new
20	subclause:
21	"(II) Under such criteria, a service or technology shall
22	not be denied treatment as a new service or technology on
23	the basis of the period of time in which the service or tech-
24	nology has been in use if such period ends before the end
25	of the 2-to-3-year period that begins on the effective date
26	of implementation of a code under ICD-9-CM (or a suc-



1	cessor coding methodology) that enables the identification
2	of specific discharges in which the service or technology has
3	been used.".
4	(2) Adjustment of threshold.—Section
5	1886(d)(5)(K)(ii)(I) (42 U.S.C.
6	1395ww(d)(5)(K)(ii)(I)) is amended by inserting
7	"(applying a threshold specified by the Secretary that
8	is 75 percent of one standard deviation for the diag-
9	nosis-related group involved)" after "is inadequate".
10	(3) Criterion for substantial improve-
11	MENT.—Section $1886(d)(5)(K)(vi)$ (42 U.S.C.
12	$1395ww(d)(5)(K)(vi)), \ as \ amended \ by \ paragraph \ (1),$
13	is further amended by adding at the end the following
14	subclause:
15	"(III) The Secretary shall by regulation provide for
16	further clarification of the criteria applied to determine
17	whether a new service or technology represents an advance
18	in medical technology that substantially improves the diag-
19	nosis or treatment of beneficiaries. Under such criteria, in
20	determining whether a new service or technology represents
21	an advance in medical technology that substantially im-
22	proves the diagnosis or treatment of beneficiaries, the Sec-
23	retary shall deem a service or technology as meeting such
24	requirement if the service or technology is a drug or biologi-
25	cal that is designated under section 506 of the Federal Food,



1	Drug, and Cosmetic Act, approved under section 314.510
2	or 601.41 of title 21, Code of Federal Regulations, or des-
3	ignated for priority review when the marketing application
4	for such drug or biological was filed or is a medical device
5	for which an exemption has been granted under section
6	520(m) of such Act, or for which priority review has been
7	provided under section 515(d)(5) of such Act. Nothing in
8	this subclause shall be construed as effecting the authority
9	of the Secretary to determine whether items and services
10	are medically necessary and appropriate under section
11	1862(a)(1).".
12	(4) Process for public input.—Section
13	1886(d)(5)(K) (42 U.S.C. $1395ww(d)(5)(K)$ ), as
14	amended by paragraph (1), is amended—
15	(A) in clause (i), by adding at the end the
16	following: "Such mechanism shall be modified to
17	meet the requirements of clause (viii)."; and
18	(B) by adding at the end the following new
19	clause:
20	"(viii) The mechanism established pursuant to clause
21	(i) shall be adjusted to provide, before publication of a pro-
22	posed rule, for public input regarding whether a new service
23	or technology not described in the second sentence of clause
24	(vi)(III) represents an advance in medical technology that



1	substantially improves the diagnosis or treatment of bene-
2	ficiaries as follows:
3	"(I) The Secretary shall make public and peri-
4	odically update a list of all the services and tech-
5	nologies for which an application for additional pay-
6	ment under this subparagraph is pending.
7	"(II) The Secretary shall accept comments, rec-
8	ommendations, and data from the public regarding
9	whether the service or technology represents a substan-
10	$tial\ improvement.$
11	"(III) The Secretary shall provide for a meeting
12	at which organizations representing hospitals, physi-
13	cians, medicare beneficiaries, manufacturers, and any
14	other interested party may present comments, rec-
15	ommendations, and data to the clinical staff of the
16	Centers for Medicare & Medicaid Services before pub-
17	lication of a notice of proposed rulemaking regarding
18	whether service or technology represents a substantial
19	improvement.".
20	(c) Preference for Use of DRG Adjustment.—
21	Section $1886(d)(5)(K)$ (42 U.S.C. $1395ww(d)(5)(K)$ ) is fur-
22	ther amended by adding at the end the following new clause:
23	"(ix) Before establishing any add-on payment under
24	this subparagraph with respect to a new technology, the

25 Secretary shall seek to identify one or more diagnosis-re-



- 1 lated groups associated with such technology, based on simi-
- 2 lar clinical or anatomical characteristics and the cost of
- 3 the technology. Within such groups the Secretary shall as-
- 4 sign an eligible new technology into a diagnosis-related
- 5 group where the average costs of care most closely approxi-
- 6 mate the costs of care of using the new technology. In such
- 7 case, the new technology would no longer meet the threshold
- 8 of exceeding 75 percent of the standard deviation for the
- 9 diagnosis-related group involved under clause (ii)(I). No
- 10 add-on payment under this subparagraph shall be made
- 11 with respect to such new technology and this clause shall
- 12 not affect the application of paragraph (4)(C)(iii).".
- 13 (d) Improvement in Payment for New Tech-
- 14 NOLOGY.—Section 1886(d)(5)(K)(ii)(III) (42 U.S.C.
- 15 1395ww(d)(5)(K)(ii)(III)) is amended by inserting after
- 16 "the estimated average cost of such service or technology"
- 17 the following: "(based on the marginal rate applied to costs
- 18 under subparagraph (A))".
- 19 (e) Establishment of New Funding for Hospital
- 20 Inpatient Technology.—Section 1886(d)(5)(K)(ii)(III)
- 21 (42 U.S.C. 1395ww(d)(5)(K)(ii)(III)) is amended by strik-
- 22 ing "subject to paragraph (4)(C)(iii),".
- 23 (f) Effective Date.—
- 24 (1) In General.—The Secretary shall imple-
- 25 ment the amendments made by this section so that



1	they apply to classification for fiscal years beginning
2	with fiscal year 2005.
3	(2) Reconsiderations of applications for
4	FISCAL YEAR 2003 THAT ARE DENIED.—In the case of
5	an application for a classification of a medical serv-
6	ice or technology as a new medical service or tech-
7	nology under section $1886(d)(5)(K)$ of the Social Se-
8	curity Act (42 U.S.C. $1395ww(d)(5)(K)$ ) that was
9	filed for fiscal year 2004 and that is denied—
10	(A) the Secretary shall automatically recon-
11	sider the application as an application for fiscal
12	year 2005 under the amendments made by this
13	section; and
14	(B) the maximum time period otherwise
15	permitted for such classification of the service or
16	technology shall be extended by 12 months.
17	SEC. 503. INCREASE IN FEDERAL RATE FOR HOSPITALS IN
18	PUERTO RICO.
19	Section $1886(d)(9)$ (42 U.S.C. $1395ww(d)(9)$ ) is
20	amended—
21	(1) in subparagraph (A)—
22	(A) in clause (i), by striking "for discharges
23	beginning on or after October 1, 1997, 50 percent
24	(and for discharges between October 1, 1987, and
25	September 30, 1997, 75 percent)" and inserting



1	"the applicable Puerto Rico percentage (specified
2	in subparagraph (E))"; and
3	(B) in clause (ii), by striking "for dis-
4	charges beginning in a fiscal year beginning on
5	or after October 1, 1997, 50 percent (and for dis-
6	charges between October 1, 1987, and September
7	30, 1997, 25 percent)" and inserting "the appli-
8	cable Federal percentage (specified in subpara-
9	$graph \ (E))$ "; and
10	(2) by adding at the end the following new sub-
11	paragraph:
12	"(E) For purposes of subparagraph (A), for discharges
13	occurring—
14	"(i) on or after October 1, 1987, and before Octo-
15	ber 1, 1997, the applicable Puerto Rico percentage is
16	75 percent and the applicable Federal percentage is
17	25 percent;
18	"(ii) on or after October 1, 1997, and before Oc-
19	tober 1, 2003, the applicable Puerto Rico percentage
20	is 50 percent and the applicable Federal percentage is
21	50 percent;
22	"(iii) during fiscal year 2004, the applicable
23	Puerto Rico percentage is 41 percent and the applica-
24	ble Federal percentage is 59 percent;



1	"(v) during fiscal year 2005, the applicable
2	Puerto Rico percentage is 33 percent and the applica-
3	ble Federal percentage is 67 percent; and
4	"(v) on or after October 1, 2005, the applicable
5	Puerto Rico percentage is 25 percent and the applica-
6	ble Federal percentage is 75 percent.".
7	SEC. 504. WAGE INDEX ADJUSTMENT RECLASSIFICATION
8	REFORM.
9	(a) In General.—Section 1886(d) (42 U.S.C.
10	1395ww(d)) is amended by adding at the end the following:
11	"(11)(A) In order to recognize commuting patterns
12	among Metropolitan Statistical Areas and between such
13	Areas and rural areas, the Secretary shall establish a proc-
14	ess, upon application of a subsection (d) hospital that estab-
15	lishes that it is a qualifying hospital described in subpara-
16	graph (B), for an increase of the wage index applied under
17	paragraph (3)(E) for the hospital in the amount computed
18	under subparagraph (D).
19	"(B) A qualifying hospital described in this subpara-
20	graph is a subsection (d) hospital—
21	"(i) the average wages of which exceed the aver-
22	age wages for the area in which the hospital is lo-
23	cated: and



1	"(ii) which has at least 10 percent of its employ-
2	ees who reside in one or more higher wage index
3	areas.
4	"(C) For purposes of this paragraph, the term higher
5	wage index area' means, with respect to a hospital, an area
6	with a wage index that exceeds that of the area in which
7	the hospital is located.
8	"(D) The increase in the wage index under subpara-
9	graph (A) for a hospital shall be equal to the percentage
10	of the employees of the hospital that resides in any higher
11	wage index area multiplied by the sum of the products, for
12	each higher wage index area of—
13	"(i) the difference between (I) the wage index for
14	such area, and (II) the wage index of the area in
15	which the hospital is located (before the application of
16	this paragraph); and
17	"(ii) the number of employees of the hospital that
18	reside in such higher wage index area divided by the
19	total number of such employees that reside in all high
20	wage index areas.
21	"(E) The process under this paragraph shall be based
22	upon the process used by the Medicare Geographic Classi-
23	fication Review Board under paragraph (10) with respect
24	to data submitted by hospitals to the Board on the location



- of residence of hospital employees and wages under the applicable schedule established for geographic reclassification. 3 "(F) A reclassification under this paragraph shall be effective for a period of 3 fiscal years, except that the Sec-5 retary shall establish procedures under which a subsection (d) hospital may elect to terminate such reclassification be-6 fore the end of such period. 8 "(G) A hospital that is reclassified under this paragraph for a period is not eligible for reclassification under 10 paragraphs (8) or (10) during that period. 11 "(H) Any increase in a wage index under this paragraph for a hospital shall not be taken into account for pur-13 poses of— 14 "(i) computing the wage index for the area in 15 which the hospital is located or any other area; or 16 "(ii) applying any budget neutrality adjustment 17 with respect to such index under paragraph (8)(D).". 18 (b) Effective Date.—The amendment made by sub-19 section (a) shall first apply to the wage index for cost re-20 porting period beginning on or after October 1, 2004.
- 21 SEC. 505. MEDPAC REPORT ON SPECIALTY HOSPITALS.
- 22 (a) MEDPAC STUDY.—The Medicare Payment Advi-
- 23 sory Commission shall conduct a study of specialty hos-
- 24 pitals compared with other similar general acute care hos-



1	pitals under the medicare program. Such study shall
2	examine—
3	(1) whether there are excessive self-referrals;
4	(2) quality of care furnished;
5	(3) the impact of specialty hospitals on such gen-
6	eral acute care hospitals; and
7	(4) differences in the scope of services, medicaid
8	utilization, and uncompensated care furnished.
9	(b) Report.—Not later than 1 year after the date of
10	the enactment of this Act, the Secretary shall submit to Con-
11	gress a report on the study conducted under subsection (a),
12	and shall include any recommendations for legislation or
13	administrative change as the Secretary determines
14	appropriate.
15	Subtitle B—Other Provisions
16	SEC. 511. PAYMENT FOR COVERED SKILLED NURSING FA-
17	CILITY SERVICES.
18	(a) Adjustment to RUGs for AIDS Residents.—
19	Paragraph (12) of section 1888(e) (42 U.S.C. 1395yy(e))
20	is amended to read as follows:
21	"(12) Adjustment for residents with
22	AIDS.—
23	"(A) In general.—Subject to subpara-
24	graph (B), in the case of a resident of a skilled
25	nursing facility who is afflicted with acquired



1	immune deficiency syndrome (AIDS), the per
2	diem amount of payment otherwise applicable
3	shall be increased by 128 percent to reflect in-
4	creased costs associated with such residents.
5	"(B) Sunset.—Subparagraph (A) shall not
6	apply on and after such date as the Secretary
7	certifies that there is an appropriate adjustment
8	in the case mix under paragraph $(4)(G)(i)$ to
9	compensate for the increased costs associated
10	with residents described in such subparagraph.".
11	(b) Effective Date.—The amendment made by
12	paragraph (1) shall apply to services furnished on or after
13	October 1, 2003.
14	SEC. 512. COVERAGE OF HOSPICE CONSULTATION SERV-
15	ICES.
16	(a) Coverage of Hospice Consultation Serv-
<ul><li>16</li><li>17</li></ul>	(a) Coverage of Hospice Consultation Serv- ices.—Section 1812(a) (42 U.S.C. 1395d(a)) is amended—
17	ICES.—Section 1812(a) (42 U.S.C. 1395d(a)) is amended—
17 18	ICES.—Section 1812(a) (42 U.S.C. 1395d(a)) is amended—  (1) by striking "and" at the end of paragraph
17 18 19	ICES.—Section 1812(a) (42 U.S.C. 1395d(a)) is amended—  (1) by striking "and" at the end of paragraph  (3);
17 18 19 20	ICES.—Section 1812(a) (42 U.S.C. 1395d(a)) is amended—  (1) by striking "and" at the end of paragraph  (3);  (2) by striking the period at the end of para-
17 18 19 20 21	ICES.—Section 1812(a) (42 U.S.C. 1395d(a)) is amended—  (1) by striking "and" at the end of paragraph  (3);  (2) by striking the period at the end of paragraph  graph (4) and inserting "; and"; and
17 18 19 20 21 22	ICES.—Section 1812(a) (42 U.S.C. 1395d(a)) is amended—  (1) by striking "and" at the end of paragraph  (3);  (2) by striking the period at the end of paragraph (4) and inserting "; and"; and  (3) by inserting after paragraph (4) the fol-



1	have not previously received services under this para-
2	graph, services that are furnished by a physician who
3	is either the medical director or an employee of a hos-
4	pice program and that consist of—
5	"(A) an evaluation of the individual's need
6	for pain and symptom management;
7	"(B) counseling the individual with respect
8	to end-of-life issues and care options; and
9	"(C) advising the individual regarding ad-
10	vanced care planning.".
11	(b) Payment.—Section 1814(i) (42 U.S.C. l395f(i)) is
12	amended by adding at the end the following new paragraph:
13	"(4) The amount paid to a hospice program with re-
14	spect to the services under section 1812(a)(5) for which pay-
15	ment may be made under this part shall be equal to an
16	amount equivalent to the amount established for an office
17	or other outpatient visit for evaluation and management
18	associated with presenting problems of moderate severity
19	under the fee schedule established under section 1848(b),
20	other than the portion of such amount attributable to the
21	practice expense component.".
22	(c) Conforming Amendment.—Section
23	1861(dd)(2)(A)(i) (42 U.S.C. $1395x(dd)(2)(A)(i)$ ) is
24	amended by inserting before the comma at the end the fol-
25	lowing: "and services described in section 1812(a)(5)".



1	(d) Effective Date.—The amendments made by this
2	section shall apply to services provided by a hospice pro-
3	gram on or after January 1, 2004.
4	TITLE VI—PROVISIONS
5	RELATING TO PART B
6	Subtitle A—Physicians' Services
7	SEC. 601. REVISION OF UPDATES FOR PHYSICIANS' SERV-
8	ICES.
9	(a) UPDATE FOR 2004 AND 2005.—
10	(1) In general.—Section 1848(d) (42 U.S.C.
11	1395w-4(d)) is amended by adding at the end the fol-
12	lowing new paragraph:
13	"(5) UPDATE FOR 2004 AND 2005.—The update to
14	the single conversion factor established in paragraph
15	(1)(C) for each of 2004 and 2005 shall be not less
16	than 1.5 percent.".
17	(2) Conforming amendment.—Paragraph
18	(4)(B) of such section is amended, in the matter be-
19	fore clause (i), by inserting "and paragraph (5)"
20	after "subparagraph $(D)$ ".
21	(3) Not treated as change in law and reg-
22	ULATION IN SUSTAINABLE GROWTH RATE DETERMINA-
23	TION.—The amendments made by this subsection shall
24	not be treated as a change in law for purposes of ap-



1	plying section $1848(f)(2)(D)$ of the Social Security
2	$Act\ (42\ U.S.C.\ 1395w-4(f)(2)(D)).$
3	(b) Use of 10-Year Rolling Average in Com-
4	Puting Gross Domestic Product.—
5	(1) In General.—Section $1848(f)(2)(C)$ (42)
6	$U.S.C.\ 1395w-4(f)(2)(C)) \ is \ amended$ —
7	(A) by striking "projected" and inserting
8	"annual average"; and
9	(B) by striking "from the previous applica-
10	ble period to the applicable period involved" and
11	inserting "during the 10-year period ending with
12	the applicable period involved".
13	(2) Effective date.—The amendment made by
14	paragraph (1) shall apply to computations of the sus-
15	tainable growth rate for years beginning with 2003.
16	SEC. 602. STUDIES ON ACCESS TO PHYSICIANS' SERVICES.
17	(a) GAO STUDY ON BENEFICIARY ACCESS TO PHYSI-
18	CIANS' SERVICES.—
19	(1) Study.—The Comptroller General of the
20	United States shall conduct a study on access of
21	medicare beneficiaries to physicians' services under
22	the medicare program. The study shall include—
23	(A) an assessment of the use by beneficiaries
24	of such services through an analysis of claims



1	submitted by physicians for such services under
2	part B of the medicare program;
3	(B) an examination of changes in the use
4	by beneficiaries of physicians' services over time;
5	(C) an examination of the extent to which
6	physicians are not accepting new medicare bene-
7	ficiaries as patients.
8	(2) Report.—Not later than 18 months after the
9	date of the enactment of this Act, the Comptroller
10	General shall submit to Congress a report on the
11	study conducted under paragraph (1). The report
12	shall include a determination whether—
13	(A) data from claims submitted by physi-
14	cians under part B of the medicare program in-
15	dicate potential access problems for medicare
16	beneficiaries in certain geographic areas; and
17	(B) access by medicare beneficiaries to phy-
18	sicians' services may have improved, remained
19	constant, or deteriorated over time.
20	(b) Study and Report on Supply of Physicians.—
21	(1) Study.—The Secretary shall request the In-
22	stitute of Medicine of the National Academy of
23	Sciences to conduct a study on the adequacy of the
24	supply of physicians (including specialists) in the
25	United States and the factors that affect such supply.



1	(2) Report to congress.—Not later than 2
2	years after the date of enactment of this section, the
3	Secretary shall submit to Congress a report on the re-
4	sults of the study described in paragraph (1), includ-
5	ing any recommendations for legislation.
6	(c) GAO Study of Medicare Payment for Inhala-
7	TION THERAPY.—
8	(1) Study.—The Comptroller General of the
9	United States shall conduct a study to examine the
10	adequacy of current reimbursements for inhalation
11	therapy under the medicare program.
12	(2) Report.—Not later than May 1, 2004, the
13	Comptroller General shall submit to Congress a report
14	on the study conducted under paragraph (1).
15	SEC. 603. MEDPAC REPORT ON PAYMENT FOR PHYSICIANS
16	SERVICES.
17	(a) Practice Expense Component.—Not later than
18	1 year after the date of the enactment of this Act, the Medi-
19	care Payment Advisory Commission shall submit to Con-
20	gress a report on the effect of refinements to the practice
21	expense component of payments for physicians' services,
22	after the transition to a full resource-based payment system
23	in 2002, under section 1848 of the Social Security Act (42
24	U.S.C. 1395w-4). Such report shall examine the following
25	matters by physician specialty:



1	(1) The effect of such refinements on payment for
2	physicians' services.
3	(2) The interaction of the practice expense com-
4	ponent with other components of and adjustments to
5	payment for physicians' services under such section.
6	(3) The appropriateness of the amount of com-
7	pensation by reason of such refinements.
8	(4) The effect of such refinements on access to
9	care by medicare beneficiaries to physicians' services.
10	(5) The effect of such refinements on physician
11	participation under the medicare program.
12	(b) Volume of Physician Services.—The Medicare
13	Payment Advisory Commission shall submit to Congress a
14	report on the extent to which increases in the volume of
15	physicians' services under part B of the medicare program
16	are a result of care that improves the health and well-being
17	of medicare beneficiaries. The study shall include the fol-
18	lowing:
19	(1) An analysis of recent and historic growth in
20	the components that the Secretary includes under the
21	sustainable growth rate (under section 1848(f) of the
22	Social Security Act).
23	(2) An examination of the relative growth of vol-
24	ume in physician services between medicare bene-
25	ficiaries and other populations.



1	(3) An analysis of the degree to which new tech-
2	nology, including coverage determinations of the Cen-
3	ters for Medicare & Medicaid Services, has affected
4	the volume of physicians' services.
5	(4) An examination of the impact on volume of
6	demographic changes.
7	(5) An examination of shifts in the site of service
8	of services that influence the number and intensity of
9	services furnished in physicians' offices and the extent
10	to which changes in reimbursement rates to other pro-
11	viders have affected these changes.
12	(6) An evaluation of the extent to which the Cen-
13	ters for Medicare & Medicaid Services takes into ac-
14	count the impact of law and regulations on the sus-
15	tainable growth rate.
16	Subtitle B—Preventive Services
17	SEC. 611. COVERAGE OF AN INITIAL PREVENTIVE PHYSICAL
18	EXAMINATION.
19	(a) Coverage.—Section 1861(s)(2) (42 U.S.C.
20	1395x(s)(2)) is amended—
21	(1) in subparagraph (U), by striking "and" at
22	$the \ end;$
23	(2) in subparagraph (V), by inserting "and" at
24	the end; and



1	(3) by adding at the end the following new sub-
2	paragraph:
3	"(W) an initial preventive physical examination
4	(as defined in subsection (ww));".
5	(b) Services Described.—Section 1861 (42 U.S.C.
6	1395x) is amended by adding at the end the following new
7	subsection:
8	"Initial Preventive Physical Examination
9	"(ww) The term 'initial preventive physical examina-
10	tion' means physicians' services consisting of a physical ex-
11	amination with the goal of health promotion and disease
12	detection and includes items and services (excluding clinical
13	laboratory tests), as determined by the Secretary, consistent
14	with the recommendations of the United States Preventive
15	Services Task Force.".
16	(c) Waiver of Deductible and Coinsurance.—
17	(1) Deductible.—The first sentence of section
18	1833(b) (42 U.S.C. 1395l(b)) is amended—
19	(A) by striking "and" before "(6)", and
20	(B) by inserting before the period at the end
21	the following: ", and (7) such deductible shall not
22	apply with respect to an initial preventive phys-
23	ical examination (as defined in section
24	1861(ww))".



1	(2) $COINSURANCE$ .— $Section 1833(a)(1) (42)$
2	$U.S.C.\ 1395l(a)(1))$ is amended—
3	(A) in clause (N), by inserting "(or 100
4	percent in the case of an initial preventive phys-
5	ical examination, as defined in section
6	1861(ww))" after "80 percent"; and
7	(B) in clause (O), by inserting "(or 100
8	percent in the case of an initial preventive phys-
9	ical examination, as defined in section
10	1861(ww))" after "80 percent".
11	(d) Payment as Physicians' Services.—Section
12	1848(j)(3) (42 U.S.C. 1395w-4(j)(3)) is amended by insert-
13	ing "(2)(W)," after "(2)(S),".
14	(e) Other Conforming Amendments.—Section
15	1862(a) (42 U.S.C. 1395y(a)) is amended—
16	(1) in paragraph (1)—
17	(A) by striking "and" at the end of sub-
18	paragraph (H);
19	(B) by striking the semicolon at the end of
20	subparagraph (I) and inserting ", and"; and
21	(C) by adding at the end the following new
22	subparagraph:
23	"( $J$ ) in the case of an initial preventive physical
24	eramination which is performed not later than 6



1	months after the date the individual's first coverage
2	period begins under part B;"; and
3	(2) in paragraph (7), by striking "or (H)" and
4	inserting " $(H)$ , or $(J)$ ".
5	(f) Effective Date.—The amendments made by this
6	section shall apply to services furnished on or after January
7	1, 2004, but only for individuals whose coverage period be-
8	gins on or after such date.
9	SEC. 612. COVERAGE OF CHOLESTEROL AND BLOOD LIPID
10	SCREENING.
11	(a) Coverage.—Section 1861(s)(2) (42 U.S.C.
12	1395x(s)(2)), as amended by section 611(a), is amended—
13	(1) in subparagraph (V), by striking "and" at
14	$the\ end;$
15	(2) in subparagraph (W), by inserting "and" at
16	the end; and
17	(3) by adding at the end the following new sub-
18	paragraph:
19	"(X) cholesterol and other blood lipid
20	screening tests (as defined in subsection (XX));".
21	(b) Services Described.—Section 1861 (42 U.S.C.
22	1395x), as amended by section 611(b), is amended by add-
23	ing at the end the following new subsection:



1	"Cholesterol and Other Blood Lipid Screening Test
2	"(xx)(1) The term 'cholesterol and other blood lipid
3	screening test' means diagnostic testing of cholesterol and
4	other lipid levels of the blood for the purpose of early detec-
5	tion of abnormal cholesterol and other lipid levels.
6	"(2) The Secretary shall establish standards, in con-
7	sultation with appropriate organizations, regarding the fre-
8	quency and type of cholesterol and other blood lipid screen-
9	ing tests, except that such frequency may not be more often
10	than once every 2 years.".
11	(c) Frequency.—Section 1862(a)(1) (42 U.S.C.
12	1395y(a)(1)), as amended by section 611(e), is amended—
13	(1) by striking "and" at the end of subpara-
14	graph(I);
15	(2) by striking the semicolon at the end of sub-
16	paragraph (I) and inserting "; and"; and
17	(3) by adding at the end the following new sub-
18	paragraph:
19	"(K) in the case of a cholesterol and other blood
20	lipid screening test (as defined in section
21	1861(xx)(1)), which is performed more frequently
22	than is covered under section $1861(xx)(2)$ .".
23	(d) Effective Date.—The amendments made by this
24	section shall apply to tests furnished on or after January
25	1, 2005.



1	SEC. 613. WAIVER OF DEDUCTIBLE FOR COLORECTAL CAN-
2	CER SCREENING TESTS.
3	(a) In General.—The first sentence of section 1833(b)
4	(42 U.S.C. 1395l(b)), as amended by section 611(c)(1), is
5	amended—
6	(1) by striking "and" before "(7)"; and
7	(2) by inserting before the period at the end the
8	following: ", and (8) such deductible shall not apply
9	with respect to colorectal cancer screening tests (as de-
10	scribed in section 1861(pp)(1))".
11	(b) Conforming Amendments.—Paragraphs
12	(2)(C)(ii) and $(3)(C)(ii)$ of section 1834(d) (42 U.S.C.
13	1395m(d)) are each amended—
14	(1) by striking "DEDUCTIBLE AND" in the head-
15	ing; and
16	(2) in subclause (I), by striking "deductible or"
17	each place it appears.
18	(c) Effective Date.—The amendment made by this
19	section shall apply to items and services furnished on or
20	after January 1, 2004.
21	SEC. 614. IMPROVED PAYMENT FOR CERTAIN MAMMOG-
22	RAPHY SERVICES.
23	(a) Exclusion from OPD Fee Schedule.—Section
24	$1833(t)(1)(B)(iv) \ (42\ U.S.C.\ 1395l(t)(1)(B)(iv)) \ is \ amend-$
25	ed by inserting before the period at the end the following:
26	"and does not include screening mammography (as defined



1	in section 1861(jj)) and unilateral and bilateral diagnostic
2	mammography".
3	(b) Adjustment to Technical Component.—For
4	diagnostic mammography performed on or after January
5	1, 2004, for which payment is made under the physician
6	fee schedule under section 1848 of the Social Security Act
7	(42 U.S.C. 1395w-4), the Secretary, based on the most re-
8	cent cost data available, shall provide for an appropriate
9	adjustment in the payment amount for the technical compo-
10	nent of the diagnostic mammography.
11	(c) Effective Date.—The amendment made by sub-
12	section (a) shall apply to mammography performed on or
13	after January 1, 2004.
14	Subtitle C—Other Services
15	SEC. 621. HOSPITAL OUTPATIENT DEPARTMENT (HOPD)
16	PAYMENT REFORM.
17	(a) Payment for Drugs.—
18	(1) Modification of ambulatory payment
19	CLASSIFICATION (APC) GROUPS.—Section 1833(t) (42
20	$U.S.C.\ 1395l(t))$ is amended—
21	(A) by redesignating paragraph (13) as
22	paragraph (14); and
23	(B) by inserting after paragraph (12) the
24	following new paragraph:
25	"(13) Drug apc payment rates.—



1	"(A) In general.—With respect to pay-
2	ment for covered OPD services that includes a
3	specified covered outpatient drug (defined in sub-
4	paragraph (B)), the amount provided for pay-
5	ment for such drug under the payment system
6	under this subsection for services furnished in—
7	"(i) 2004, 2005, or 2006, shall in no
8	case—
9	"(I) exceed 95 percent of the aver-
10	age wholesale price for the drug; or
11	"(II) be less than the transition
12	percentage (under subparagraph (C))
13	of the average wholesale price for the
14	drug; or
15	"(ii) a subsequent year, shall be equal
16	to the average price for the drug for that
17	area and year established under the com-
18	petitive acquisition program under section
19	1847A as calculated and applied by the Sec-
20	retary for purposes of this paragraph.
21	"(B) Specified covered outpatient
22	DRUG DEFINED.—
23	"(i) In general.—In this paragraph,
24	the term 'specified covered outpatient drug'
25	means, subject to clause (ii), a covered out-



1	patient drug (as defined in 1927(k)(2), that
2	is—
3	$``(I)\ a\ radiopharmaceutical;\ or$
4	"(II) a drug or biological for
5	which payment was made under para-
6	graph (6) (relating to pass-through
7	payments) on or before December 31,
8	2002.
9	"(ii) Exception.—Such term does not
10	include—
11	"(I) a drug for which payment is
12	first made on or after January 1,
13	2003, under paragraph (6); or
14	"(II) a drug for a which a tem-
15	porary HCPCS code has not been as-
16	signed.
17	"(C) Transition towards historical av-
18	ERAGE ACQUISITION COST.—The transition per-
19	centage under this subparagraph for drugs fur-
20	nished in a year is determined in accordance
21	with the following table:

## The transition percentage for—

For the year—	Single source drugs are—	Innovator mul- tiple source drugs are—	Generic drugs are—
2004	83%	81.5%	46%
2005	77%	75%	46%
2006	71%	68%	46%



1	"(D) Payment for New Drugs until
2	TEMPORARY HCPCS CODE ASSIGNED.—With re-
3	spect to payment for covered OPD services that
4	includes a covered outpatient drug (as defined in
5	1927(k)) for a which a temporary HCPCS code
6	has not been assigned, the amount provided for
7	payment for such drug under the payment sys-
8	tem under this subsection shall be equal to 95
9	percent of the average wholesale price for the
10	drug.
11	"(E) Classes of drugs.—For purposes of
12	this paragraph, each of the following shall be
13	treated as a separate class of drugs:
14	"(i) Sole source drugs.—A sole
15	source drug which for purposes of this para-
16	graph means a drug or biological that is
17	not a multiple source drug (as defined in
18	subclauses (I) and (II) of section
19	1927(k)(7)(A)(i)) and is not a drug ap-
20	proved under an abbreviated new drug ap-
21	plication under section 355(j) of the Federal
22	Food, Drug, and Cosmetic Act.
23	"(ii) Innovator multiple source
24	DRUGS.—Innovator multiple source drugs

(as defined in section 1927(k)(7)(A)(ii)).



25

1	"(iii) Noninnovator multiple
2	SOURCE DRUGS.—Noninnovator multiple
3	source drugs (as defined in section
4	1927(k)(7)(A)(iii)).
5	"(F) Inapplicability of expenditures
6	IN DETERMINING CONVERSION FACTORS.—Addi-
7	tional expenditures resulting from this para-
8	graph and paragraph (14)(C) in a year shall not
9	be taken into account in establishing the conver-
10	sion factor for that year.".
11	(2) Reduction in threshold for separate
12	APCS FOR DRUGS.—Section 1833(t)(14), as redesig-
13	nated by paragraph (1)(A), is amended by adding at
14	the end the following new subparagraph:
15	"(B) Threshold for establishment of
16	SEPARATE APCS FOR DRUGS.—The Secretary
17	shall reduce the threshold for the establishment of
18	separate ambulatory procedure classification
19	groups (APCs) with respect to drugs to \$50 per
20	administration.".
21	(3) Exclusion of separate drug apcs from
22	OUTLIER PAYMENTS.—Section 1833(t)(5) is amended
23	by adding at the end the following new subparagraph:
24	"(E) Exclusion of separate drug apcs
25	From outlier payments.—No additional pay-



1	ment shall be made under subparagraph (A) in
2	the case of ambulatory procedure codes estab-
3	lished separately for drugs.".
4	(4) Payment for pass through drugs.—
5	Clause (i) of section $1833(t)(6)(D)$ (42 U.S.C.
6	1395l(t)(6)(D)) is amended by inserting after "under
7	section 1842(o)" the following: "(or if the drug is cov-
8	ered under a competitive acquisition contract under
9	section 1847A for an area, an amount determined by
10	the Secretary equal to the average price for the drug
11	for that area and year established under such section
12	as calculated and applied by the Secretary for pur-
13	poses of this paragraph)".
14	(5) Effective date.—The amendments made
15	by this subsection shall apply to services furnished on
16	or after January 1, 2004.
17	(b) Special Payment for Brachytherapy.—
18	(1) In general.—Section 1833(t)(14), as so re-
19	designated and amended by subsection $(a)(2)$ , is
20	amended by adding at the end the following new sub-
21	paragraph:
22	"(C) PAYMENT FOR DEVICES OF
23	BRACHYTHERAPY AT CHARGES ADJUSTED TO
24	COST.—Notwithstanding the preceding provi-

sions of this subsection, for a device of



25

1	brachytherapy furnished on or after January 1,
2	2004, and before January 1, 2007, the payment
3	basis for the device under this subsection shall be
4	equal to the hospital's charges for each device
5	furnished, adjusted to cost.".
6	(2) Specification of groups for
7	BRACHYTHERAPY DEVICES.—Section $1833(t)(2)$ (42)
8	$U.S.C.\ 1395l(t)(2)$ is amended—
9	(A) in subparagraph (F), by striking "and"
10	at the end;
11	(B) in subparagraph (G), by striking the
12	period at the end and inserting "; and"; and
13	(C) by adding at the end the following new
14	subparagraph:
15	"(H) with respect to devices of
16	brachytherapy, the Secretary shall create addi-
17	tional groups of covered OPD services that clas-
18	sify such devices separately from the other serv-
19	ices (or group of services) paid for under this
20	subsection in a manner reflecting the number,
21	isotope, and radioactive intensity of such devices
22	furnished, including separate groups for palla-
23	dium-103 and iodine-125 devices.".
24	(3) GAO REPORT.—The Comptroller General of
25	the United States shall conduct a study to determine



1	appropriate payment amounts under section
2	1833(t)(13)(B) of the Social Security Act, as added
3	by paragraph (1), for devices of brachytherapy. Not
4	later than January 1, 2005, the Comptroller General
5	shall submit to Congress and the Secretary a report
6	on the study conducted under this paragraph, and
7	shall include specific recommendations for appro-
8	priate payments for such devices.
9	(c) Application of Functional Equivalence
10	Test.—
11	(1) In General.—Section 1833(t)(6) (42 U.S.C.
12	1395l(t)(6)) is amended by adding at the end the fol-
13	lowing new subparagraph:
14	"(F) Limitation on application of func-
15	TIONAL EQUIVALENCE STANDARD.—The Sec-
16	retary may not apply a 'functional equivalence'
17	payment standard (including such standard pro-
18	mulgated on November 1, 2002) or any other
19	similar standard in order to deem a particular
20	drug or biological to be identical to or similar to
21	another drug or biological with respect to its
22	mechanism of action or clinical effect to deny
23	pass-through status to new drugs or biologics or
24	to remove such status of an existing eligible drug
25	or biologic under this paragraph unless—



1	"(i) the Secretary develops by regula-
2	tion (after providing notice and a period
3	for public comment) criteria for the appli-
4	cation of such standard; and
5	"(ii) such criteria provide for coordi-
6	nation with the Federal Food and Drug Ad-
7	ministration and require scientific studies
8	that show the clinical relationship between
9	the drugs or biologicals treated as function-
10	ally equivalent.".
11	(2) Effective date.—The amendment made by
12	paragraph (1) shall apply to the application of a
13	functional equivalence standard to a drug or biologi-
14	cal on or after the date of the enactment of this Act,
15	unless such application was being made to such drug
16	or biological prior to June 13, 2003.
17	(d) Hospital Acquisition Cost Study.—
18	(1) In general.—The Secretary shall conduct a
19	study on the costs incurred by hospitals in acquiring
20	covered outpatient drugs for which payment is made
21	under section 1833(t) of the Social Security Act (42
22	$U.S.C. \ 1395l(t)).$
23	(2) Drugs covered.—The study in paragraph
24	(1) shall not include those drugs for which the acqui-
25	sition costs is less than \$50 per administration.



1	(3) Representative sample of hospitals.—
2	In conducting the study under paragraph (1), the
3	Secretary shall collect data from a statistically valid
4	sample of hospitals with an urban/rural stratifica-
5	tion.
6	(4) Report.—Not later than January 1, 2006,
7	the Secretary shall submit to Congress a report on the
8	study conducted under paragraph (1), and shall in-
9	clude recommendations with respect to the following:
10	(A) Whether the study should be repeated,
11	and if so, how frequently.
12	(B) Whether the study produced useful data
13	on hospital acquisition cost.
14	(C) Whether data produced in the study is
15	appropriate for use in making adjustments to
16	payments for drugs and biologicals under section
17	1847A of the Social Security Act.
18	(D) Whether separate estimates can made of
19	overhead costs, including handing and admin-
20	istering costs for drugs.
21	SEC. 622. PAYMENT FOR AMBULANCE SERVICES.
22	(a) Phase-In Providing Floor Using Blend of
23	FEE SCHEDULE AND REGIONAL FEE SCHEDULES.—Sec-
24	tion 1834(l) (42 U.S.C. 1395m(l)), as amended by section
25	410(a), is amended—



1	(1) in paragraph (2)( $E$ ), by inserting "consistent
2	with paragraph (11)" after "in an efficient and fair
3	manner"; and
4	(2) by adding at the end the following new para-
5	graph:
6	"(11) Phase-in providing floor using blend
7	OF FEE SCHEDULE AND REGIONAL FEE SCHED-
8	ULES.—In carrying out the phase-in under para-
9	$graph\ (2)(E)\ for\ each\ level\ of\ service\ furnished\ in\ a$
10	year, the portion of the payment amount that is based
11	on the fee schedule shall not be less than the following
12	blended rate of the fee schedule under paragraph (1)
13	and of a regional fee schedule for the region involved:
14	"(A) For 2004, the blended rate shall be
15	based 20 percent on the fee schedule under para-
16	graph (1) and 80 percent on the regional fee
17	schedule.
18	"(B) For 2005, the blended rate shall be
19	based 40 percent on the fee schedule under para-
20	graph (1) and 60 percent on the regional fee
21	schedule.
22	"(C) For 2006, the blended rate shall be
23	based 60 percent on the fee schedule under para-
24	graph (1) and 40 percent on the regional fee
25	schedule.



1	"(D) For 2007, 2008, and 2009, the blended
2	rate shall be based 80 percent on the fee schedule
3	under paragraph (1) and 20 percent on the re-
4	gional fee schedule.
5	"(E) For 2010 and each succeeding year,
6	the blended rate shall be based 100 percent on the
7	fee schedule under paragraph (1).
8	For purposes of this paragraph, the Secretary shall
9	establish a regional fee schedule for each of the 9 Cen-
10	sus divisions using the methodology (used in estab-
11	lishing the fee schedule under paragraph (1)) to cal-
12	culate a regional conversion factor and a regional
13	mileage payment rate and using the same payment
14	adjustments and the same relative value units as used
15	in the fee schedule under such paragraph.".
16	(b) Adjustment in Payment for Certain Long
17	TRIPS.—Section 1834(1), as amended by subsection (a), is
18	further amended by adding at the end the following new
19	paragraph:
20	"(12) Adjustment in payment for certain
21	LONG TRIPS.—In the case of ground ambulance serv-
22	ices furnished on or after January 1, 2004, and before
23	January 1, 2009, regardless of where the transpor-
24	tation originates, the fee schedule established under
25	this subsection shall provide that, with respect to the



1 payment rate for mileage for a trip above 50 miles 2 the per mile rate otherwise established shall be in-3 creased by 1/4 of the payment per mile otherwise ap-4 plicable to such miles.". 5 (c) GAO REPORT ON COSTS AND ACCESS.—Not later than December 31, 2005, the Comptroller General of the 6 United States shall submit to Congress an initial report 8 on how costs differ among the types of ambulance providers and on access, supply, and quality of ambulance services 10 in those regions and States that have a reduction in payment under the medicare ambulance fee schedule (under sec-12 tion 1834(l) of the Social Security Act, as amended by this section). Not later than December 31, 2007, the Comptroller General shall submit to Congress a final report on such ac-14 15 cess and supply. 16 (d) Effective Date.—The amendments made by this section shall apply to ambulance services furnished on or 18 after January 1, 2004. 19 SEC. 623. RENAL DIALYSIS SERVICES. 20 (a) Demonstration of Alternative Delivery 21 Models.— 22 (1) Use of advisory board.—In carrying out 23 the demonstration project relating to improving care 24 for people with end-stage renal disease through alter-

native delivery models (as published in the Federal



1	Register of June 4, 2003), the Secretary shall estab-
2	lish an advisory board comprised of representatives
3	described in paragraph (2) to provide advice and rec-
4	ommendations with respect to the establishment and
5	operation of such demonstration project.
6	(2) Representatives re-
7	ferred to in paragraph (1) include representatives of
8	the following:
9	(A) Patient organizations.
10	(B) Clinicians.
11	(C) The medicare payment advisory com-
12	mission, established under section 1805 of the
13	Social Security Act (42 U.S.C. 1395b-6).
14	(D) The National Kidney Foundation.
15	(E) The National Institute of Diabetes and
16	Digestive and Kidney Diseases of National Insti-
17	tutes of Health.
18	(F) End-stage renal disease networks.
19	(G) Medicare contractors to monitor quality
20	of care.
21	(I) providers of services and renal dialysis
22	facilities furnishing end-stage renal disease serv-
23	ices.
24	$(J)\ Economists.$
25	(K) Researchers.



1	(b) Restoring Composite Rate Exceptions for
2	PEDIATRIC FACILITIES.—
3	(1) In general.—Section 422(a)(2) of BIPA is
4	amended—
5	(A) in subparagraph (A), by striking "and
6	(C)" and inserting ", (C), and (D)";
7	(B) in subparagraph (B), by striking "In
8	the case" and inserting "Subject to subpara-
9	graph (D), in the case"; and
10	(C) by adding at the end the following new
11	subparagraph:
12	"(D) Inapplicability to pediatric fa-
13	CILITIES.—Subparagraphs (A) and (B) shall not
14	apply, as of October 1, 2002, to pediatric facili-
15	ties that do not have an exception rate described
16	in subparagraph (C) in effect on such date. For
17	purposes of this subparagraph, the term 'pedi-
18	atric facility' means a renal facility at least 50
19	percent of whose patients are individuals under
20	18 years of age.".
21	(2) Conforming amendment.—The fourth sen-
22	tence of section $1881(b)(7)$ (42 U.S.C. $1395rr(b)(7)$ ),
23	as amended by subsection (b), is further amended by
24	striking "Until" and inserting "Subject to section
25	422(a)(2) of the Medicare, Medicaid, and SCHIP



- 1 Benefits Improvement and Protection Act of 2000,
- 2 and until".
- 3 (c) Increase in Renal Dialysis Composite Rate
- 4 FOR SERVICES FURNISHED IN 2004.—Notwithstanding any
- 5 other provision of law, with respect to payment under part
- 6 B of title XVIII of the Social Security Act for renal dialysis
- 7 services furnished in 2004, the composite payment rate oth-
- 8 erwise established under section 1881(b)(7) of such Act (42
- 9 U.S.C. 1395rr(b)(7)) shall be increased by 1.6 percent.
- 10 SEC. 624. ONE-YEAR MORATORIUM ON THERAPY CAPS; PRO-
- 11 VISIONS RELATING TO REPORTS.
- 12 (a) 1-Year Moratorium on Therapy Caps.—Sec-
- 13  $tion \ 1833(g)(4) \ (42 \ U.S.C. \ 1395l(g)(4))$  is amended by
- 14 striking "and 2002" and inserting "2002, and 2004".
- 15 (b) Prompt Submission of Overdue Reports on
- 16 Payment and Utilization of Outpatient Therapy
- 17 Services.—Not later than December 31, 2003, the Sec-
- 18 retary shall submit to Congress the reports required under
- 19 section 4541(d)(2) of the Balanced Budget Act of 1997 (re-
- 20 lating to alternatives to a single annual dollar cap on out-
- 21 patient therapy) and under section 221(d) of the Medicare,
- 22 Medicaid, and SCHIP Balanced Budget Refinement Act of
- 23 1999 (relating to utilization patterns for outpatient ther-
- 24 *apy*).



1	(c) Identification of Conditions and Diseases
2	Justifying Waiver of Therapy Cap.—
3	(1) Study.—The Secretary shall request the In-
4	stitute of Medicine of the National Academy of
5	Sciences to identify conditions or diseases that should
6	justify conducting an assessment of the need to waive
7	the therapy caps under section $1833(g)(4)$ of the So-
8	cial Security Act (42 U.S.C. $1395l(g)(4)$ ).
9	(2) Reports to congress.—
10	(A) Preliminary report.—Not later than
11	July 1, 2004, the Secretary shall submit to Con-
12	gress a preliminary report on the conditions and
13	diseases identified under paragraph (1).
14	(B) Final report.—Not later than Sep-
15	tember 1, 2004, the Secretary shall submit to
16	Congress a final report on such conditions and
17	diseases.
18	(C) Recommendations.—Not later than
19	October 1, 2004, the Secretary shall submit to
20	Congress a recommendation of criteria, with re-
21	spect to such conditions and disease, under
22	which a waiver of the therapy caps would apply.
23	(d) GAO STUDY OF PATIENT ACCESS TO PHYSICAL
24	Therapist Services.—



1	(1) Study.—The Comptroller General of the
2	United States shall conduct a study on access to phys-
3	ical therapist services in States authorizing such serv-
4	ices without a physician referral and in States that
5	require such a physician referral. The study shall—
6	(A) examine the use of and referral patterns
7	for physical therapist services for patients age 50
8	and older in States that authorize such services
9	without a physician referral and in States that
10	require such a physician referral;
11	(B) examine the use of and referral patterns
12	for physical therapist services for patients who
13	are medicare beneficiaries;
14	(C) examine the potential effect of prohib-
15	iting a physician from referring patients to
16	physical therapy services owned by the physician
17	and provided in the physician's office;
18	(D) examine the delivery of physical thera-
19	pists' services within the facilities of Department
20	of Defense; and
21	(E) analyze the potential impact on medi-
22	care beneficiaries and on expenditures under the
23	medicare program of eliminating the need for a

physician referral and physician certification for



1	physical therapist services under the medicare
2	program.
3	(2) Report.—The Comptroller General shall
4	submit to Congress a report on the study conducted
5	under paragraph (1) by not later than 1 year after
6	the date of the enactment of this Act.
7	SEC. 625. ADJUSTMENT TO PAYMENTS FOR SERVICES FUR-
8	NISHED IN AMBULATORY SURGICAL CEN-
9	TERS.
10	Section $1833(i)(2)(C)$ (42 U.S.C. $1395l(i)(2)(C)$ ) is
11	amended in the last sentence by inserting "and each of fis-
12	cal years 2004 through 2008" after "In each of the fiscal
13	years 1998 through 2002".
14	SEC. 626. PAYMENT FOR CERTAIN SHOES AND INSERTS
15	UNDER THE FEE SCHEDULE FOR ORTHOTICS
16	AND PROSTHETICS.
17	(a) In General.—Section 1833(o) (42 U.S.C.
18	1395l(o)) is amended—
19	(1) in paragraph (1), by striking "no more than
20	the limits established under paragraph (2)" and in-
21	serting "no more than the amount of payment appli-
22	cable under paragraph (2)"; and
23	(2) in paragraph (2), to read as follows:
24	"(2)(A) Except as provided by the Secretary under
25	subparagraphs (B) and (C), the amount of payment under



- 1 this paragraph for custom molded shoes, extra depth shoes,
- 2 and inserts shall be the amount determined for such items
- 3 by the Secretary under section 1834(h).
- 4 "(B) The Secretary or a carrier may establish pay-
- 5 ment amounts for shoes and inserts that are lower than the
- 6 amount established under section 1834(h) if the Secretary
- 7 finds that shoes and inserts of an appropriate quality are
- 8 readily available at or below the amount established under
- 9 such section.
- 10 "(C) In accordance with procedures established by the
- 11 Secretary, an individual entitled to benefits with respect
- 12 to shoes described in section 1861(s)(12) may substitute
- 13 modification of such shoes instead of obtaining one (or
- 14 more, as specified by the Secretary) pair of inserts (other
- 15 than the original pair of inserts with respect to such shoes).
- 16 In such case, the Secretary shall substitute, for the payment
- 17 amount established under section 1834(h), a payment
- 18 amount that the Secretary estimates will assure that there
- 19 is no net increase in expenditures under this subsection as
- 20 a result of this subparagraph.".
- 21 (b) Conforming Amendments.—(1) Section
- 22 1834(h)(4)(C) (42 U.S.C. 1395m(h)(4)(C)) is amended by
- 23 inserting "(and includes shoes described in section
- 24 1861(s)(12))" after "in section 1861(s)(9)".



1	(2) Section $1842(s)(2)$ (42 U.S.C. $1395u(s)(2)$ ) is
2	amended by striking subparagraph (C).
3	(c) Effective Date.—The amendments made by this
4	section shall apply to items furnished on or after January
5	1, 2004.
6	SEC. 627. WAIVER OF PART B LATE ENROLLMENT PENALTY
7	FOR CERTAIN MILITARY RETIREES; SPECIAL
8	ENROLLMENT PERIOD.
9	(a) Waiver of Penalty.—
10	(1) In General.—Section 1839(b) (42 U.S.C.
11	1395r(b)) is amended by adding at the end the fol-
12	lowing new sentence: "No increase in the premium
13	shall be effected for a month in the case of an indi-
14	vidual who is 65 years of age or older, who enrolls
15	under this part during 2001, 2002, 2003, or 2004 and
16	who demonstrates to the Secretary before December
17	31, 2004, that the individual is a covered beneficiary
18	(as defined in section 1072(5) of title 10, United
19	States Code). The Secretary of Health and Human
20	Services shall consult with the Secretary of Defense in
21	identifying individuals described in the previous sen-
22	tence.".
23	(2) Effective date.—The amendment made by
24	paragraph (1) shall apply to premiums for months

beginning with January 2004. The Secretary of



1	Health and Human Services shall establish a method
2	for providing rebates of premium penalties paid for
3	months on or after January 2004 for which a penalty
4	does not apply under such amendment but for which
5	a penalty was previously collected.
6	(b) Medicare Part B Special Enrollment Pe-
7	RIOD.—
8	(1) In general.—In the case of any individual
9	who, as of the date of the enactment of this Act, is
10	65 years of age or older, is eligible to enroll but is not
11	enrolled under part B of title XVIII of the Social Se-
12	curity Act, and is a covered beneficiary (as defined
13	in section 1072(5) of title 10, United States Code), the
14	Secretary of Health and Human Services shall pro-
15	vide for a special enrollment period during which the
16	individual may enroll under such part. Such period
17	shall begin as soon as possible after the date of the en-
18	actment of this Act and shall end on December 31,
19	2004.
20	(2) Coverage period.—In the case of an indi-
21	vidual who enrolls during the special enrollment pe-
22	riod provided under paragraph (1), the coverage pe-
23	riod under part B of title XVIII of the Social Secu-
24	rity Act shall begin on the first day of the month fol-

lowing the month in which the individual enrolls.



1	SEC. 628. PART B DEDUCTIBLE.
2	Section 1833(b) (42 U.S.C. 1395l(b)) is amended—
3	(1) by striking "1991 and" and inserting
4	"1991,"; and
5	(2) by striking "and subsequent years" and in-
6	serting "and each subsequent year through 2003, and
7	for a subsequent year after 2003 the amount of such
8	deductible for the previous year increased by the an-
9	nual percentage increase in the monthly actuarial
10	rate under section 1839(a)(1) ending with such subse-
11	quent year (rounded to the nearest \$1)".
12	SEC. 629. EXTENSION OF COVERAGE OF INTRAVENOUS IM-
13	MUNE GLOBULIN (IVIG) FOR THE TREATMENT
14	OF PRIMARY IMMUNE DEFICIENCY DISEASES
15	IN THE HOME.
16	(a) In General.—Section 1861 (42 U.S.C. 1395x), as
17	amended by sections 611(a) and 612(a) is amended—
18	(1) in subsection $(s)(2)$ —
19	(A) by striking "and" at the end of sub-
20	paragraph (W);
21	(B) by adding "and" at the end of subpara-
22	graph(X); and
23	(C) by adding at the end the following new
24	subparagraph:
25	"(Y) intravenous immune globulin for the

treatment of primary immune deficiency diseases



1	in the home (as defined in subsection (yy));";
2	and
3	(2) by adding at the end the following new sub-
4	section:
5	"Intravenous Immune Globulin
6	"(yy) The term 'intravenous immune globulin' means
7	an approved pooled plasma derivative for the treatment in
8	the patient's home of a patient with a diagnosed primary
9	immune deficiency disease, but not including items or serv-
10	ices related to the administration of the derivative, if a phy-
11	sician determines administration of the derivative in the
12	patient's home is medically appropriate.".
13	(b) Payment as a Drug or Biological.—Section
14	1833(a)(1)(S) (42 U.S.C. 1395l(a)(1)(S)) is amended by in-
15	serting "(including intravenous immune globulin (as de-
16	fined in section 1861(yy)))" after "with respect to drugs
17	and biologicals".
18	(c) Effective Date.—The amendments made by this
19	section shall apply to items furnished administered on or
20	after January 1, 2004.
21	TITLE VII—PROVISIONS
22	RELATING TO PARTS A AND B
23	Subtitle A—Home Health Services
24	SEC. 701. UPDATE IN HOME HEALTH SERVICES.
25	(a) Change to Calender Year Update.—



1	(1) In General.—Section 1895(b) (42 U.S.C.
2	1395fff(b)(3)) is amended—
3	(A) in paragraph $(3)(B)(i)$ —
4	(i) by striking "each fiscal year (begin-
5	ning with fiscal year 2002)" and inserting
6	"fiscal year 2002 and for fiscal year 2003
7	and for each subsequent year (beginning
8	with 2004)"; and
9	(ii) by inserting "or year" after "the
10	fiscal year";
11	(B) in paragraph $(3)(B)(ii)(II)$ , by striking
12	"any subsequent fiscal year" and inserting
13	"2004 and any subsequent year";
14	(C) in paragraph $(3)(B)(iii)$ , by inserting
15	"or year" after "fiscal year" each place it ap-
16	pears;
17	(D) in paragraph $(3)(B)(iv)$ —
18	(i) by inserting "or year" after "fiscal
19	year" each place it appears; and
20	(ii) by inserting "or years" after "fis-
21	cal years"; and
22	(E) in paragraph (5), by inserting "or
23	year" after "fiscal year".
24	(2) Transition rule.—The standard prospec-
25	tive payment amount (or amounts) under section



1	1895(b)(3) of the Social Security Act for the calendar
2	quarter beginning on October 1, 2003, shall be such
3	amount (or amounts) for the previous calendar quar-
4	ter.
5	(b) Changes in Updates for 2004, 2005, and
6	2006.—Section $1895(b)(3)(B)(ii)$ (42 U.S.C.
7	1395fff(b)(3)(B)(ii)), as amended by subsection $(a)(1)(B)$ ,
8	is amended—
9	(1) by striking "or" at the end of subclause (I);
10	(2) by redesignating subclause (II) as subclause
11	(III);
12	(3) in subclause (III), as so redesignated, by
13	striking "2004" and inserting "2007"; and
14	(4) by inserting after subclause (I) the following
15	new subclause:
16	"(II) each of 2004, 2005, and
17	2006 the home health market basket
18	percentage increase minus 0.4 percent-
19	age points; or".
20	SEC. 702. ESTABLISHMENT OF REDUCED COPAYMENT FOR A
21	HOME HEALTH SERVICE EPISODE OF CARE
22	FOR CERTAIN BENEFICIARIES.
23	(a) Part A.—



1	(1) In General.—Section 1813(a) (42 U.S.C.
2	1395e(a)) is amended by adding at the end the fol-
3	lowing new paragraph:
4	"(5)(A)(i) Subject to clause (ii), the amount payable
5	for home health services furnished to the individual under
6	this title for each episode of care beginning in a year (begin-
7	ning with 2004) shall be reduced by a copayment equal to
8	the copayment amount specified in subparagraph (B)(ii)
9	for such year.
10	"(ii) The copayment under clause (i) shall not apply—
11	"(I) in the case of an individual who has been
12	determined to be entitled to medical assistance under
13	section $1902(a)(10)(A)$ or $1902(a)(10)(C)$ or to be a
14	qualified medicare beneficiary (as defined in section
15	1905(p)(1)), a specified low-income medicare bene-
16	ficiary described in section $1902(a)(10)(E)(iii)$ , or a
17	qualifying individual described in section
18	$1902(a)(10)(E)(iv)(I); \ and$
19	"(II) in the case of an episode of care which con-
20	sists of 4 or fewer visits.
21	" $(B)(i)$ The Secretary shall estimate, before the begin-
22	ning of each year (beginning with 2004), the national aver-
23	age payment under this title per episode for home health
24	services projected for the year involved.



1	"(ii) For each year the copayment amount under this
2	clause is equal to 1.5 percent of the national average pay-
3	ment estimated for the year involved under clause (i). Any
4	amount determined under the preceding sentence which is
5	not a multiple of \$5 shall be rounded to the nearest multiple
6	of $$5$ .
7	"(iii) There shall be no administrative or judicial re-
8	view under section 1869, 1878, or otherwise of the esti-
9	mation of average payment under clause (i).".
10	(2) Timely implementation.—Unless the Sec-
11	retary of Health and Human Services otherwise pro-
12	vides on a timely basis, the copayment amount speci-
13	fied under section $1813(a)(5)(B)(ii)$ of the Social Se-
14	curity Act (as added by paragraph (1)) for 2004 shall
15	be deemed to be \$40.
16	(b) Conforming Provisions.—
17	(1) Section $1833(a)(2)(A)$ (42 U.S.C.
18	1395l(a)(2)(A)) is amended by inserting 'less the co-
19	payment amount applicable under section
20	1813(a)(5)" after "1895".
21	(2) Section $1866(a)(2)(A)(i)$ (42 U.S.C.
22	1395cc(a)(2)(A)(i)) is amended—
23	(A) by striking "or coinsurance" and in-
24	serting ", coinsurance, or copayment"; and



1	(B) by striking "or $(a)(4)$ " and inserting
2	" $(a)(4)$ , or $(a)(5)$ ".
3	SEC. 703. MEDPAC STUDY ON MEDICARE MARGINS OF HOME
4	HEALTH AGENCIES.
5	(a) Study.—The Medicare Payment Advisory Com-
6	mission shall conduct a study of payment margins of home
7	health agencies under the home health prospective payment
8	system under section 1895 of the Social Security Act (42
9	U.S.C. 1395fff). Such study shall examine whether system-
10	atic differences in payment margins are related to dif-
11	ferences in case mix (as measured by home health resource
12	groups (HHRGs)) among such agencies. The study shall use
13	the partial or full-year cost reports filed by home health
14	agencies.
15	(b) Report.—Not later than 2 years after the date
16	of the enactment of this Act, the Commission shall submit
17	to Congress a report on the study under subsection (a).
18	Subtitle B—Direct Graduate
19	Medical Education
20	SEC. 711. EXTENSION OF UPDATE LIMITATION ON HIGH
21	COST PROGRAMS.
22	Section $1886(h)(2)(D)(iv)$ (42 U.S.C.
23	1395ww(h)(2)(D)(iv)) is amended—
24	(1) in subclause (I)—



1	(A) by inserting "AND 2004 THROUGH 2013"
2	after "AND 2002"; and
3	(B) by inserting "or during the period be-
4	ginning with fiscal year 2004 and ending with
5	fiscal year 2013" after "during fiscal year 2001
6	or fiscal year 2002"; and
7	(2) in subclause (II)—
8	(A) by striking "fiscal year 2004, or fiscal
9	year 2005," and
10	(B) by striking "For a" and inserting "For
11	the".
12	Subtitle C—Chronic Care
13	Improvement
14	SEC. 721. VOLUNTARY CHRONIC CARE IMPROVEMENT
15	UNDER TRADITIONAL FEE-FOR-SERVICE.
16	Title XVIII, as amended by section 105(a), is amended
17	by inserting after section 1807 the following new section:
18	"CHRONIC CARE IMPROVEMENT
19	"Sec. 1808. (a) In General.—
20	"(1) In general.—The Secretary shall establish
21	a process for providing chronic care improvement
22	programs in each CCIA region for medicare bene-
23	ficiaries who are not enrolled under part C or E and
24	who have certain chronic conditions, such as conges-
	tone in the configuration of t
25	tive heart failure, diabetes, chronic obstructive pul-



1	identified by the Secretary as appropriate for chronic
2	care improvement. Such a process shall begin to be
3	implemented no later than 1 year after the date of the
4	enactment of this section.
5	"(2) Terminology.—For purposes of this sec-
6	tion:
7	"(A) CCIA REGION.—The term 'CCIA re-
8	gion' means a chronic care improvement admin-
9	istrative region delineated under subsection
10	(b)(2).
11	"(B) Chronic care improvement pro-
12	GRAM.—The terms 'chronic care improvement
13	program' and 'program' means such a program
14	provided by a contractor under this section.
15	"(C) Contractor.—The term contractor
16	means an entity with a contract to provide a
17	chronic care improvement program in a CCIA
18	region under this section.
19	"(D) Individual plan.—The term 'indi-
20	vidual plan' means a chronic care improvement
21	plan established under subsection $(c)(5)$ for an
22	individual.
23	"(3) Construction.—Nothing in this section
24	shall be construed as expanding the amount, dura-
25	tion, or scope of benefits under this title.



1	"(b) Competitive Bidding Process.—
2	"(1) In general.—Under this section the Sec-
3	retary shall award contracts to qualified entities for
4	chronic care improvement programs for each CCIA
5	region under this section through a competitive bid-
6	ding process.
7	"(2) Process.—Under such process—
8	"(A) the Secretary shall delineate the
9	United States into multiple chronic care im-
10	provement administrative regions; and
11	"(B) the Secretary shall select at least 2
12	winning bidders in each CCIA region on the
13	basis of the ability of each bidder to carry out
14	a chronic care improvement program in accord-
15	ance with this section, in order to achieve im-
16	proved health and financial outcomes.
17	"(3) Eligible contractor.—A contractor may
18	be a disease improvement organization, health in-
19	surer, provider organization, a group of physicians,
20	or any other legal entity that the Secretary deter-
21	mines appropriate.
22	"(c) Chronic Care Improvement Programs.—
23	"(1) In general.—Each contract under this
24	section shall provide for the operation of a chronic



care improvement program by a contractor in a
CCIA region consistent with this subsection.
"(2) Identification of prospective program
PARTICIPANTS.—Each contractor shall have a method
for identifying medicare beneficiaries in the region to
whom it will offer services under its program. The
contractor shall identify such beneficiaries through
claims or other data and other means permitted con-
sistent with applicable disclosure provisions.
"(3) Initial contact by secretary.—The Sec-
retary shall communicate with each beneficiary iden-
tified under paragraph (2) as a prospective partici-
pant in one or more programs concerning participa-
tion in a program. Such communication may be
made by the Secretary (or on behalf of the Secretary)
and shall include information on the following:
"(A) A description of the advantages to the
beneficiary in participating in a program.
"(B) Notification that the contractor offer-
ing a program may contact the beneficiary di-
rectly concerning such participation.
"(C) Notification that participation in a
program is voluntary.
"(D) A description of the method for the

beneficiary to select the single program in which



1	the beneficiary wishes to participate and for de-
2	clining to participate and a method for obtain-
3	ing additional information concerning such par-
4	ticipation.
5	"(4) Participation.—A medicare beneficiary
6	may participate in only one program under this sec-
7	tion and may terminate participation at any time in
8	a manner specified by the Secretary.
9	"(5) Individual chronic care improvement
10	PLANS.—
11	"(A) In GENERAL.—For each beneficiary
12	participating in a program of a contractor
13	under this section, the contractor shall develop
14	with the beneficiary an individualized, goal-ori-
15	ented chronic care improvement plan.
16	"(B) Elements of individual plan.—
17	Each individual plan developed under subpara-
18	graph (A) shall include a single point of contact
19	to coordinate care and the following, as appro-
20	priate:
21	"(i) Self-improvement education for
22	the beneficiary and support education for
23	health care providers, primary caregivers,
24	and family members.



1	"(ii) Coordination of health care serv-
2	ices, such as application of a prescription
3	drug regimen and home health services.
4	"(iii) Collaboration with physicians
5	and other providers to enhance communica-
6	tion of relevant clinical information.
7	"(iv) The use of monitoring tech-
8	nologies that enable patient guidance
9	through the exchange of pertinent clinical
10	information, such as vital signs, sympto-
11	matic information, and health self-assess-
12	ment.
13	"(v) The provision of information
14	about hospice care, pain and palliative
15	care, and end-of-life care.
16	"(C) Contractor responsibilities.—In
17	establishing and carrying out individual plans
18	under a program, a contractor shall, directly or
19	through subcontractors—
20	"(i) guide participants in managing
21	their health, including all their co-
22	morbidities, and in performing activities as
23	specified under the elements of the plan:



1	"(ii) use decision support tools such as
2	evidence-based practice guidelines or other
3	criteria as determined by the Secretary; and
4	"(iii) develop a clinical information
5	database to track and monitor each partici-
6	pant across settings and to evaluate out-
7	comes.
8	"(6) Additional requirements.—The Sec-
9	retary may establish additional requirements for pro-
10	grams and contractors under this section.
11	"(7) Accreditation.—The Secretary may pro-
12	vide that programs that are accredited by qualified
13	organizations may be deemed to meet such require-
14	ments under this section as the Secretary may speci-
15	fy.
16	"(c) Contract Terms.—
17	"(1) In general.—A contract under this section
18	shall contain such terms and conditions as the Sec-
19	retary may specify consistent with this section. The
20	Secretary may not enter into a contract with an enti-
21	ty under this section unless the entity meets such clin-
22	ical, quality improvement, financial, and other re-
23	quirements as the Secretary deems to be appropriate

for the population to be served.



1	"(2) Use of subcontractors permitted.—A
2	contractor may carry out a program directly or
3	through contracts with subcontractors.
4	"(3) Budget neutral payment condition.—
5	In entering into a contract with an entity under this
6	subsection, the Secretary shall establish payment rates
7	that assure that there will be no net aggregate in-
8	crease in payments under this title over any period
9	of 3 years or longer, as agreed to by the Secretary.
10	Under this section, the Secretary shall assure that
11	medicare program outlays plus administrative ex-
12	penses (that would not have been paid under this title
13	without implementation of this section), including
14	contractor fees, shall not exceed the expenditures that
15	would have been incurred under this title for a com-
16	parable population in the absence of the program
17	under this section for the 3-year contract period.
18	"(4) At risk relationship.—For purposes of
19	section $1128B(b)(3)(F)$ , a contract under this section
20	shall be treated as a risk-sharing arrangement re-
21	ferred to in such section.
22	"(5) Performance standards.—Payment to
23	contractors under this section shall be subject to the
24	contractor's meeting of clinical and financial ner-

 $formance\ standards\ set\ by\ the\ Secretary.$ 



1	"(6) Contractor outcomes report.—Each
2	contractor offering a program shall monitor and re-
3	port to the Secretary, in a manner specified by the
4	Secretary, the quality of care and efficacy of such
5	program in terms of—
6	"(A) process measures, such as reductions
7	in errors of treatment and rehospitalization
8	rates;
9	"(B) beneficiary and provider satisfaction;
10	"(C) health outcomes; and
11	$``(D)\ financial\ outcomes.$
12	"(7) Phased in implementation.—Nothing in
13	this section shall be construed as preventing the Sec-
14	retary from phasing in the implementation of pro-
15	grams.
16	"(d) Biannual Outcomes Reports.—The Secretary
17	shall submit to the Congress biannual reports on the imple-
18	mentation of this section. Each such report shall include
19	information on—
20	"(1) the scope of implementation (in terms of
21	both regions and chronic conditions);
22	"(2) program design; and
23	"(3) improvements in health outcomes and fi-
24	nancial efficiencies that result from such implementa-
25	tion.



1	"(e) CLINICAL TRIALS.—The Secretary shall conduct
2	randomized clinical trials, that compare program partici-
3	pants with medicare beneficiaries who are offered, but de-
4	cline, to participate, in order to assess the potential of pro-
5	grams to—
6	"(1) reduce costs under this title; and
7	"(2) improve health outcomes under this title.
8	"(f) Authorization of Appropriations.—There are
9	authorized to be appropriated to the Secretary, in appro-
10	priate part from the Hospital Insurance Trust Fund and
11	the Supplementary Medical Insurance Trust Fund, such
12	sums as may be necessary to provide for contracts with
13	chronic care improvement programs under this section.
14	"(g) Limitation on Funding.—In no case shall the
15	funding under this section exceed \$100,000,000 over a pe-
16	riod of 3 years.".
17	SEC. 722. CHRONIC CARE IMPROVEMENT UNDER MEDICARE
18	ADVANTAGE AND ENHANCED FEE-FOR-SERV-
19	ICE PROGRAMS.
20	(a) Under Medicare Advantage Program.—Sec-
21	tion 1852 (42 U.S.C. 1395w-22) is amended—
22	(1) by amending subsection (e) to read as fol-
23	lows:



1	"(e) Implementation of Chronic Care Improve-
2	MENT PROGRAMS FOR BENEFICIARIES WITH MULTIPLE OR
3	Sufficiently Severe Chronic Conditions.—
4	"(1) In General.—Each Medicare Advantage
5	organization with respect to each Medicare Advantage
6	plan it offers shall have in effect, for enrollees with
7	multiple or sufficiently severe chronic conditions, a
8	chronic care improvement program that is designed to
9	manage the needs of such enrollees and that meets the
10	requirements of this subsection.
11	"(2) Enrollee with multiple or suffi-
12	CIENTLY SEVERE CHRONIC CONDITIONS.—For pur-
13	poses of this subsection, the term 'enrollee with mul-
14	tiple or sufficiently severe chronic conditions' means,
15	with respect to an enrollee in a Medicare Advantage
16	plan of a Medicare Advantage organization, an en-
17	rollee in the plan who has one or more chronic condi-
18	tions, such as congestive heart failure, diabetes,
19	COPD, stroke, or other disease as identified by the or-
20	ganization as appropriate for chronic care improve-
21	ment.
22	"(3) General requirements.—
23	"(A) In general.—Each chronic care im-
24	provement program under this subsection shall

be conducted consistent with this subsection.



1	"(B) Identification of enrollees.—
2	Each such program shall have a method for
3	monitoring and identifying enrollees with mul-
4	tiple or sufficiently severe chronic conditions
5	that meet the organization's criteria for partici-
6	pation under the program.
7	"(C) Development of plans.—For an en-
8	rollee identified under subparagraph (B) for par-
9	ticipation in a program, the program shall de-
10	velop, with the enrollee's consent, an individual-
11	ized, goal-oriented chronic care improvement
12	plan for chronic care improvement.
13	"(D) Elements of plans.—Each chronic
14	care improvement plan developed under subpara-
15	graph (C) shall include a single point of contact
16	to coordinate care and the following, as appro-
17	priate:
18	"(i) Self-improvement education for
19	the enrollee and support education for
20	health care providers, primary caregivers,
21	and family members.
22	"(ii) Coordination of health care serv-
23	ices, such as application of a prescription
24	drug regimen and home health services.



1	"(iii) Collaboration with physicians
2	and other providers to enhance communica-
3	tion of relevant clinical information.
4	"(iv) The use of monitoring tech-
5	nologies that enable patient guidance
6	through the exchange of pertinent clinical
7	information, such as vital signs, sympto-
8	matic information, and health self-assess-
9	ment.
10	"(v) The provision of information
11	about hospice care, pain and palliative
12	care, and end-of-life care.
13	"(E) Organization responsibilities.—
14	In establishing and carrying out chronic care
15	improvement plans for participants under this
16	paragraph, a Medicare Advantage organization
17	shall, directly or through subcontractors—
18	"(i) guide participants in managing
19	their health, including all their co-
20	morbidities, and in performing the activi-
21	ties as specified under the elements of the
22	plan;
23	"(ii) use decision support tools such as
24	evidence-based practice guidelines or other
25	criteria as determined by the Secretary; and



1	"(iii) develop a clinical information
2	database to track and monitor each partici-
3	pant across settings and to evaluate out-
4	comes.
5	"(3) Additional requirements.—The Sec-
6	retary may establish additional requirements for
7	chronic care improvement programs under this sec-
8	tion.
9	"(4) Accreditation.—The Secretary may pro-
10	vide that chronic care improvement programs that
11	are accredited by qualified organizations may be
12	deemed to meet such requirements under this sub-
13	section as the Secretary may specify.
14	"(5) Outcomes report.—Each Medicare Ad-
15	vantage organization with respect to its chronic care
16	improvement program under this subsection shall
17	monitor and report to the Secretary information on
18	the quality of care and efficacy of such program as
19	the Secretary may require."; and
20	(2) by amending subparagraph (I) of subsection
21	(c)(1) to read as follows:
22	"(I) Chronic care improvement pro-
23	GRAM.—A description of the organization's
24	chronic care improvement program under sub-
25	section (e).".



1	(b) Application under Enhanced Fee-for-Serv-
2	ICE Program.—Section 1860E-2(c)(3), as inserted by sec-
3	tion 201(a), is amended by inserting ", including subsection
4	(e) (relating to implementation of chronic care improve-
5	ment programs)" after "The provisions of section 1852".
6	(c) Effective Date.—The amendments made by this
7	section shall apply for contract years beginning on or after
8	1 year after the date of the enactment of this Act.
9	SEC. 723. INSTITUTE OF MEDICINE REPORT.
10	(a) Study.—
11	(1) In General.—The Secretary of Health and
12	Human Services shall contract with the Institute of
13	Medicine of the National Academy of Sciences to con-
14	duct a study of the barriers to effective integrated care
15	improvement for medicare beneficiaries with multiple
16	or severe chronic conditions across settings and over
17	time and to submit a report under subsection (b).
18	(2) Specific items.—The study shall examine
19	the statutory and regulatory barriers to coordinating
20	care across settings for medicare beneficiaries in tran-
21	sition from one setting to another (such as between
22	hospital, nursing facility, home health, hospice, and
23	home). The study shall specifically identify the fol-



lowing:

1	(A) Clinical, financial, or administrative
2	requirements in the medicare program that
3	present barriers to effective, seamless transitions
4	across care settings.
5	(B) Policies that impede the establishment
6	of administrative and clinical information sys-
7	tems to track health status, utilization, cost, and
8	quality data across settings.
9	(C) State-level requirements that may
10	present barriers to better care for medicare bene-
11	ficiaries.
12	(3) Consultation.—The study under this sub-
13	section shall be conducted in consultation with experts
14	in the field of chronic care, consumers, and family
15	caregivers, working to integrate care delivery and cre-
16	ate more seamless transitions across settings and over
17	time.
18	(b) Report.—The report under this subsection shall
19	be submitted to the Secretary and Congress not later than
20	18 months after the date of the enactment of this Act.
21	SEC. 724. MEDPAC REPORT.
22	(a) Evaluation.—shall conduct an evaluation that
23	includes a description of the status of the implementation
24	of chronic care improvement programs under section 1808

25 of the Social Security Act, the quality of health care services



- 1 provided to individuals in such program, the health status
- 2 of the participants of such program, and the cost savings
- 3 attributed to implementation of such program.
- 4 (b) Report.—Not later than 2 years after the date
- 5 of implementation of such chronic care improvement pro-
- 6 grams, the Commission shall submit a report on such eval-
- 7 uation.

## 8 Subtitle D—Other Provisions

- 9 SEC. 731. MODIFICATIONS TO MEDICARE PAYMENT ADVI-
- 10 **SORY COMMISSION (MEDPAC).**
- 11 (a) Examination of Budget Consequences.—Sec-
- 12 tion 1805(b) (42 U.S.C. 1395b-6(b)) is amended by adding
- 13 at the end the following new paragraph:
- 14 "(8) Examination of budget con-
- 15 SEQUENCES.—Before making any recommendations,
- 16 the Commission shall examine the budget con-
- 17 sequences of such recommendations, directly or
- 18 through consultation with appropriate expert enti-
- 19 *ties.*".
- 20 (b) Consideration of Efficient Provision of
- 21 Services.—Section 1805(b)(2)(B)(i) (42 U.S.C. 1395b-
- 22 6(b)(2)(B)(i) is amended by inserting "the efficient provi-
- 23 sion of" after "expenditures for".
- 24 (c) Application of Disclosure Requirements.—



1	(1) In General.—Section $1805(c)(2)(D)$ (42)
2	$U.S.C.\ 1395b-6(c)(2)(D))$ is amended by adding at
3	the end the following: "Members of the Commission
4	shall be treated as employees of the Congress for pur-
5	poses of applying title I of the Ethics in Government
6	Act of 1978 (Public Law 95-521).".
7	(2) Effective date.—The amendment made by
8	paragraph (1) shall take effect on January 1, 2004.
9	(d) Additional Reports.—
10	(1) Data needs and sources.—The Medicare
11	Payment Advisory Commission shall conduct a study,
12	and submit a report to Congress by not later than
13	June 1, 2004, on the need for current data, and
14	sources of current data available, to determine the sol-
15	vency and financial circumstances of hospitals and
16	other medicare providers of services. The Commission
17	shall examine data on uncompensated care, as well as
18	the share of uncompensated care accounted for by the
19	expenses for treating illegal aliens.
20	(2) Use of tax-related returns.—Using re-
21	turn information provided under Form 990 of the In-
22	ternal Revenue Service, the Commission shall submit
23	to Congress, by not later than June 1, 2004, a report



on the following:

1	(A) Investments, endowments, and fund-	
2	raising of hospitals participating under the	
3	medicare program and related foundations.	
4	(B) Access to capital financing for private	
5	and for not-for-profit hospitals.	
6	SEC. 732. DEMONSTRATION PROJECT FOR MEDICAL ADULT	
7	DAY CARE SERVICES.	
8	(a) Establishment.—Subject to the succeeding provi-	
9	sions of this section, the Secretary of Health and Human	
10	Services shall establish a demonstration project (in this sec-	
11	tion referred to as the "demonstration project") under	
12	which the Secretary shall, as part of a plan of an episode	
13	of care for home health services established for a medicare	
14	beneficiary, permit a home health agency, directly or under	
15	arrangements with a medical adult day care facility, to	
16	provide medical adult day care services as a substitute for	
17	a portion of home health services that would otherwise be	
18	provided in the beneficiary's home.	
19	(b) Payment.—	
20	(1) In General.—The amount of payment for	
21	an episode of care for home health services, a portion	
22	of which consists of substitute medical adult day care	
23	services, under the demonstration project shall be	
24	made at a rate equal to 95 percent of the amount that	
25	would otherwise apply for such home health services	



1	under section 1895 of the Social Security Act (42
2	u.s.c. 1395fff). In no case may a home health agency,
3	or a medical adult day care facility under arrange-
4	ments with a home health agency, separately charge
5	a beneficiary for medical adult day care services fur-
6	nished under the plan of care.
7	(2) Budget neutrality for demonstration
8	PROJECT.—Notwithstanding any other provision of
9	law, the Secretary shall provide for an appropriate

PROJECT.—Notwithstanding any other provision of law, the Secretary shall provide for an appropriate reduction in the aggregate amount of additional payments made under section 1895 of the Social Security Act (42 U.S.C. 1395fff) to reflect any increase in amounts expended from the Trust Funds as a result of the demonstration project conducted under this section.

- 16 (c) Demonstration Project Sites.—The project es-17 tablished under this section shall be conducted in not more 18 than 5 States selected by the Secretary that license or certify 19 providers of services that furnish medical adult day care 20 services.
- 21 (d) DURATION.—The Secretary shall conduct the dem-22 onstration project for a period of 3 years.
- 23 (e) VOLUNTARY PARTICIPATION.—Participation of 24 medicare beneficiaries in the demonstration project shall be 25 voluntary. The total number of such beneficiaries that may



participate in the project at any given time may not exceed 2 15,000. 3 (f) Preference in Selecting Agencies.—In selecting home health agencies to participate under the dem-5 onstration project, the Secretary shall give preference to those agencies that are currently licensed or certified 6 through common ownership and control to furnish medical 8 adult day care services. 9 (q) Waiver Authority.—The Secretary may waive 10 such requirements of title XVIII of the Social Security Act as may be necessary for the purposes of carrying out the 12 demonstration project, other than waiving the requirement that an individual be homebound in order to be eligible for benefits for home health services. 14 15 (h) Evaluation and Report.—The Secretary shall conduct an evaluation of the clinical and cost effectiveness 16 of the demonstration project. Not later 30 months after the 17 commencement of the project, the Secretary shall submit to 18 19 Congress a report on the evaluation, and shall include in 20 the report the following: 21 (1) An analysis of the patient outcomes and costs 22 of furnishing care to the medicare beneficiaries par-23 ticipating in the project as compared to such out-24 comes and costs to beneficiaries receiving only home

health services for the same health conditions.



1	(2) Such recommendations regarding the exten-
2	sion, expansion, or termination of the project as the
3	Secretary determines appropriate.
4	(i) Definitions.—In this section:
5	(1) Home Health Agency.—The term "home
6	health agency" has the meaning given such term in
7	section 1861(o) of the Social Security Act (42 U.S.C.
8	1395x(0)).
9	(2) Medical adult day care facility.—The
10	term "medical adult day care facility" means a facil-
11	ity that—
12	(A) has been licensed or certified by a State
13	to furnish medical adult day care services in the
14	State for a continuous 2-year period;
15	(B) is engaged in providing skilled nursing
16	services and other therapeutic services directly or
17	under arrangement with a home health agency;
18	(C) meets such standards established by the
19	Secretary to assure quality of care and such
20	other requirements as the Secretary finds nec-
21	essary in the interest of the health and safety of
22	individuals who are furnished services in the fa-
23	cility; and
24	(D) provides medical adult day care serv-
25	ices.



1	(3) Medical adult day care services.—The	
2	term "medical adult day care services" means—	
3	(A) home health service items and services	
4	described in paragraphs (1) through (7) of sec-	
5	tion 1861(m) furnished in a medical adult day	
6	$care\ facility;$	
7	(B) a program of supervised activities fur-	
8	nished in a group setting in the facility that—	
9	(i) meet such criteria as the Secretary	
10	determines appropriate; and	
11	(ii) is designed to promote physical	
12	and mental health of the individuals; and	
13	(C) such other services as the Secretary may	
14	specify.	
15	(4) Medicare beneficiary.—The term "medi-	
16	care beneficiary" means an individual entitled to	
17	benefits under part A of this title, enrolled under part	
18	B of this title, or both.	
19	SEC. 733. IMPROVEMENTS IN NATIONAL AND LOCAL COV-	
20	ERAGE DETERMINATION PROCESS TO RE-	
21	SPOND TO CHANGES IN TECHNOLOGY.	
22	(a) National and Local Coverage Determination	
23	Process.—	
24	(1) In General.—Section 1862 (42 U.S.C.	
25	1395y) is amended—	



1	(A) in the third sentence of subsection (a)
2	by inserting "consistent with subsection (k)"
3	after "the Secretary shall ensure"; and
4	(B) by adding at the end the following new
5	subsection:
6	"(k) National and Local Coverage Determina-
7	TION PROCESS.—
8	"(1) Criteria and evidence used in making
9	NATIONAL COVERAGE DETERMINATIONS.—The Sec-
10	retary shall make available to the public the criteria
11	the Secretary uses in making national coverage deter-
12	minations, including how evidence to demonstrate
13	that a procedure or device is reasonable and necessary
14	$is\ considered.$
15	"(2) Timeframe for decisions on requests
16	FOR NATIONAL COVERAGE DETERMINATIONS.—In the
17	case of a request for a national coverage determina-
18	tion that—
19	"(A) does not require a technology assess-
20	ment from an outside entity or deliberation from
21	the Medicare Coverage Advisory Committee, the
22	decision on the request shall be made not later
23	than 6 months after the date of the request; or
24	"(B) requires such an assessment or delib-
25	eration and in which a clinical trial is not re-



1	quested, the decision on the request shall be made
2	not later than 12 months after the date of the re-
3	quest.
4	"(3) Process for public comment in Na-
5	TIONAL COVERAGE DETERMINATIONS.—At the end of
6	the 6-month period that begins on the date a request
7	for a national coverage determination is made, the
8	Secretary shall—
9	"(A) make a draft of proposed decision on
10	the request available to the public through the
11	Medicare Internet site of the Department of
12	Health and Human Services or other appro-
13	priate means;
14	"(B) provide a 30-day period for public
15	comment on such draft;
16	"(C) make a final decision on the request
17	within 60 days of the conclusion of the 30-day
18	period referred to under subparagraph (B);
19	"(D) include in such final decision sum-
20	maries of the public comments received and re-
21	$sponses\ thereto;$
22	"(E) make available to the public the clin-
23	ical evidence and other data used in making
24	such a decision when the decision differs from



1	the recommendations of the Medicare Coverage
2	Advisory Committee; and
3	"(F) in the case of a decision to grant the
4	coverage determination, assign a temporary or
5	permanent code during the 60-day period re-
6	ferred to in subparagraph (C).
7	"(4) Consultation with outside experts in
8	CERTAIN NATIONAL COVERAGE DETERMINATIONS.—
9	With respect to a request for a national coverage de-
10	termination for which there is not a review by the
11	Medicare Coverage Advisory Committee, the Secretary
12	shall consult with appropriate outside clinical ex-
13	perts.
14	"(5) Local coverage determination proc-
15	ESS.—With respect to local coverage determinations
16	made on or after January 1, 2004—
17	"(A) Plan to promote consistency of
18	COVERAGE DETERMINATIONS.—The Secretary
19	shall develop a plan to evaluate new local cov-
20	erage determinations to determine which deter-
21	minations should be adopted nationally and to
22	what extent greater consistency can be achieved
23	among local coverage determinations.
24	"(B) Consultation.—The Secretary shall
25	require the fiscal intermediaries or carriers pro-



1	viding services within the same area to consult
2	on all new local coverage determinations within
3	the area.
4	"(C) Dissemination of information.—
5	The Secretary should serve as a center to dis-
6	seminate information on local coverage deter-
7	minations among fiscal intermediaries and car-
8	riers to reduce duplication of effort.
9	"(6) National and local coverage deter-
10	MINATION DEFINED.—For purposes of this subsection,
11	the terms 'national coverage determination' and 'local
12	coverage determination' have the meaning given such
13	terms in paragraphs $(1)(B)$ and $(2)(B)$ , respectively,
14	of section 1869(f).".
15	(2) Effective date.—The amendments made
16	by paragraph (1) shall apply to national and local
17	coverage determinations as of January 1, 2004.
18	(b) Medicare Coverage of Routine Costs Associ-
19	ATED WITH CERTAIN CLINICAL TRIALS.—
20	(1) In general.—With respect to the coverage of
21	routine costs of care for beneficiaries participating in
22	a qualifying clinical trial, as set forth on the date of
23	the enactment of this Act in National Coverage Deter-
24	mination 30-1 of the Medicare Coverage Issues Man-

ual, the Secretary shall deem clinical trials conducted



1	in accordance with an investigational device exemp-
2	tion approved under section 520(g) of the Federal
3	Food, Drug, and Cosmetic Act (42 U.S.C. 360j(g)) to
4	be automatically qualified for such coverage.
5	(2) Rule of construction.—Nothing in this
6	subsection shall be construed as authorizing or requir-
7	ing the Secretary to modify the regulations set forth
8	on the date of the enactment of this Act at subpart
9	B of part 405 of title 42, Code of Federal Regulations,
10	or subpart A of part 411 of such title, relating to cov-
11	erage of, and payment for, a medical device that is
12	the subject of an investigational device exemption by
13	the Food and Drug Administration (except as may be
14	necessary to implement paragraph (1)).
15	(3) Effective date.—This subsection shall
16	apply to clinical trials begun before, on, or after the
17	date of the enactment of this Act and to items and
18	services furnished on or after such date.
19	(c) Issuance of Temporary National Codes.—Not
20	later than January 1, 2004, the Secretary shall implement
21	revised procedures for the issuance of temporary national
22	HCPCS codes under part B of title XVIII of the Social Se-
23	$curity\ Act.$



1	SEC. 734. TREATMENT OF CERTAIN PHYSICIAN PATHOLOGY
2	SERVICES.
3	(a) In General.—Section 1848(i) (42 U.S.C. 1395w-
4	4(i)) is amended by adding at the end the following new
5	paragraph:
6	"(4) Treatment of certain inpatient physi-
7	CIAN PATHOLOGY SERVICES.—
8	"(A) In General.—With respect to services
9	furnished on or after January 1, 2001, and be-
10	fore January 1, 2006, if an independent labora-
11	tory furnishes the technical component of a phy-
12	sician pathology service to a fee-for-service medi-
13	care beneficiary who is an inpatient or out-
14	patient of a covered hospital, the Secretary shall
15	treat such component as a service for which pay-
16	ment shall be made to the laboratory under this
17	section and not as an inpatient hospital service
18	for which payment is made to the hospital under
19	section 1886(d) or as a hospital outpatient serv-
20	ice for which payment is made to the hospital
21	under section $1833(t)$ .
22	"(B) Definitions.—In this paragraph:
23	"(i) Covered hospital.—
24	"(I) In general.—The term 'cov-
25	ered hospital' means, with respect to
26	an inpatient or outpatient, a hospital



1	that had an arrangement with an
2	independent laboratory that was in ef-
3	fect as of July 22, 1999, under which
4	a laboratory furnished the technical
5	component of physician pathology serv-
6	ices to fee-for-service medicare bene-
7	ficiaries who were hospital inpatients
8	or outpatients, respectively, and sub-
9	mitted claims for payment for such
10	component to a carrier with a contract
11	under section 1842 and not to the hos-
12	pital.
13	"(II) Change in ownership
14	DOES NOT AFFECT DETERMINATION.—
15	A change in ownership with respect to
16	a hospital on or after the date referred
17	to in subclause (I) shall not affect the
18	determination of whether such hospital
19	is a covered hospital for purposes of
20	such subclause.
21	"(ii) Fee-for-service medicare
22	BENEFICIARY.—The term 'fee-for-service
23	medicare beneficiary' means an individual

who is entitled to benefits under part A, or



1	enrolled under this part, or both, but is not
2	enrolled in any of the following:
3	``(I)  A  Medicare + Choice  plan
4	under part C.
5	"(II) A plan offered by an eligible
6	organization under section 1876.
7	"(III) A program of all-inclusive
8	care for the elderly (PACE) under sec-
9	$tion\ 1894.$
10	"(IV) A social health maintenance
11	organization (SHMO) demonstration
12	project established under section
13	4018(b) of the Omnibus Budget Rec-
14	onciliation Act of 1987 (Public Law
15	100–203).".
16	(b) Conforming Amendment.—Section 542 of the
17	Medicare, Medicaid, and SCHIP Benefits Improvement
18	and Protection Act of 2000 (114 Stat. 2763A-550), as en-
19	acted into law by section 1(a)(6) of Public Law 106-554,
20	is repealed.
21	(c) Effective Dates.—The amendments made by
22	this section shall take effect as if included in the enactment
23	of the Medicare, Medicaid, and SCHIP Benefits Improve-
24	ment and Protection Act of 2000 (Amendix F 114 Stat



1	2763A-463), as enacted into law by section 1(a)(6) of Pub-
2	lic Law 106–554.
3	TITLE VIII—MEDICARE
4	BENEFITS ADMINISTRATION
5	SEC. 801. ESTABLISHMENT OF MEDICARE BENEFITS ADMIN-
6	ISTRATION.
7	(a) In General.—Title XVIII (42 U.S.C. 1395 et
8	seq.), as amended by sections 105 and 721, is amended by
9	inserting after 1808 the following new section:
10	"MEDICARE BENEFITS ADMINISTRATION
11	"Sec. 1809. (a) Establishment.—There is estab-
12	lished within the Department of Health and Human Serv-
13	ices an agency to be known as the Medicare Benefits Admin-
14	istration.
15	"(b) Administrator; Deputy Administrator;
16	Chief Actuary.—
17	"(1) Administrator.—
18	"(A) In General.—The Medicare Benefits
19	Administration shall be headed by an adminis-
20	trator to be known as the 'Medicare Benefits Ad-
21	ministrator' (in this section referred to as the
22	'Administrator') who shall be appointed by the
23	President, by and with the advice and consent of
24	the Senate. The Administrator shall be in direct
25	line of authority to the Secretary.



1	"(B) Compensation.—The Administrator
2	shall be paid at the rate of basic pay payable for
3	level III of the Executive Schedule under section
4	5314 of title 5, United States Code.
5	"(C) Term of office.—The Administrator
6	shall be appointed for a term of 4 years. In any
7	case in which a successor does not take office at
8	the end of an Administrator's term of office, that
9	Administrator may continue in office until the
10	entry upon office of such a successor. An Admin-
11	istrator appointed to a term of office after the
12	commencement of such term may serve under
13	such appointment only for the remainder of such
14	term.
15	"(D) General authority.—The Adminis-
16	trator shall be responsible for the exercise of all
17	powers and the discharge of all duties of the Ad-
18	ministration, and shall have authority and con-
19	trol over all personnel and activities thereof.
20	"(E) Rulemaking authority.—The Ad-
21	ministrator may prescribe such rules and regula-
22	tions as the Administrator determines necessary
23	or appropriate to carry out the functions of the
24	Administration. The regulations prescribed by

the Administrator shall be subject to the rule-



1	making procedures established under section 553
2	of title 5, United States Code. The Administrator
3	shall provide for the issuance of new regulations
4	to carry out parts C, D, and E.
5	"(F) Authority to establish organiza-
6	TIONAL UNITS.—The Administrator may estab-
7	lish, alter, consolidate, or discontinue such orga-
8	nizational units or components within the Ad-
9	ministration as the Administrator considers nec-
10	essary or appropriate, except as specified in this
11	section.
12	"(G) Authority to delegate.—The Ad-
13	ministrator may assign duties, and delegate, or
14	authorize successive redelegations of, authority to
15	act and to render decisions, to such officers and
16	employees of the Administration as the Adminis-
17	trator may find necessary. Within the limita-
18	tions of such delegations, redelegations, or as-
19	signments, all official acts and decisions of such
20	officers and employees shall have the same force
21	and effect as though performed or rendered by
22	$the \ Administrator.$
23	"(2) Deputy administrator.—
24	"(A) In General.—There shall be a Dep-

uty Administrator of the Medicare Benefits Ad-



1	ministration who shall be appointed by the
2	President, by and with the advice and consent of
3	the Senate.
4	"(B) Compensation.—The Deputy Admin-
5	istrator shall be paid at the rate of basic pay
6	payable for level IV of the Executive Schedule
7	under section 5315 of title 5, United States Code.
8	"(C) Term of office.—The Deputy Ad-
9	ministrator shall be appointed for a term of 4
10	years. In any case in which a successor does not
11	take office at the end of a Deputy Administra-
12	tor's term of office, such Deputy Administrator
13	may continue in office until the entry upon of-
14	fice of such a successor. A Deputy Administrator
15	appointed to a term of office after the commence-
16	ment of such term may serve under such ap-
17	pointment only for the remainder of such term.
18	"(D) Duties.—The Deputy Administrator
19	shall perform such duties and exercise such pow-
20	ers as the Administrator shall from time to time
21	assign or delegate. The Deputy Administrator
22	shall be Acting Administrator of the Administra-
23	tion during the absence or disability of the Ad-
24	ministrator and, unless the President designates

another officer of the Government as Acting Ad-



1	ministrator, in the event of a vacancy in the of-
2	fice of the Administrator.
3	"(3) Chief actuary.—
4	"(A) In general.—There is established in
5	the Administration the position of Chief Actu-
6	ary. The Chief Actuary shall be appointed by,
7	and in direct line of authority to, the Adminis-
8	trator of such Administration. The Chief Actu-
9	ary shall be appointed from among individuals
10	who have demonstrated, by their education and
11	experience, superior expertise in the actuarial
12	sciences. The Chief Actuary may be removed only
13	for cause.
14	"(B) Compensation.—The Chief Actuary
15	shall be compensated at the highest rate of basic
16	pay for the Senior Executive Service under sec-
17	tion 5382(b) of title 5, United States Code.
18	"(C) Duties.—The Chief Actuary shall ex-
19	ercise such duties as are appropriate for the of-
20	fice of the Chief Actuary and in accordance with
21	professional standards of actuarial independence.
22	"(4) Secretarial coordination of program
23	ADMINISTRATION.—The Secretary shall ensure appro-
24	priate coordination between the Administrator and

the Administrator of the Centers for Medicare & Med-



1	icaid Services in carrying out the programs under
2	$this\ title.$
3	"(c) Duties; Administrative Provisions.—
4	"(1) Duties.—
5	"(A) General duties.—The Adminis-
6	trator shall carry out parts C, D, and E,
7	including—
8	"(i) negotiating, entering into, and en-
9	forcing, contracts with plans for the offering
10	of Medicare Advantage plans under part C
11	and EFFS plans under part E, including
12	the offering of qualified prescription drug
13	coverage under such plans; and
14	"(ii) negotiating, entering into, and
15	enforcing, contracts with PDP sponsors for
16	the offering of prescription drug plans
17	under part D.
18	"(B) Other duties.—The Administrator
19	shall carry out any duty provided for under part
20	C, part D, or part E, including demonstration
21	projects carried out in part or in whole under
22	such parts, the programs of all-inclusive care for
23	the elderly (PACE program) under section 1894,
24	the social health maintenance organization
25	(SHMO) demonstration projects (referred to in



1	section 4104(c) of the Balanced Budget Act of
2	1997), medicare cost contractors under section
3	1876(h), and through a Medicare Advantage
4	project that demonstrates the application of capi-
5	tation payment rates for frail elderly medicare
6	beneficiaries through the use of a interdiscipli-
7	nary team and through the provision of primary
8	care services to such beneficiaries by means of
9	such a team at the nursing facility involved).
10	"(C) Prescription drug card.—The Ad-
11	ministrator shall carry out section 1807 (relat-
12	ing to the medicare prescription drug discount
13	card endorsement program).
14	"(D) Noninterference.—In carrying out
15	its duties with respect to the provision of quali-
16	fied prescription drug coverage to beneficiaries
17	under this title, the Administrator may not—
18	"(i) require a particular formulary or
19	institute a price structure for the reimburse-
20	ment of covered outpatient drugs;
21	"(ii) interfere in any way with nego-
22	tiations between PDP sponsors and Medi-
23	care Advantage organizations and EFFS

organizations and drug manufacturers,



1	wholesalers, or other suppliers of covered
2	outpatient drugs; and
3	"(iii) otherwise interfere with the com-
4	petitive nature of providing such coverage
5	through such sponsors and organizations.
6	"(E) Annual reports.—Not later March
7	31 of each year, the Administrator shall submit
8	to Congress and the President a report on the
9	administration of parts C, D, and E during the
10	previous fiscal year.
11	"(2) Staff.—
12	"(A) In General.—The Administrator,
13	with the approval of the Secretary, may employ,
14	without regard to chapter 31 of title 5, United
15	States Code, other than sections 3102 through
16	3108, 3110 through 3113, 3136m and 3151, such
17	officers and employees as are necessary to ad-
18	minister the activities to be carried out through
19	the Medicare Benefits Administration. The Ad-
20	ministrator shall employ staff with appropriate
21	and necessary expertise in negotiating contracts
22	in the private sector.
23	"(B) Flexibility with respect to com-



24

PENSATION.—

1	"(i) In General.—The staff of the
2	Medicare Benefits Administration shall,
3	subject to clause (ii), be paid without regard
4	to the provisions of chapter 51 (other than
5	section 5101) and chapter 53 (other than
6	section 5301) of such title (relating to clas-
7	sification and schedule pay rates).
8	"(ii) Maximum rate.—In no case
9	may the rate of compensation determined
10	under clause (i) exceed the rate of basic pay
11	payable for level IV of the Executive Sched-
12	ule under section 5315 of title 5, United
13	States Code.
14	"(C) Limitation on full-time equiva-
15	LENT STAFFING FOR CURRENT CMS FUNCTIONS
16	BEING TRANSFERRED.—The Administrator may
17	not employ under this paragraph a number of
18	full-time equivalent employees, to carry out func-
19	tions that were previously conducted by the Cen-
20	ters for Medicare & Medicaid Services and that
21	are conducted by the Administrator by reason of
22	this section, that exceeds the number of such full-
23	time equivalent employees authorized to be em-

ployed by the Centers for Medicare & Medicaid



1	Services to conduct such functions as of the date
2	of the enactment of this Act.
3	"(3) Redelegation of certain functions of
4	THE CENTERS FOR MEDICARE & MEDICAID SERV-
5	ICES.—
6	"(A) In general.—The Secretary, the Ad-
7	ministrator, and the Administrator of the Cen-
8	ters for Medicare & Medicaid Services shall es-
9	tablish an appropriate transition of responsi-
10	bility in order to redelegate the administration
11	of part C from the Secretary and the Adminis-
12	trator of the Centers for Medicare & Medicaid
13	Services to the Administrator as is appropriate
14	to carry out the purposes of this section.
15	"(B) Transfer of data and informa-
16	TION.—The Secretary shall ensure that the Ad-
17	ministrator of the Centers for Medicare & Med-
18	icaid Services transfers to the Administrator of
19	the Medicare Benefits Administration such infor-
20	mation and data in the possession of the Admin-
21	istrator of the Centers for Medicare & Medicaid
22	Services as the Administrator of the Medicare
23	Benefits Administration requires to carry out the

duties described in paragraph (1).



1	"(C) Construction.—Insofar as a respon-
2	sibility of the Secretary or the Administrator of
3	the Centers for Medicare & Medicaid Services is
4	redelegated to the Administrator under this sec-
5	tion, any reference to the Secretary or the Ad-
6	ministrator of the Centers for Medicare & Med-
7	icaid Services in this title or title XI with re-
8	spect to such responsibility is deemed to be a ref-
9	erence to the Administrator.
10	"(d) Office of Beneficiary Assistance.—
11	"(1) Establishment.—The Secretary shall es-
12	tablish within the Medicare Benefits Administration
13	an Office of Beneficiary Assistance to coordinate
14	functions relating to outreach and education of medi-
15	care beneficiaries under this title, including the func-
16	tions described in paragraph (2). The Office shall be
17	separate operating division within the Administra-
18	tion.
19	"(2) Dissemination of information on bene-
20	FITS AND APPEALS RIGHTS.—
21	"(A) Dissemination of Benefits infor-
22	MATION.—The Office of Beneficiary Assistance
23	shall disseminate, directly or through contract, to
24	medicare beneficiaries, by mail, by posting on

the Internet site of the Medicare Benefits Admin-



1	istration and through a toll-free telephone num-
2	ber, information with respect to the following:
3	"(i) Benefits, and limitations on pay-
4	ment (including cost-sharing, stop-loss pro-
5	visions, and formulary restrictions) under
6	parts C, D, and E.
7	"(ii) Benefits, and limitations on pay-
8	ment under parts A and B, including infor-
9	mation on medicare supplemental policies
10	under section 1882.
11	Such information shall be presented in a manner
12	so that medicare beneficiaries may compare ben-
13	efits under parts A, B, D, and medicare supple-
14	mental policies with benefits under Medicare Ad-
15	vantage plans under part C and EFFS plans
16	$under\ part\ E.$
17	"(B) Dissemination of appeals rights
18	Information.—The Office of Beneficiary Assist-
19	ance shall disseminate to medicare beneficiaries
20	in the manner provided under subparagraph (A)
21	a description of procedural rights (including
22	grievance and appeals procedures) of bene-
23	ficiaries under the original medicare fee-for-serv-
24	ice program under parts A and B, the Medicare

Advantage program under part C, the Voluntary



1	Prescription Drug Benefit Program under part
2	D, and the Enhanced Fee-for-Service program
3	$under\ part\ E.$
4	"(e) Medicare Policy Advisory Board.—
5	"(1) Establishment.—There is established
6	within the Medicare Benefits Administration the
7	Medicare Policy Advisory Board (in this section re-
8	ferred to the 'Board'). The Board shall advise, consult
9	with, and make recommendations to the Adminis-
10	trator of the Medicare Benefits Administration with
11	respect to the administration of parts C, D, and E,
12	including the review of payment policies under such
13	parts.
14	"(2) Reports.—
15	"(A) In general.—With respect to matters
16	of the administration of parts C, D, and E the
17	Board shall submit to Congress and to the Ad-
18	ministrator of the Medicare Benefits Administra-
19	tion such reports as the Board determines appro-
20	priate. Each such report may contain such rec-
21	ommendations as the Board determines appro-
22	priate for legislative or administrative changes
23	to improve the administration of such parts, in-

cluding the topics described in subparagraph



1	(B). Each such report shall be published in the
2	Federal Register.
3	"(B) Topics described.—Reports required
4	under subparagraph (A) may include the fol-
5	lowing topics:
6	"(i) Fostering competition.—Rec-
7	ommendations or proposals to increase com-
8	petition under parts C, D, and E for serv-
9	ices furnished to medicare beneficiaries.
10	"(ii) Education and enrollment.—
11	Recommendations for the improvement to
12	efforts to provide medicare beneficiaries in-
13	formation and education on the program
14	under this title, and specifically parts C, D,
15	and E, and the program for enrollment
16	under the title.
17	"(iii) Implementation of risk-ad-
18	Justment.—Evaluation of the implementa-
19	tion under section $1853(a)(3)(C)$ of the risk
20	adjustment methodology to payment rates
21	under that section to Medicare Advantage
22	organizations offering Medicare Advantage
23	plans (and the corresponding payment pro-
24	visions under part E) that accounts for

variations in per capita costs based on



1	health status, geography, and other demo-
2	graphic factors.
3	"(iv) Rural access.—Recommenda-
4	tions to improve competition and access to
5	plans under parts C, D, and E in rural
6	areas.
7	"(C) MAINTAINING INDEPENDENCE OF
8	BOARD.—The Board shall directly submit to
9	Congress reports required under subparagraph
10	(A). No officer or agency of the United States
11	may require the Board to submit to any officer
12	or agency of the United States for approval,
13	comments, or review, prior to the submission to
14	Congress of such reports.
15	"(3) Duty of administrator of medicare
16	Benefits administration.—With respect to any re-
17	port submitted by the Board under paragraph (2)(A),
18	not later than 90 days after the report is submitted,
19	the Administrator of the Medicare Benefits Adminis-
20	tration shall submit to Congress and the President and
21	analysis of recommendations made by the Board in
22	such report. Each such analysis shall be published in
23	the Federal Register.



1	"(A) Appointment.—Subject to the suc-
2	ceeding provisions of this paragraph, the Board
3	shall consist of seven members to be appointed as
4	follows:
5	"(i) Three members shall be appointed
6	by the President.
7	"(ii) Two members shall be appointed
8	by the Speaker of the House of Representa-
9	tives, with the advice of the chairmen and
10	the ranking minority members of the Com-
11	mittees on Ways and Means and on Energy
12	and Commerce of the House of Representa-
13	tives.
14	"(iii) Two members shall be appointed
15	by the President pro tempore of the Senate
16	with the advice of the chairman and the
17	ranking minority member of the Senate
18	Committee on Finance.
19	"(B) Qualifications.—The members shall
20	be chosen on the basis of their integrity, impar-
21	tiality, and good judgment, and shall be individ-
22	uals who are, by reason of their education and
23	experience in health care benefits management,
24	exceptionally qualified to perform the duties of

members of the Board.



1	"(C) Prohibition on inclusion of fed-
2	ERAL EMPLOYEES.—No officer or employee of the
3	United States may serve as a member of the
4	Board.
5	"(5) Compensation.—Members of the Board
6	shall receive, for each day (including travel time) they
7	are engaged in the performance of the functions of the
8	board, compensation at rates not to exceed the daily
9	equivalent to the annual rate in effect for level IV of
10	the Executive Schedule under section 5315 of title 5,
11	United States Code.
12	"(6) Terms of office.—
13	"(A) In general.—The term of office of
14	members of the Board shall be 3 years.
15	"(B) Terms of initial appointees.—As
16	designated by the President at the time of ap-
17	pointment, of the members first appointed—
18	"(i) one shall be appointed for a term
19	of 1 year;
20	"(ii) three shall be appointed for terms
21	of 2 years; and
22	"(iii) three shall be appointed for
23	terms of 3 years.



1	"(C) REAPPOINTMENTS.—Any person ap-
2	pointed as a member of the Board may not serve
3	for more than 8 years.
4	"(D) VACANCY.—Any member appointed to
5	fill a vacancy occurring before the expiration of
6	the term for which the member's predecessor was
7	appointed shall be appointed only for the re-
8	mainder of that term. A member may serve after
9	the expiration of that member's term until a suc-
10	cessor has taken office. A vacancy in the Board
11	shall be filled in the manner in which the origi-
12	nal appointment was made.
13	"(7) Chair.—The Chair of the Board shall be
14	elected by the members. The term of office of the Chair
15	shall be 3 years.
16	"(8) Meetings.—The Board shall meet at the
17	call of the Chair, but in no event less than three times
18	during each fiscal year.
19	"(9) Director and staff.—
20	"(A) Appointment of director.—The
21	Board shall have a Director who shall be ap-
22	pointed by the Chair.
23	"(B) In General.—With the approval of
24	the Board, the Director may appoint, without re-
25	gard to chapter 31 of title 5, United States Code,



1	such additional personnel as the Director con-
2	siders appropriate.
3	"(C) Flexibility with respect to com-
4	PENSATION.—
5	"(i) In General.—The Director and
6	staff of the Board shall, subject to clause
7	(ii), be paid without regard to the provi-
8	sions of chapter 51 and chapter 53 of such
9	title (relating to classification and schedule
10	pay rates).
11	"(ii) Maximum rate.—In no case
12	may the rate of compensation determined
13	under clause (i) exceed the rate of basic pay
14	payable for level IV of the Executive Sched-
15	ule under section 5315 of title 5, United
16	States Code.
17	"(D) Assistance from the adminis-
18	TRATOR OF THE MEDICARE BENEFITS ADMINIS-
19	TRATION.—The Administrator of the Medicare
20	Benefits Administration shall make available to
21	the Board such information and other assistance
22	as it may require to carry out its functions.
23	"(10) Contract authority.—The Board may
24	contract with and compensate government and pri-
25	vate agencies or persons to carry out its duties under



1	this subsection, without regard to section 3709 of the
2	Revised Statutes (41 U.S.C. 5).
3	"(f) Funding.—There is authorized to be appro-
4	priated, in appropriate part from the Federal Hospital In-
5	surance Trust Fund and from the Federal Supplementary
6	Medical Insurance Trust Fund (including the Medicare
7	Prescription Drug Account), such sums as are necessary to
8	carry out this section.".
9	(b) Effective Date.—
10	(1) In general.—The amendment made by sub-
11	section (a) shall take effect on the date of the enact-
12	ment of this Act.
13	(2) Duties with respect to eligibility de-
14	TERMINATIONS AND ENROLLMENT.—The Adminis-
15	trator of the Medicare Benefits Administration shall
16	carry out enrollment under title XVIII of the Social
17	Security Act, make eligibility determinations under
18	such title, and carry out parts C and E of such title
19	for years beginning or after January 1, 2006.
20	(3) Transition.—Before the date the Adminis-
21	trator of the Medicare Benefits Administration is ap-
22	pointed and assumes responsibilities under this sec-
23	tion and section 1807 of the Social Security Act, the

Secretary of Health and Human Services shall pro-



1	vide for the conduct of any responsibilities of such
2	Administrator that are otherwise provided under law.
3	(c) Miscellaneous Administrative Provisions.—
4	(1) Administrator as member of the board
5	OF TRUSTEES OF THE MEDICARE TRUST FUNDS.—
6	Section 1817(b) and section 1841(b) (42 U.S.C.
7	1395i(b), 1395t(b)) are each amended by striking
8	"and the Secretary of Health and Human Services,
9	all ex officio," and inserting "the Secretary of Health
10	and Human Services, and the Administrator of the
11	Medicare Benefits Administration, all ex officio,".
12	(2) Increase in grade to executive level
13	III FOR THE ADMINISTRATOR OF THE CENTERS FOR
14	MEDICARE & MEDICAID SERVICES; LEVEL FOR MEDI-
15	CARE BENEFITS ADMINISTRATOR.—
16	(A) In general.—Section 5314 of title 5,
17	United States Code, by adding at the end the fol-
18	lowing:
19	"Administrator of the Centers for Medicare &
20	Medicaid Services.
21	"Administrator of the Medicare Benefits Admin-
22	istration.".
23	(B) Conforming amendment.—Section
24	5315 of such title is amended by striking "Ad-



1	ministrator of the Health Care Financing Ad-
2	ministration.".
3	(C) Effective date.—The amendments
4	made by this paragraph take effect on January
5	1, 2004.
6	TITLE IX—REGULATORY REDUC-
7	TION AND CONTRACTING RE-
8	FORM
9	Subtitle A—Regulatory Reform
10	SEC. 901. CONSTRUCTION; DEFINITION OF SUPPLIER.
11	(a) Construction.—Nothing in this title shall be
12	construed—
13	(1) to compromise or affect existing legal rem-
14	edies for addressing fraud or abuse, whether it be
15	criminal prosecution, civil enforcement, or adminis-
16	trative remedies, including under sections 3729
17	through 3733 of title 31, United States Code (known
18	as the False Claims Act); or
19	(2) to prevent or impede the Department of
20	Health and Human Services in any way from its on-
21	going efforts to eliminate waste, fraud, and abuse in
22	the medicare program.
23	Furthermore, the consolidation of medicare administrative
24	contracting set forth in this Act does not constitute consoli-
25	dation of the Federal Hospital Insurance Trust Fund and



- 1 the Federal Supplementary Medical Insurance Trust Fund
- 2 or reflect any position on that issue.
- 3 (b) Definition of Supplier.—Section 1861 (42)
- 4 U.S.C. 1395x) is amended by inserting after subsection (c)
- 5 the following new subsection:
- 6 "Supplier
- 7 "(d) The term 'supplier' means, unless the context oth-
- 8 erwise requires, a physician or other practitioner, a facility,
- 9 or other entity (other than a provider of services) that fur-
- 10 nishes items or services under this title.".
- 11 SEC. 902. ISSUANCE OF REGULATIONS.
- 12 (a) Regular Timeline for Publication of Final
- 13 *Rules.*—
- 14 (1) In General.—Section 1871(a) (42 U.S.C.
- 15 1395hh(a)) is amended by adding at the end the fol-
- 16 lowing new paragraph:
- 17 "(3)(A) The Secretary, in consultation with the Direc-
- 18 tor of the Office of Management and Budget, shall establish
- 19 and publish a regular timeline for the publication of final
- 20 regulations based on the previous publication of a proposed
- 21 regulation or an interim final regulation.
- 22 "(B) Such timeline may vary among different regula-
- 23 tions based on differences in the complexity of the regula-
- 24 tion, the number and scope of comments received, and other
- 25 relevant factors, but shall not be longer than 3 years except



- 1 under exceptional circumstances. If the Secretary intends
- 2 to vary such timeline with respect to the publication of a
- 3 final regulation, the Secretary shall cause to have published
- 4 in the Federal Register notice of the different timeline by
- 5 not later than the timeline previously established with re-
- 6 spect to such regulation. Such notice shall include a brief
- 7 explanation of the justification for such variation.
- 8 "(C) In the case of interim final regulations, upon the
- 9 expiration of the regular timeline established under this
- 10 paragraph for the publication of a final regulation after
- 11 opportunity for public comment, the interim final regula-
- 12 tion shall not continue in effect unless the Secretary pub-
- 13 lishes (at the end of the regular timeline and, if applicable,
- 14 at the end of each succeeding 1-year period) a notice of con-
- 15 tinuation of the regulation that includes an explanation of
- 16 why the regular timeline (and any subsequent 1-year exten-
- 17 sion) was not complied with. If such a notice is published,
- 18 the regular timeline (or such timeline as previously ex-
- 19 tended under this paragraph) for publication of the final
- 20 regulation shall be treated as having been extended for 1
- 21 additional year.
- 22 "(D) The Secretary shall annually submit to Congress
- 23 a report that describes the instances in which the Secretary
- 24 failed to publish a final regulation within the applicable



1	regular timeline under this paragraph and that provides
2	an explanation for such failures.".
3	(2) Effective date.—The amendment made by
4	paragraph (1) shall take effect on the date of the en-
5	actment of this Act. The Secretary shall provide for
6	an appropriate transition to take into account the
7	backlog of previously published interim final regula-
8	tions.
9	(b) Limitations on New Matter in Final Regula-
10	TIONS.—
11	(1) In general.—Section 1871(a) (42 U.S.C.
12	1395hh(a)), as amended by subsection (a), is amended
13	by adding at the end the following new paragraph:
14	"(4) If the Secretary publishes a final regulation that
15	includes a provision that is not a logical outgrowth of a
16	previously published notice of proposed rulemaking or in-
17	terim final rule, such provision shall be treated as a pro-
18	posed regulation and shall not take effect until there is the
19	further opportunity for public comment and a publication
20	of the provision again as a final regulation.".
21	(2) Effective date.—The amendment made by
22	paragraph (1) shall apply to final regulations pub-
23	lished on or after the date of the enactment of this
24	Act.



1	SEC. 903. COMPLIANCE WITH CHANGES IN REGULATIONS
2	AND POLICIES.
3	(a) No Retroactive Application of Substantive
4	CHANGES.—
5	(1) In General.—Section 1871 (42 U.S.C.
6	1395hh), as amended by section 902(a), is amended
7	by adding at the end the following new subsection:
8	" $(e)(1)(A)$ A substantive change in regulations, man-
9	ual instructions, interpretative rules, statements of policy,
10	or guidelines of general applicability under this title shall
11	not be applied (by extrapolation or otherwise) retroactively
12	to items and services furnished before the effective date of
13	the change, unless the Secretary determines that—
14	"(i) such retroactive application is necessary to
15	comply with statutory requirements; or
16	"(ii) failure to apply the change retroactively
17	would be contrary to the public interest.".
18	(2) Effective date.—The amendment made by
19	paragraph (1) shall apply to substantive changes
20	issued on or after the date of the enactment of this
21	Act.
22	(b) Timeline for Compliance With Substantive
23	Changes After Notice.—
24	(1) In General.—Section 1871(e)(1), as added
25	by subsection (a), is amended by adding at the end
26	the following:



1 "(B)(i) Except as provided in clause (ii), a substantive 2 change referred to in subparagraph (A) shall not become 3 effective before the end of the 30-day period that begins on 4 the date that the Secretary has issued or published, as the 5 case may be, the substantive change. 6 "(ii) The Secretary may provide for such a substantive change to take effect on a date that precedes the end of the 8 30-day period under clause (i) if the Secretary finds that waiver of such 30-day period is necessary to comply with 10 statutory requirements or that the application of such 30day period is contrary to the public interest. If the Secretary provides for an earlier effective date pursuant to this 12 clause, the Secretary shall include in the issuance or publication of the substantive change a finding described in the 14 first sentence, and a brief statement of the reasons for such 16 finding. 17 "(C) No action shall be taken against a provider of 18 services or supplier with respect to noncompliance with 19 such a substantive change for items and services furnished before the effective date of such a change.". 20 21 (2) Effective date.—The amendment made by 22 paragraph (1) shall apply to compliance actions un-23 dertaken on or after the date of the enactment of this 24 Act.



1	(1) In general.—Section 1871(e), as added by
2	subsection (a), is further amended by adding at the
3	end the following new paragraph:
4	"(2)(A) If—
5	"(i) a provider of services or supplier follows the
6	written guidance (which may be transmitted elec-
7	tronically) provided by the Secretary or by a medi-
8	care contractor (as defined in section $1889(g)$ ) acting
9	within the scope of the contractor's contract authority,
10	with respect to the furnishing of items or services and
11	submission of a claim for benefits for such items or
12	services with respect to such provider or supplier;
13	"(ii) the Secretary determines that the provider
14	of services or supplier has accurately presented the
15	circumstances relating to such items, services, and
16	claim to the contractor in writing; and
17	"(iii) the guidance was in error;
18	the provider of services or supplier shall not be subject to
19	any sanction (including any penalty or requirement for re-
20	payment of any amount) if the provider of services or sup-
21	plier reasonably relied on such guidance.
22	"(B) Subparagraph (A) shall not be construed as pre-
23	venting the recoupment or repayment (without any addi-
24	tional penalty) relating to an overpayment insofar as the



1	overpayment was solely the result of a clerical or technical
2	operational error.".
3	(2) Effective date.—The amendment made by
4	paragraph (1) shall take effect on the date of the en-
5	actment of this Act but shall not apply to any sanc-
6	tion for which notice was provided on or before the
7	date of the enactment of this Act.
8	SEC. 904. REPORTS AND STUDIES RELATING TO REGU
9	LATORY REFORM.
10	(a) GAO STUDY ON ADVISORY OPINION AUTHORITY.—
11	(1) STUDY.—The Comptroller General of the
12	United States shall conduct a study to determine the
13	feasibility and appropriateness of establishing in the
14	Secretary authority to provide legally binding advi-
15	sory opinions on appropriate interpretation and ap-
16	plication of regulations to carry out the medicare pro-
17	gram under title XVIII of the Social Security Act
18	Such study shall examine the appropriate timeframe
19	for issuing such advisory opinions, as well as the need
20	for additional staff and funding to provide such opin-
21	ions.
22	(2) Report.—The Comptroller General shall
23	submit to Congress a report on the study conducted
24	under paragraph (1) by not later than one year after

 $the \ date \ of \ the \ enactment \ of \ this \ Act.$ 



1	(b) Report on Legal and Regulatory Inconsist-
2	Encies.—Section 1871 (42 U.S.C. 1395hh), as amended by
3	section 2(a), is amended by adding at the end the following
4	new subsection:
5	"(f)(1) Not later than 2 years after the date of the en-
6	actment of this subsection, and every 2 years thereafter, the
7	Secretary shall submit to Congress a report with respect
8	to the administration of this title and areas of inconsistency
9	or conflict among the various provisions under law and reg-
10	ulation.
11	"(2) In preparing a report under paragraph (1), the
12	Secretary shall collect—
13	"(A) information from individuals entitled to
14	benefits under part A or enrolled under part B, or
15	both, providers of services, and suppliers and from the
16	Medicare Beneficiary Ombudsman and the Medicare
17	Provider Ombudsman with respect to such areas of
18	inconsistency and conflict; and
19	"(B) information from medicare contractors that
20	tracks the nature of written and telephone inquiries.
21	"(3) A report under paragraph (1) shall include a de-
22	scription of efforts by the Secretary to reduce such inconsist-
23	ency or conflicts, and recommendations for legislation or
24	administrative action that the Secretary determines appro-

25 priate to further reduce such inconsistency or conflicts.".



## Subtitle B—Contracting Reform 1 SEC. 911. INCREASED FLEXIBILITY IN MEDICARE ADMINIS-3 TRATION. 4 (a) Consolidation and Flexibility in Medicare ADMINISTRATION.— 5 6 (1) In General.—Title XVIII is amended by in-7 serting after section 1874 the following new section: 8 "CONTRACTS WITH MEDICARE ADMINISTRATIVE 9 CONTRACTORS10 "Sec. 1874A. (a) AUTHORITY.— 11 "(1) Authority to enter into contracts.— 12 The Secretary may enter into contracts with any eli-13 gible entity to serve as a medicare administrative 14 contractor with respect to the performance of any or 15 all of the functions described in paragraph (4) or 16 parts of those functions (or, to the extent provided in 17 a contract, to secure performance thereof by other en-18 tities). 19 "(2) Eligibility of entities.—An entity is el-20 igible to enter into a contract with respect to the per-21 formance of a particular function described in para-22 graph (4) only if— "(A) the entity has demonstrated capability 23

to carry out such function;



1	"(B) the entity complies with such conflict
2	of interest standards as are generally applicable
3	to Federal acquisition and procurement;
4	"(C) the entity has sufficient assets to fi-
5	nancially support the performance of such func-
6	tion; and
7	"(D) the entity meets such other require-
8	ments as the Secretary may impose.
9	"(3) Medicare administrative contractor
10	DEFINED.—For purposes of this title and title XI—
11	"(A) In general.—The term 'medicare ad-
12	ministrative contractor' means an agency, orga-
13	nization, or other person with a contract under
14	this section.
15	"(B) Appropriate medicare administra-
16	TIVE CONTRACTOR.—With respect to the perform-
17	ance of a particular function in relation to an
18	individual entitled to benefits under part $A$ or
19	enrolled under part B, or both, a specific pro-
20	vider of services or supplier (or class of such pro-
21	viders of services or suppliers), the 'appropriate'
22	medicare administrative contractor is the medi-
23	care administrative contractor that has a con-
24	tract under this section with respect to the per-
25	formance of that function in relation to that in-



1	dividual, provider of services or supplier or class
2	of provider of services or supplier.
3	"(4) Functions described.—The functions re-
4	ferred to in paragraphs (1) and (2) are payment
5	functions, provider services functions, and functions
6	relating to services furnished to individuals entitled
7	to benefits under part A or enrolled under part B, or
8	both, as follows:
9	"(A) DETERMINATION OF PAYMENT
10	AMOUNTS.—Determining (subject to the provi-
11	sions of section 1878 and to such review by the
12	Secretary as may be provided for by the con-
13	tracts) the amount of the payments required pur-
14	suant to this title to be made to providers of
15	services, suppliers and individuals.
16	"(B) Making payments.—Making pay-
17	ments described in subparagraph (A) (including
18	receipt, disbursement, and accounting for funds
19	in making such payments).
20	"(C) Beneficiary education and assist-
21	ANCE.—Providing education and outreach to in-
22	dividuals entitled to benefits under part A or en-
23	rolled under part B, or both, and providing as-
24	sistance to those individuals with enerific issues

concerns or problems.



1	"(D) Provider consultative serv-
2	ICES.—Providing consultative services to institu-
3	tions, agencies, and other persons to enable them
4	to establish and maintain fiscal records nec-
5	essary for purposes of this title and otherwise to
6	qualify as providers of services or suppliers.
7	"(E) Communication with providers.—
8	Communicating to providers of services and sup-
9	pliers any information or instructions furnished
10	to the medicare administrative contractor by the
11	Secretary, and facilitating communication be-
12	tween such providers and suppliers and the Sec-
13	retary.
14	"(F) Provider education and technical
15	ASSISTANCE.—Performing the functions relating
16	to provider education, training, and technical
17	assistance.
18	"(G) Additional functions.—Performing
19	such other functions as are necessary to carry
20	out the purposes of this title.
21	"(5) Relationship to mip contracts.—
22	"(A) Nonduplication of duties.—In en-
23	tering into contracts under this section, the Sec-
24	retary shall assure that functions of medicare

administrative contractors in carrying out ac-



1	tivities under parts A and B do not duplicate
2	activities carried out under the Medicare Integ-
3	rity Program under section 1893. The previous
4	sentence shall not apply with respect to the ac-
5	tivity described in section 1893(b)(5) (relating to
6	prior authorization of certain items of durable
7	$medical\ equipment\ under\ section\ 1834(a)(15)).$
8	"(B) Construction.—An entity shall not
9	be treated as a medicare administrative con-
10	tractor merely by reason of having entered into
11	a contract with the Secretary under section
12	1893.
13	"(6) Application of federal acquisition
14	REGULATION.—Except to the extent inconsistent with
15	a specific requirement of this title, the Federal Acqui-
16	sition Regulation applies to contracts under this title.
17	"(b) Contracting Requirements.—
18	"(1) Use of competitive procedures.—
19	"(A) In general.—Except as provided in
20	laws with general applicability to Federal acqui-
21	sition and procurement or in subparagraph (B),
22	the Secretary shall use competitive procedures
23	when entering into contracts with medicare ad-

ministrative contractors under this section, tak-



1 ing into account performance quality as well as 2 price and other factors. "(B) Renewal of contracts.—The Sec-3 4 retary may renew a contract with a medicare 5 administrative contractor under this section 6 from term to term without regard to section 5 of 7 title 41, United States Code, or any other provi-8 sion of law requiring competition, if the medi-9 care administrative contractor has met or ex-10 ceeded the performance requirements applicable 11 with respect to the contract and contractor, ex-12 cept that the Secretary shall provide for the application of competitive procedures under such a 13 14 contract not less frequently than once every five 15 years. 16 "(C) Transfer of functions.—The Sec-17 retary may transfer functions among medicare 18 administrative contractors consistent with the 19 provisions of this paragraph. The Secretary shall 20 ensure that performance quality is considered in 21 such transfers. The Secretary shall provide pub-22 lic notice (whether in the Federal Register or 23 otherwise) of any such transfer (including a de-24 scription of the functions so transferred, a de-

scription of the providers of services and sup-



1	pliers affected by such transfer, and contact in-
2	formation for the contractors involved).
3	"(D) Incentives for quality.—The Sec-
4	retary shall provide incentives for medicare ad-
5	ministrative contractors to provide quality serv-
6	ice and to promote efficiency.
7	"(2) Compliance with requirements.—No
8	contract under this section shall be entered into with
9	any medicare administrative contractor unless the
10	Secretary finds that such medicare administrative
11	contractor will perform its obligations under the con-
12	tract efficiently and effectively and will meet such re-
13	quirements as to financial responsibility, legal au-
14	thority, quality of services provided, and other mat-
15	ters as the Secretary finds pertinent.
16	"(3) Performance requirements.—
17	"(A) Development of specific perform-
18	ANCE REQUIREMENTS.—In developing contract
19	performance requirements, the Secretary shall
20	develop performance requirements applicable to
21	functions described in subsection $(a)(4)$ .
22	"(B) Consultation.— In developing such
23	requirements, the Secretary may consult with
24	providers of services and suppliers, organizations

representing individuals entitled to benefits



1	under part A or enrolled under part B, or both,
2	and organizations and agencies performing func-
3	tions necessary to carry out the purposes of this
4	section with respect to such performance require-
5	ments.
6	"(C) Inclusion in contracts.—All con-
7	tractor performance requirements shall be set
8	forth in the contract between the Secretary and
9	the appropriate medicare administrative con-
10	tractor. Such performance requirements—
11	"(i) shall reflect the performance re-
12	quirements developed under subparagraph
13	(A), but may include additional perform-
14	$ance\ requirements;$
15	"(ii) shall be used for evaluating con-
16	tractor performance under the contract; and
17	"(iii) shall be consistent with the writ-
18	ten statement of work provided under the
19	contract.
20	"(4) Information requirements.—The Sec-
21	retary shall not enter into a contract with a medicare
22	administrative contractor under this section unless
23	the contractor agrees—
24	"(A) to furnish to the Secretary such timely
25	information and reports as the Secretary may



1	find necessary in performing his functions under
2	this title; and
3	"(B) to maintain such records and afford
4	such access thereto as the Secretary finds nec-
5	essary to assure the correctness and verification
6	of the information and reports under subpara-
7	graph (A) and otherwise to carry out the pur-
8	poses of this title.
9	"(5) Surety Bond.—A contract with a medi-
10	care administrative contractor under this section may
11	require the medicare administrative contractor, and
12	any of its officers or employees certifying payments or
13	disbursing funds pursuant to the contract, or other-
14	wise participating in carrying out the contract, to
15	give surety bond to the United States in such amount
16	as the Secretary may deem appropriate.
17	"(c) Terms and Conditions.—
18	"(1) In general.—A contract with any medi-
19	care administrative contractor under this section may
20	contain such terms and conditions as the Secretary
21	finds necessary or appropriate and may provide for
22	advances of funds to the medicare administrative con-
23	tractor for the making of payments by it under sub-



section (a)(4)(B).

1	"(2) Prohibition on mandates for certain
2	DATA COLLECTION.—The Secretary may not require,
3	as a condition of entering into, or renewing, a con-
4	tract under this section, that the medicare adminis-
5	trative contractor match data obtained other than in
6	its activities under this title with data used in the ad-
7	ministration of this title for purposes of identifying
8	situations in which the provisions of section 1862(b)
9	may apply.
10	"(d) Limitation on Liability of Medicare Admin-
11	ISTRATIVE CONTRACTORS AND CERTAIN OFFICERS.—
12	"(1) Certifying officer.—No individual des-
13	ignated pursuant to a contract under this section as
14	a certifying officer shall, in the absence of the reckless
15	disregard of the individual's obligations or the intent
16	by that individual to defraud the United States, be
17	liable with respect to any payments certified by the
18	individual under this section.
19	"(2) Disbursing officer.—No disbursing offi-
20	cer shall, in the absence of the reckless disregard of the
21	officer's obligations or the intent by that officer to de-
22	fraud the United States, be liable with respect to any
23	payment by such officer under this section if it was
24	based upon an authorization (which meets the appli-

cable requirements for such internal controls estab-



1	lished by the Comptroller General) of a certifying offi-
2	cer designated as provided in paragraph (1) of this
3	subsection.
4	"(3) Liability of medicare administrative
5	CONTRACTOR.—
6	"(A) In general.—No medicare administrative
7	contractor shall be liable to the United States for a
8	payment by a certifying or disbursing officer unless,
9	in connection with such payment, the medicare ad-
10	ministrative contractor acted with reckless disregard
11	of its obligations under its medicare administrative
12	contract or with intent to defraud the United States.
13	"(B) Relationship to false claims act.—
14	Nothing in this subsection shall be construed to limit
15	liability for conduct that would constitute a violation
16	of sections 3729 through 3731 of title 31, United
17	States Code (commonly known as the False Claims
18	Act').
19	"(4) Indemnification by secretary.—
20	"(A) In general.—Subject to subpara-
21	graphs (B) and (D), in the case of a medicare
22	administrative contractor (or a person who is a
23	director, officer, or employee of such a contractor
24	or who is engaged by the contractor to partici-

pate directly in the claims administration proc-



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1	ess) who is made a party to any judicial or ad-
2	ministrative proceeding arising from or relating
3	directly to the claims administration process
4	under this title, the Secretary may, to the extent
5	the Secretary determines to be appropriate and
6	as specified in the contract with the contractor,
7	indemnify the contractor and such persons.
8	"(B) Conditions.—The Secretary may not
9	provide indemnification under subparagraph (A)
10	insofar as the liability for such costs arises di-
11	rectly from conduct that is determined by the ju-
12	dicial proceeding or by the Secretary to be crimi-
13	nal in nature, fraudulent, or grossly negligent. If
14	indemnification is provided by the Secretary
15	with respect to a contractor before a determina-
16	tion that such costs arose directly from such con-
17	duct, the contractor shall reimburse the Secretary
18	for costs of indemnification.
19	"(C) Scope of indemnification.—Indem-
20	nification by the Secretary under subparagraph
21	(A) may include payment of judgments, settle-
22	ments (subject to subparagraph (D)), awards,
23	and costs (including reasonable legal expenses).
24	"(D) Written approval for settle-



1	in subparagraph (A) may not propose to nego-
2	tiate a settlement or compromise of a proceeding
3	described in such subparagraph without the
4	prior written approval of the Secretary to nego-
5	tiate such settlement or compromise. Any indem-
6	nification under subparagraph (A) with respect
7	to amounts paid under a settlement or com-
8	promise of a proceeding described in such sub-
9	paragraph are conditioned upon prior written
10	approval by the Secretary of the final settlement
11	or compromise.
12	"(E) Construction.—Nothing in this
13	paragraph shall be construed—
14	"(i) to change any common law immu-
15	nity that may be available to a medicare
16	administrative contractor or person de-
17	scribed in subparagraph (A); or
18	"(ii) to permit the payment of costs
19	not otherwise allowable, reasonable, or allo-
20	cable under the Federal Acquisition Regula-
21	tions.".
22	(2) Consideration of incorporation of cur-
23	RENT LAW STANDARDS.—In developing contract per-
24	formance requirements under section 1874A(b) of the
25	Social Security Act, as inserted by paragraph (1), the



1	Secretary shall consider inclusion of the performance
2	standards described in sections 1816(f)(2) of such Act
3	(relating to timely processing of reconsiderations and
4	applications for exemptions) and section
5	1842(b)(2)(B) of such Act (relating to timely review
6	of determinations and fair hearing requests), as such
7	sections were in effect before the date of the enactment
8	$of\ this\ Act.$
9	(b) Conforming Amendments to Section 1816 (Re-
10	LATING TO FISCAL INTERMEDIARIES).—Section 1816 (42
11	U.S.C. 1395h) is amended as follows:
12	(1) The heading is amended to read as follows:
13	"PROVISIONS RELATING TO THE ADMINISTRATION OF PART
14	A".
15	(2) Subsection (a) is amended to read as follows:
16	"(a) The administration of this part shall be conducted
17	through contracts with medicare administrative contractors
18	under section 1874A.".
19	(3) Subsection (b) is repealed.
20	(4) Subsection (c) is amended—
21	(A) by striking paragraph (1); and
22	(B) in each of paragraphs $(2)(A)$ and
23	(3)(A), by striking "agreement under this sec-
24	tion" and inserting "contract under section
25	1874A that provides for making payments under



this part".

1	(5) Subsections (d) through (i) are repealed.
2	(6) Subsections (j) and (k) are each amended—
3	(A) by striking "An agreement with an
4	agency or organization under this section" and
5	inserting "A contract with a medicare adminis-
6	trative contractor under section 1874A with re-
7	spect to the administration of this part'; and
8	(B) by striking "such agency or organiza-
9	tion" and inserting "such medicare administra-
10	tive contractor" each place it appears.
11	(7) Subsection (l) is repealed.
12	(c) Conforming Amendments to Section 1842 (Re-
13	LATING TO CARRIERS).—Section 1842 (42 U.S.C. 1395u)
14	is amended as follows:
15	(1) The heading is amended to read as follows:
16	"PROVISIONS RELATING TO THE ADMINISTRATION OF PART
17	$B^{\prime\prime}.$
18	(2) Subsection (a) is amended to read as follows:
19	"(a) The administration of this part shall be conducted
20	$through\ contracts\ with\ medicare\ administrative\ contractors$
21	under section 1874A.".
22	(3) Subsection (b) is amended—
23	(A) by striking paragraph (1);
24	(B) in paragraph (2)—
25	(i) by striking subparagraphs (A) and
26	(B);



1	(ii) in subparagraph (C), by striking
2	"carriers" and inserting "medicare admin-
3	istrative contractors"; and
4	(iii) by striking subparagraphs (D)
5	and $(E)$ ;
6	(C) in paragraph (3)—
7	(i) in the matter before subparagraph
8	(A), by striking "Each such contract shall
9	provide that the carrier" and inserting
10	"The Secretary";
11	(ii) by striking "will" the first place it
12	appears in each of subparagraphs (A), (B),
13	(F), (G), (H), and (L) and inserting
14	"shall";
15	(iii) in subparagraph (B), in the mat-
16	ter before clause (i), by striking "to the pol-
17	icyholders and subscribers of the carrier"
18	and inserting "to the policyholders and sub-
19	scribers of the medicare administrative con-
20	tractor";
21	(iv) by striking subparagraphs (C),
22	(D), and $(E)$ ;
23	(v) in subparagraph (H)—
24	(I) by striking "if it makes deter-
25	minations or payments with respect to



1	physicians' services," in the matter
2	preceding clause (i); and
3	(II) by striking "carrier" and in-
4	serting "medicare administrative con-
5	tractor" in clause (i);
6	(vi) by striking subparagraph (I);
7	(vii) in subparagraph (L), by striking
8	the semicolon and inserting a period;
9	(viii) in the first sentence, after sub-
10	paragraph (L), by striking "and shall con-
11	tain" and all that follows through the pe-
12	riod; and
13	(ix) in the seventh sentence, by insert-
14	ing "medicare administrative contractor,"
15	after "carrier,"; and
16	(D) by striking paragraph (5);
17	(E) in paragraph $(6)(D)(iv)$ , by striking
18	"carrier" and inserting "medicare administra-
19	tive contractor"; and
20	(F) in paragraph (7), by striking "the car-
21	rier" and inserting "the Secretary" each place it
22	appears.
23	(4) Subsection (c) is amended—
24	(A) by striking paragraph (1):



1	(B) in paragraph $(2)(A)$ , by striking "con-
2	tract under this section which provides for the
3	disbursement of funds, as described in subsection
4	(a)(1)(B)," and inserting "contract under section
5	1874A that provides for making payments under
6	this part";
7	(C) in paragraph (3)(A), by striking "sub-
8	section $(a)(1)(B)$ " and inserting "section
9	1874A(a)(3)(B)";
10	(D) in paragraph (4), in the matter pre-
11	ceding subparagraph (A), by striking "carrier"
12	and inserting "medicare administrative con-
13	tractor"; and
14	(E) by striking paragraphs (5) and (6).
15	(5) Subsections (d), (e), and (f) are repealed.
16	(6) Subsection (g) is amended by striking "car-
17	rier or carriers" and inserting "medicare administra-
18	tive contractor or contractors".
19	(7) Subsection (h) is amended—
20	(A) in paragraph (2)—
21	(i) by striking "Each carrier having
22	an agreement with the Secretary under sub-
23	section (a)" and inserting "The Secretary";
24	and



1	(ii) by striking "Each such carrier"
2	and inserting "The Secretary";
3	(B) in paragraph $(3)(A)$ —
4	(i) by striking "a carrier having an
5	agreement with the Secretary under sub-
6	section (a)" and inserting "medicare ad-
7	ministrative contractor having a contract
8	under section 1874A that provides for mak-
9	ing payments under this part"; and
10	(ii) by striking "such carrier" and in-
11	serting "such contractor";
12	(C) in paragraph $(3)(B)$ —
13	(i) by striking "a carrier" and insert-
14	ing "a medicare administrative contractor"
15	each place it appears; and
16	(ii) by striking "the carrier" and in-
17	serting "the contractor" each place it ap-
18	pears; and
19	(D) in paragraphs $(5)(A)$ and $(5)(B)(iii)$ ,
20	by striking "carriers" and inserting "medicare
21	administrative contractors" each place it ap-
22	pears.
23	(8) Subsection (1) is amended—



1	(A) in paragraph (1)(A)(iii), by striking
2	"carrier" and inserting "medicare administra-
3	tive contractor"; and
4	(B) in paragraph (2), by striking "carrier"
5	and inserting "medicare administrative con-
6	tractor".
7	(9) Subsection $(p)(3)(A)$ is amended by striking
8	"carrier" and inserting "medicare administrative
9	contractor".
10	(10) Subsection $(q)(1)(A)$ is amended by striking
11	"carrier".
12	(d) Effective Date; Transition Rule.—
13	(1) Effective date.—
14	(A) In general.—Except as otherwise pro-
15	vided in this subsection, the amendments made
16	by this section shall take effect on October 1,
17	2005, and the Secretary is authorized to take
18	such steps before such date as may be necessary
19	to implement such amendments on a timely
20	basis.
21	(B) Construction for current con-
22	TRACTS.—Such amendments shall not apply to
23	contracts in effect before the date specified under
24	subparagraph (A) that continue to retain the

terms and conditions in effect on such date (ex-



1	cept as otherwise provided under this Act, other
2	than under this section) until such date as the
3	contract is let out for competitive bidding under
4	such amendments.
5	(C) Deadline for competitive bid-
6	DING.—The Secretary shall provide for the let-
7	ting by competitive bidding of all contracts for
8	functions of medicare administrative contractors
9	for annual contract periods that begin on or
10	after October 1, 2010.
11	(D) Waiver of provider nomination
12	PROVISIONS DURING TRANSITION.—During the
13	period beginning on the date of the enactment of
14	this Act and before the date specified under sub-
15	paragraph (A), the Secretary may enter into
16	new agreements under section 1816 of the Social
17	Security Act (42 U.S.C. 1395h) without regard
18	to any of the provider nomination provisions of
19	such section.
20	(2) General transition rules.—The Sec-
21	retary shall take such steps, consistent with para-
22	graph (1)(B) and (1)(C), as are necessary to provide
23	for an appropriate transition from contracts under

section 1816 and section 1842 of the Social Security



1	Act (42 U.S.C. 1395h, 1395u) to contracts under sec-
2	tion 1874A, as added by subsection (a)(1).
3	(3) Authorizing continuation of mip func-
4	TIONS UNDER CURRENT CONTRACTS AND AGREE-
5	MENTS AND UNDER ROLLOVER CONTRACTS.—The pro-
6	visions contained in the exception in section
7	1893(d)(2) of the Social Security Act (42 U.S.C.
8	1395ddd(d)(2)) shall continue to apply notwith-
9	standing the amendments made by this section, and
10	any reference in such provisions to an agreement or
11	contract shall be deemed to include a contract under
12	section 1874A of such Act, as inserted by subsection
13	(a)(1), that continues the activities referred to in such
14	provisions.
15	(e) References.—On and after the effective date pro-
16	vided under subsection (d)(1), any reference to a fiscal
17	intermediary or carrier under title XI or XVIII of the So-
18	cial Security Act (or any regulation, manual instruction,
19	interpretative rule, statement of policy, or guideline issued
20	to carry out such titles) shall be deemed a reference to a
21	medicare administrative contractor (as provided under sec-
22	tion 1874A of the Social Security Act).
23	(f) Reports on Implementation.—
24	(1) Plan for implementation.—By not later
25	than October 1, 2004, the Secretary shall submit a re-



1	port to Congress and the Comptroller General of the
2	United States that describes the plan for implementa-
3	tion of the amendments made by this section. The
4	Comptroller General shall conduct an evaluation of
5	such plan and shall submit to Congress, not later
6	than 6 months after the date the report is received, a
7	report on such evaluation and shall include in such
8	report such recommendations as the Comptroller Gen-
9	eral deems appropriate.
10	(2) Status of implementation.—The Sec-
11	retary shall submit a report to Congress not later
12	than October 1, 2008, that describes the status of im-
13	plementation of such amendments and that includes
14	a description of the following:
15	(A) The number of contracts that have been
16	competitively bid as of such date.
17	(B) The distribution of functions among
18	contracts and contractors.
19	(C) A timeline for complete transition to
20	full competition.
21	(D) A detailed description of how the Sec-
22	retary has modified oversight and management
23	of medicare contractors to adapt to full competi-



tion.

1	SEC. 912. REQUIREMENTS FOR INFORMATION SECURITY
2	FOR MEDICARE ADMINISTRATIVE CONTRAC-
3	TORS.
4	(a) In General.—Section 1874A, as added by section
5	911(a)(1), is amended by adding at the end the following
6	new subsection:
7	"(e) Requirements for Information Security.—
8	"(1) Development of information security
9	PROGRAM.—A medicare administrative contractor
10	that performs the functions referred to in subpara-
11	graphs (A) and (B) of subsection (a)(4) (relating to
12	determining and making payments) shall implement
13	a contractor-wide information security program to
14	provide information security for the operation and
15	assets of the contractor with respect to such functions
16	under this title. An information security program
17	under this paragraph shall meet the requirements for
18	information security programs imposed on Federal
19	agencies under paragraphs (1) through (8) of section
20	3544(b) of title 44, United States Code (other than the
21	requirements under paragraphs $(2)(D)(i)$ , $(5)(A)$ , and
22	(5)(B) of such section).
23	"(2) Independent audits.—
24	"(A) PERFORMANCE OF ANNUAL EVALUA-
25	TIONS.—Each year a medicare administrative
26	contractor that performs the functions referred to



1	in subparagraphs (A) and (B) of subsection
2	(a)(4) (relating to determining and making pay-
3	ments) shall undergo an evaluation of the infor-
4	mation security of the contractor with respect to
5	such functions under this title. The evaluation
6	shall—
7	"(i) be performed by an entity that
8	meets such requirements for independence as
9	the Inspector General of the Department of
10	Health and Human Services may establish;
11	and
12	"(ii) test the effectiveness of informa-
13	tion security control techniques of an ap-
14	propriate subset of the contractor's informa-
15	tion systems (as defined in section 3502(8)
16	of title 44, United States Code) relating to
17	such functions under this title and an as-
18	sessment of compliance with the require-
19	ments of this subsection and related infor-
20	mation security policies, procedures, stand-
21	ards and guidelines, including policies and
22	procedures as may be prescribed by the Di-
23	rector of the Office of Management and

Budget and applicable information security



1	standards promulgated under section 11331
2	of title 40, United States Code.
3	"(B) Deadline for initial evalua-
4	TION.—
5	"(i) New contractors.—In the case
6	of a medicare administrative contractor
7	covered by this subsection that has not pre-
8	viously performed the functions referred to
9	in subparagraphs (A) and (B) of subsection
10	(a)(4) (relating to determining and making
11	payments) as a fiscal intermediary or car-
12	rier under section 1816 or 1842, the first
13	independent evaluation conducted pursuant
14	subparagraph (A) shall be completed prior
15	to commencing such functions.
16	"(ii) Other contractors.—In the
17	case of a medicare administrative con-
18	tractor covered by this subsection that is not
19	described in clause (i), the first independent
20	evaluation conducted pursuant subpara-
21	graph (A) shall be completed within 1 year
22	after the date the contractor commences
23	functions referred to in clause (i) under this
24	section.
25	"(C) Reports on evaluations.—



1	"(i) To the department of health
2	AND HUMAN SERVICES.—The results of
3	independent evaluations under subpara-
4	graph (A) shall be submitted promptly to
5	the Inspector General of the Department of
6	Health and Human Services and to the Sec-
7	retary.
8	"(ii) To congress.—The Inspector
9	General of Department of Health and
10	Human Services shall submit to Congress
11	annual reports on the results of such eval-
12	uations, including assessments of the scope
13	and sufficiency of such evaluations.
14	"(iii) AGENCY REPORTING.—The Sec-
15	retary shall address the results of such eval-
16	uations in reports required under section
17	3544(c) of title 44, United States Code.".
18	(b) Application of Requirements to Fiscal
19	Intermediaries and Carriers.—
20	(1) In general.—The provisions of section
21	1874A(e)(2) of the Social Security Act (other than
22	subparagraph (B)), as added by subsection (a), shall
23	apply to each fiscal intermediary under section 1816
24	of the Social Security Act (42 U.S.C. 1395h) and
25	each carrier under section 1842 of such Act (42



1	U.S.C. 1395u) in the same manner as they apply to
2	medicare administrative contractors under such pro-
3	visions.
4	(2) Deadline for initial evaluation.—In the
5	case of such a fiscal intermediary or carrier with an
6	agreement or contract under such respective section in
7	effect as of the date of the enactment of this Act, the
8	first evaluation under section 1874A(e)(2)(A) of the
9	Social Security Act (as added by subsection (a)), pur-
10	suant to paragraph (1), shall be completed (and a re-
11	port on the evaluation submitted to the Secretary) by
12	not later than 1 year after such date.
13	Subtitle C—Education and
14	Outreach
	Outreach SEC. 921. PROVIDER EDUCATION AND TECHNICAL ASSIST-
14	
14 15	SEC. 921. PROVIDER EDUCATION AND TECHNICAL ASSIST
<ul><li>14</li><li>15</li><li>16</li></ul>	SEC. 921. PROVIDER EDUCATION AND TECHNICAL ASSISTANCE.
<ul><li>14</li><li>15</li><li>16</li><li>17</li></ul>	SEC. 921. PROVIDER EDUCATION AND TECHNICAL ASSISTANCE.  (a) COORDINATION OF EDUCATION FUNDING.—
14 15 16 17 18	SEC. 921. PROVIDER EDUCATION AND TECHNICAL ASSISTANCE.  (a) COORDINATION OF EDUCATION FUNDING.—  (1) IN GENERAL.—Title XVIII is amended by in-
14 15 16 17 18 19	SEC. 921. PROVIDER EDUCATION AND TECHNICAL ASSISTANCE.  (a) COORDINATION OF EDUCATION FUNDING.—  (1) IN GENERAL.—Title XVIII is amended by inserting after section 1888 the following new section:
<ul><li>14</li><li>15</li><li>16</li><li>17</li><li>18</li><li>19</li><li>20</li></ul>	SEC. 921. PROVIDER EDUCATION AND TECHNICAL ASSISTANCE.  (a) COORDINATION OF EDUCATION FUNDING.—  (1) IN GENERAL.—Title XVIII is amended by inserting after section 1888 the following new section:  "PROVIDER EDUCATION AND TECHNICAL ASSISTANCE"
14 15 16 17 18 19 20 21	SEC. 921. PROVIDER EDUCATION AND TECHNICAL ASSISTANCE.  (a) COORDINATION OF EDUCATION FUNDING.—  (1) IN GENERAL.—Title XVIII is amended by inserting after section 1888 the following new section:  "PROVIDER EDUCATION AND TECHNICAL ASSISTANCE"  "SEC. 1889. (a) COORDINATION OF EDUCATION FUND-
14 15 16 17 18 19 20 21 22	SEC. 921. PROVIDER EDUCATION AND TECHNICAL ASSISTANCE.  (a) COORDINATION OF EDUCATION FUNDING.—  (1) IN GENERAL.—Title XVIII is amended by inserting after section 1888 the following new section:  "PROVIDER EDUCATION AND TECHNICAL ASSISTANCE"  "SEC. 1889. (a) COORDINATION OF EDUCATION FUNDING.—The Secretary shall coordinate the educational activities.
14 15 16 17 18 19 20 21 22 23	ANCE.  (a) Coordination of Education Funding.—  (1) In General.—Title XVIII is amended by inserting after section 1888 the following new section:  "Provider Education and Technical Assistance  "Sec. 1889. (a) Coordination of Education Funding.—The Secretary shall coordinate the educational activities provided through medicare contractors (as defined in



1	(2) Effective date.—The amendment made by
2	paragraph (1) shall take effect on the date of the en-
3	actment of this Act.
4	(3) Report.—Not later than October 1, 2004,
5	the Secretary shall submit to Congress a report that
6	includes a description and evaluation of the steps
7	taken to coordinate the funding of provider education
8	under section 1889(a) of the Social Security Act, as
9	added by paragraph (1).
10	(b) Incentives To Improve Contractor Perform-
11	ANCE.—
12	(1) In General.—Section 1874A, as added by
13	section 911(a)(1) and as amended by section 912(a),
14	is amended by adding at the end the following new
15	subsection:
16	"(f) Incentives To Improve Contractor Perform-
17	ANCE IN PROVIDER EDUCATION AND OUTREACH.—The Sec-
18	retary shall use specific claims payment error rates or simi-
19	lar methodology of medicare administrative contractors in
20	the processing or reviewing of medicare claims in order to
21	give such contractors an incentive to implement effective
22	education and outreach programs for providers of services
23	and suppliers.".
24	(2) Application to fiscal intermediaries
25	AND CARRIERS.—The provisions of section 1874A(f) of



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1	the Social Security Act, as added by paragraph (1),
2	shall apply to each fiscal intermediary under section
3	1816 of the Social Security Act (42 U.S.C. 1395h)
4	and each carrier under section 1842 of such Act (42
5	U.S.C. 1395u) in the same manner as they apply to
5	medicare administrative contractors under such pro-
7	visions.

- (3) GAO REPORT ON ADEQUACY OF METHOD-OLOGY.—Not later than October 1, 2004, the Comptroller General of the United States shall submit to Congress and to the Secretary a report on the adequacy of the methodology under section 1874A(f) of the Social Security Act, as added by paragraph (1), and shall include in the report such recommendations as the Comptroller General determines appropriate with respect to the methodology.
- (4) Report on use of methodology in assessing contractor performance in implementing effective education and outreach programs, including whether to use such methodology as a basis for performance bonuses. The report shall include an analysis of the



1	sources of identified errors and potential changes in
2	systems of contractors and rules of the Secretary that
3	could reduce claims error rates.
4	(c) Provision of Access to and Prompt Re-
5	SPONSES FROM MEDICARE ADMINISTRATIVE CONTRAC-
6	TORS.—
7	(1) In General.—Section 1874A, as added by
8	section 911(a)(1) and as amended by section 912(a)
9	and subsection (b), is further amended by adding at
10	the end the following new subsection:
11	"(g) Communications with Beneficiaries, Pro-
12	VIDERS OF SERVICES AND SUPPLIERS.—
13	"(1) Communication strategy.—The Secretary
14	shall develop a strategy for communications with in-
15	dividuals entitled to benefits under part A or enrolled
16	under part B, or both, and with providers of services
17	and suppliers under this title.
18	"(2) Response to written inquiries.—Each
19	medicare administrative contractor shall, for those
20	providers of services and suppliers which submit
21	claims to the contractor for claims processing and for
22	those individuals entitled to benefits under part $A$ or
23	enrolled under part B, or both, with respect to whom
24	claims are submitted for claims processing, provide

general written responses (which may be through elec-



1	tronic transmission) in a clear, concise, and accurate
2	manner to inquiries of providers of services, suppliers
3	and individuals entitled to benefits under part $A$ or
4	enrolled under part B, or both, concerning the pro-
5	grams under this title within 45 business days of the
6	date of receipt of such inquiries.
7	"(3) Response to toll-free lines.—The Sec-
8	retary shall ensure that each medicare administrative
9	contractor shall provide, for those providers of services
10	and suppliers which submit claims to the contractor
11	for claims processing and for those individuals enti-
12	tled to benefits under part A or enrolled under part
13	B, or both, with respect to whom claims are submitted
14	for claims processing, a toll-free telephone number at
15	which such individuals, providers of services and sup-
16	pliers may obtain information regarding billing, cod-
17	ing, claims, coverage, and other appropriate informa-
18	tion under this title.
19	"(4) Monitoring of contractor re-
20	SPONSES.—
21	"(A) In General.—Each medicare admin-
22	istrative contractor shall, consistent with stand-
23	ards developed by the Secretary under subpara-
24	graph(B)—



1	"(i) maintain a system for identifying
2	who provides the information referred to in
3	paragraphs (2) and (3); and
4	"(ii) monitor the accuracy, consist-
5	ency, and timeliness of the information so
6	provided.
7	"(B) Development of standards.—
8	"(i) In general.—The Secretary shall
9	establish and make public standards to
10	monitor the accuracy, consistency, and
11	timeliness of the information provided in
12	response to written and telephone inquiries
13	under this subsection. Such standards shall
14	be consistent with the performance require-
15	$ments\ established\ under\ subsection\ (b)(3).$
16	"(ii) Evaluation.—In conducting
17	evaluations of individual medicare admin-
18	istrative contractors, the Secretary shall
19	take into account the results of the moni-
20	toring conducted under subparagraph (A)
21	taking into account as performance require-
22	ments the standards established under
23	clause (i). The Secretary shall, in consulta-
24	tion with organizations representing pro-

viders of services, suppliers, and individuals



1	entitled to benefits under part A or enrolled
2	under part B, or both, establish standards
3	relating to the accuracy, consistency, and
4	timeliness of the information so provided.
5	"(C) Direct monitoring.—Nothing in this
6	paragraph shall be construed as preventing the
7	Secretary from directly monitoring the accuracy,
8	consistency, and timeliness of the information so
9	provided.".
10	(2) Effective date.—The amendment made by
11	paragraph (1) shall take effect October 1, 2004.
12	(3) Application to fiscal intermediaries
13	AND CARRIERS.—The provisions of section $1874A(g)$
14	of the Social Security Act, as added by paragraph
15	(1), shall apply to each fiscal intermediary under sec-
16	tion 1816 of the Social Security Act (42 U.S.C.
17	1395h) and each carrier under section 1842 of such
18	Act (42 U.S.C. 1395u) in the same manner as they
19	apply to medicare administrative contractors under
20	such provisions.
21	(d) Improved Provider Education and Train-
22	ING.—
23	(1) In general.—Section 1889, as added by
24	subsection (a), is amended by adding at the end the
25	following new subsections:



1	"(b) Enhanced Education and Training.—
2	"(1) Additional resources.—There are au-
3	thorized to be appropriated to the Secretary (in ap-
4	propriate part from the Federal Hospital Insurance
5	Trust Fund and the Federal Supplementary Medical
6	Insurance Trust Fund) \$25,000,000 for each of fiscal
7	years 2005 and 2006 and such sums as may be nec-
8	essary for succeeding fiscal years.
9	"(2) USE.—The funds made available under
10	paragraph (1) shall be used to increase the conduct by
11	medicare contractors of education and training of
12	providers of services and suppliers regarding billing,
13	coding, and other appropriate items and may also be
14	used to improve the accuracy, consistency, and timeli-
15	ness of contractor responses.
16	"(c) Tailoring Education and Training Activi-
17	ties for Small Providers or Suppliers.—
18	"(1) In general.—Insofar as a medicare con-
19	tractor conducts education and training activities, it
20	shall tailor such activities to meet the special needs
21	of small providers of services or suppliers (as defined
22	in paragraph (2)).
23	"(2) Small provider of services or sup-
24	PLIER.—In this subsection, the term 'small provider
25	of services or supplier' means—



1	"(A) a provider of services with fewer than
2	25 full-time-equivalent employees; or
3	"(B) a supplier with fewer than 10 full-
4	time-equivalent employees.".
5	(2) Effective date.—The amendment made by
6	paragraph (1) shall take effect on October 1, 2004.
7	(e) Requirement To Maintain Internet Sites.—
8	(1) In general.—Section 1889, as added by
9	subsection (a) and as amended by subsection (d), is
10	further amended by adding at the end the following
11	new subsection:
12	"(d) Internet Sites; FAQs.—The Secretary, and
13	each medicare contractor insofar as it provides services (in-
14	cluding claims processing) for providers of services or sup-
15	pliers, shall maintain an Internet site which—
16	"(1) provides answers in an easily accessible for-
17	mat to frequently asked questions, and
18	"(2) includes other published materials of the
19	contractor,
20	that relate to providers of services and suppliers under the
21	programs under this title (and title XI insofar as it relates
22	to such programs).".
23	(2) Effective date.—The amendment made by
24	paragraph (1) shall take effect on October 1, 2004.
25	(f) Additional Provider Education Provisions.—



1	(1) In general.—Section 1889, as added by
2	subsection (a) and as amended by subsections (d) and
3	(e), is further amended by adding at the end the fol-
4	lowing new subsections:
5	"(e) Encouragement of Participation in Edu-
6	CATION PROGRAM ACTIVITIES.—A medicare contractor
7	may not use a record of attendance at (or failure to attend)
8	educational activities or other information gathered during
9	an educational program conducted under this section or
10	otherwise by the Secretary to select or track providers of
11	services or suppliers for the purpose of conducting any type
12	of audit or prepayment review.
13	"(f) Construction.—Nothing in this section or sec-
14	tion 1893(g) shall be construed as providing for disclosure
15	by a medicare contractor of information that would com-
16	promise pending law enforcement activities or reveal find-
17	ings of law enforcement-related audits.
18	"(g) Definitions.—For purposes of this section, the
19	term 'medicare contractor' includes the following:
20	"(1) A medicare administrative contractor with
21	a contract under section 1874A, including a fiscal
22	intermediary with a contract under section 1816 and
23	a carrier with a contract under section 1842.
24	"(2) An eligible entity with a contract under sec-
25	tion 1893.



1	Such term does not include, with respect to activities of a
2	specific provider of services or supplier an entity that has
3	no authority under this title or title IX with respect to such
4	activities and such provider of services or supplier.".
5	(2) Effective date.—The amendment made by
6	paragraph (1) shall take effect on the date of the en-
7	actment of this Act.
8	SEC. 922. SMALL PROVIDER TECHNICAL ASSISTANCE DEM-
9	ONSTRATION PROGRAM.
10	(a) Establishment.—
11	(1) In General.—The Secretary shall establish
12	a demonstration program (in this section referred to
13	as the "demonstration program") under which tech-
14	nical assistance described in paragraph (2) is made
15	available, upon request and on a voluntary basis, to
16	small providers of services or suppliers in order to
17	improve compliance with the applicable requirements
18	of the programs under medicare program under title
19	XVIII of the Social Security Act (including provi-
20	sions of title XI of such Act insofar as they relate to
21	such title and are not administered by the Office of
22	the Inspector General of the Department of Health
23	and Human Services).
24	(2) Forms of technical assistance.—The

technical assistance described in this paragraph is—



1	(A) evaluation and recommendations re-
2	garding billing and related systems; and
3	(B) information and assistance regarding
4	policies and procedures under the medicare pro-
5	gram, including coding and reimbursement.
6	(3) Small providers of services or sup-
7	PLIERS.—In this section, the term "small providers of
8	services or suppliers" means—
9	(A) a provider of services with fewer than
10	25 full-time-equivalent employees; or
11	(B) a supplier with fewer than 10 full-time-
12	equivalent employees.
13	(b) Qualification of Contractors.—In conducting
14	the demonstration program, the Secretary shall enter into
15	contracts with qualified organizations (such as peer review
16	organizations or entities described in section $1889(g)(2)$ of
17	the Social Security Act, as inserted by section 5(f)(1)) with
18	appropriate expertise with billing systems of the full range
19	of providers of services and suppliers to provide the tech-
20	nical assistance. In awarding such contracts, the Secretary
21	shall consider any prior investigations of the entity's work
22	by the Inspector General of Department of Health and
23	Human Services or the Comptroller General of the United
24	States.



1	(c) Description of Technical Assistance.—The
2	technical assistance provided under the demonstration pro-
3	gram shall include a direct and in-person examination of
4	billing systems and internal controls of small providers of
5	services or suppliers to determine program compliance and
6	to suggest more efficient or effective means of achieving such
7	compliance.
8	(d) Avoidance of Recovery Actions for Prob-
9	LEMS IDENTIFIED AS CORRECTED.—The Secretary shall
10	provide that, absent evidence of fraud and notwithstanding
11	any other provision of law, any errors found in a compli-
12	ance review for a small provider of services or supplier that
13	participates in the demonstration program shall not be sub-
14	ject to recovery action if the technical assistance personnel
15	under the program determine that—
16	(1) the problem that is the subject of the compli-
17	ance review has been corrected to their satisfaction
18	within 30 days of the date of the visit by such per-
19	sonnel to the small provider of services or supplier;
20	and
21	(2) such problem remains corrected for such pe-
22	riod as is appropriate.
23	The previous sentence applies only to claims filed as part
24	of the demonstration program and lasts only for the dura-



- 1 tion of such program and only as long as the small provider
- 2 of services or supplier is a participant in such program.
- 3 (e) GAO EVALUATION.—Not later than 2 years after
- 4 the date of the date the demonstration program is first im-
- 5 plemented, the Comptroller General, in consultation with
- 6 the Inspector General of the Department of Health and
- 7 Human Services, shall conduct an evaluation of the dem-
- 8 onstration program. The evaluation shall include a deter-
- 9 mination of whether claims error rates are reduced for
- 10 small providers of services or suppliers who participated
- 11 in the program and the extent of improper payments made
- 12 as a result of the demonstration program. The Comptroller
- 13 General shall submit a report to the Secretary and the Con-
- 14 gress on such evaluation and shall include in such report
- 15 recommendations regarding the continuation or extension
- 16 of the demonstration program.
- 17 (f) Financial Participation by Providers.—The
- 18 provision of technical assistance to a small provider of serv-
- 19 ices or supplier under the demonstration program is condi-
- 20 tioned upon the small provider of services or supplier pay-
- 21 ing an amount estimated (and disclosed in advance of a
- 22 provider's or supplier's participation in the program) to
- 23 be equal to 25 percent of the cost of the technical assistance.
- 24 (g) Authorization of Appropriations.—There are
- 25 authorized to be appropriated to the Secretary (in appro-



1	priate part from the Federal Hospital Insurance Trust
2	Fund and the Federal Supplementary Medical Insurance
3	Trust Fund) to carry out the demonstration program—
4	(1) for fiscal year 2005, \$1,000,000, and
5	(2) for fiscal year 2006, \$6,000,000.
6	SEC. 923. MEDICARE PROVIDER OMBUDSMAN; MEDICARE
7	BENEFICIARY OMBUDSMAN.
8	(a) Medicare Provider Ombudsman.—Section 1868
9	(42 U.S.C. 1395ee) is amended—
10	(1) by adding at the end of the heading the fol-
11	lowing: "; medicare provider ombudsman";
12	(2) by inserting "Practicing Physicians Advi-
13	SORY COUNCIL.—(1)" after "(a)";
14	(3) in paragraph (1), as so redesignated under
15	paragraph (2), by striking "in this section" and in-
16	serting "in this subsection";
17	(4) by redesignating subsections (b) and (c) as
18	paragraphs (2) and (3), respectively; and
19	(5) by adding at the end the following new sub-
20	section:
21	"(b) Medicare Provider Ombudsman.—The Sec-
22	retary shall appoint within the Department of Health and
23	Human Services a Medicare Provider Ombudsman. The
24	Omhudsman shall—



1	"(1) provide assistance, on a confidential basis,
2	to providers of services and suppliers with respect to
3	complaints, grievances, and requests for information
4	concerning the programs under this title (including
5	provisions of title XI insofar as they relate to this
6	title and are not administered by the Office of the In-
7	spector General of the Department of Health and
8	Human Services) and in the resolution of unclear or
9	conflicting guidance given by the Secretary and medi-
10	care contractors to such providers of services and sup-
11	pliers regarding such programs and provisions and
12	requirements under this title and such provisions;
13	and
14	"(2) submit recommendations to the Secretary
15	for improvement in the administration of this title
16	and such provisions, including—
17	"(A) recommendations to respond to recur-
18	ring patterns of confusion in this title and such
19	provisions (including recommendations regard-
20	ing suspending imposition of sanctions where
21	there is widespread confusion in program ad-
22	ministration), and
23	"(B) recommendations to provide for an ap-
24	propriate and consistent response (including not

providing for audits) in cases of self-identified



1	overpayments by providers of services and sup-
2	pliers.
3	The Ombudsman shall not serve as an advocate for any in-
4	creases in payments or new coverage of services, but may
5	identify issues and problems in payment or coverage poli-
6	cies.".
7	(b) Medicare Beneficiary Ombudsman.—Title
8	XVIII, as previously amended, is amended by inserting
9	after section 1809 the following new section:
10	"MEDICARE BENEFICIARY OMBUDSMAN
11	"Sec. 1810. (a) In General.—The Secretary shall
12	appoint within the Department of Health and Human
13	Services a Medicare Beneficiary Ombudsman who shall
14	have expertise and experience in the fields of health care
15	and education of (and assistance to) individuals entitled
16	to benefits under this title.
17	"(b) Duties.—The Medicare Beneficiary Ombudsman
18	shall—
19	"(1) receive complaints, grievances, and requests
20	for information submitted by individuals entitled to
21	benefits under part A or enrolled under part B, or
22	both, with respect to any aspect of the medicare pro-
23	gram;
24	"(2) provide assistance with respect to com-
25	plaints, grievances, and requests referred to in para-
26	graph (1), including—



1	"(A) assistance in collecting relevant infor-
2	mation for such individuals, to seek an appeal of
3	a decision or determination made by a fiscal
4	$intermediary,\ carrier,\ Medicare + Choice\ organi-$
5	zation, or the Secretary;
6	"(B) assistance to such individuals with
7	any problems arising from disenrollment from a
8	Medicare+Choice plan under part C; and
9	"(C) assistance to such individuals in pre-
10	senting information under section 1860D-
11	2(b)(4)(D)(v); and
12	"(3) submit annual reports to Congress and the
13	Secretary that describe the activities of the Office and
14	that include such recommendations for improvement
15	in the administration of this title as the Ombudsman
16	determines appropriate.
17	The Ombudsman shall not serve as an advocate for any in-
18	creases in payments or new coverage of services, but may
19	identify issues and problems in payment or coverage poli-
20	cies.
21	"(c) Working With Health Insurance Coun-
22	SELING PROGRAMS.—To the extent possible, the Ombuds-
23	man shall work with health insurance counseling programs
24	(receiving funding under section 4360 of Omnibus Budget
25	Reconciliation Act of 1990) to facilitate the provision of in-



- 1 formation to individuals entitled to benefits under part A
- 2 or enrolled under part B, or both regarding
- 3 Medicare+Choice plans and changes to those plans. Noth-
- 4 ing in this subsection shall preclude further collaboration
- 5 between the Ombudsman and such programs.".
- 6 (c) Deadline for Appointment.—The Secretary
- 7 shall appoint the Medicare Provider Ombudsman and the
- 8 Medicare Beneficiary Ombudsman, under the amendments
- 9 made by subsections (a) and (b), respectively, by not later
- 10 than 1 year after the date of the enactment of this Act.
- 11 (d) Funding.—There are authorized to be appro-
- 12 priated to the Secretary (in appropriate part from the Fed-
- 13 eral Hospital Insurance Trust Fund and the Federal Sup-
- 14 plementary Medical Insurance Trust Fund) to carry out
- 15 the provisions of subsection (b) of section 1868 of the Social
- 16 Security Act (relating to the Medicare Provider Ombuds-
- 17 man), as added by subsection (a)(5) and section 1807 of
- 18 such Act (relating to the Medicare Beneficiary Ombuds-
- 19 man), as added by subsection (b), such sums as are nec-
- 20 essary for fiscal year 2004 and each succeeding fiscal year.
- 21 (e) Use of Central, Toll-Free Number (1–800–
- 22 MEDICARE).—
- 23 (1) Phone triage system; listing in medi-
- 24 CARE HANDBOOK INSTEAD OF OTHER TOLL-FREE
- 25 NUMBERS.—Section 1804(b) (42 U.S.C. 1395b–2(b))



is amended by adding at the end the following: "The
Secretary shall provide, through the toll-free number
1-800-MEDICARE, for a means by which individ-
uals seeking information about, or assistance with,
such programs who phone such toll-free number are
transferred (without charge) to appropriate entities
for the provision of such information or assistance.
Such toll-free number shall be the toll-free number
listed for general information and assistance in the
annual notice under subsection (a) instead of the list-
ing of numbers of individual contractors.".

## (2) Monitoring accuracy.—

(A) STUDY.—The Comptroller General of the United States shall conduct a study to monitor the accuracy and consistency of information provided to individuals entitled to benefits under part A or enrolled under part B, or both, through the toll-free number 1–800–MEDICARE, including an assessment of whether the information provided is sufficient to answer questions of such individuals. In conducting the study, the Comptroller General shall examine the education and training of the individuals providing information through such number.



1	(B) Report.—Not later than 1 year after
2	the date of the enactment of this Act, the Comp-
3	troller General shall submit to Congress a report
4	on the study conducted under subparagraph (A).
5	SEC. 924. BENEFICIARY OUTREACH DEMONSTRATION PRO-
6	GRAM.
7	(a) In General.—The Secretary shall establish a
8	demonstration program (in this section referred to as the
9	"demonstration program") under which medicare special-
10	ists employed by the Department of Health and Human
11	Services provide advice and assistance to individuals enti-
12	tled to benefits under part A of title XVIII of the Social
13	Security Act, or enrolled under part B of such title, or both,
14	regarding the medicare program at the location of existing
15	$local\ of fices\ of\ the\ Social\ Security\ Administration.$
16	(b) Locations.—
17	(1) In general.—The demonstration program
18	shall be conducted in at least 6 offices or areas. Sub-
19	ject to paragraph (2), in selecting such offices and
20	areas, the Secretary shall provide preference for offices
21	with a high volume of visits by individuals referred
22	to in subsection (a).
23	(2) Assistance for rural beneficiaries.—
24	The Secretary shall provide for the selection of at
25	least 2 rural areas to participate in the demonstra-



1	tion program. In conducting the demonstration pro-
2	gram in such rural areas, the Secretary shall provide
3	for medicare specialists to travel among local offices
4	in a rural area on a scheduled basis.
5	(c) Duration.—The demonstration program shall be
6	conducted over a 3-year period.
7	(d) Evaluation and Report.—
8	(1) EVALUATION.—The Secretary shall provide
9	for an evaluation of the demonstration program. Such
10	evaluation shall include an analysis of—
11	(A) utilization of, and satisfaction of those
12	individuals referred to in subsection (a) with, the
13	assistance provided under the program; and
14	(B) the cost-effectiveness of providing bene-
15	ficiary assistance through out-stationing medi-
16	care specialists at local offices of the Social Secu-
17	$rity\ Administration.$
18	(2) Report.—The Secretary shall submit to
19	Congress a report on such evaluation and shall in-
20	clude in such report recommendations regarding the
21	feasibility of permanently out-stationing medicare
22	specialists at local offices of the Social Security Ad-
23	ministration.



1	SEC. 925. INCLUSION OF ADDITIONAL INFORMATION IN NO-
2	TICES TO BENEFICIARIES ABOUT SKILLED
3	NURSING FACILITY BENEFITS.
4	(a) In General.—The Secretary shall provide that in
5	medicare beneficiary notices provided (under section
6	1806(a) of the Social Security Act, 42 U.S.C. 1395b-7(a))
7	with respect to the provision of post-hospital extended care
8	services under part A of title XVIII of the Social Security
9	Act, there shall be included information on the number of
10	days of coverage of such services remaining under such part
11	for the medicare beneficiary and spell of illness involved.
12	(b) Effective Date.—Subsection (a) shall apply to
13	notices provided during calendar quarters beginning more
14	than 6 months after the date of the enactment of this Act.
15	SEC. 926. INFORMATION ON MEDICARE-CERTIFIED SKILLED
16	NURSING FACILITIES IN HOSPITAL DIS-
17	CHARGE PLANS.
18	(a) Availability of Data.—The Secretary shall pub-
19	licly provide information that enables hospital discharge
20	planners, medicare beneficiaries, and the public to identify
21	skilled nursing facilities that are participating in the medi-
22	care program.
23	(b) Inclusion of Information in Certain Hos-
24	PITAL DISCHARGE PLANS.—
25	(1) In General.—Section $1861(ee)(2)(D)$ (42)
26	$U.S.C.\ 1395x(ee)(2)(D))$ is amended—



1	(A) by striking "hospice services" and in-
2	serting "hospice care and post-hospital extended
3	care services"; and
4	(B) by inserting before the period at the end
5	the following: "and, in the case of individuals
6	who are likely to need post-hospital extended care
7	services, the availability of such services through
8	facilities that participate in the program under
9	this title and that serve the area in which the
10	patient resides".
11	(2) Effective date.—The amendments made
12	by paragraph (1) shall apply to discharge plans made
13	on or after such date as the Secretary shall specify,
14	but not later than 6 months after the date the Sec-
15	retary provides for availability of information under
16	subsection (a).
17	Subtitle D—Appeals and Recovery
18	SEC. 931. TRANSFER OF RESPONSIBILITY FOR MEDICARE
19	APPEALS.
20	(a) Transition Plan.—
21	(1) In general.—Not later than October 1,
22	2004, the Commissioner of Social Security and the
23	Secretary shall develop and transmit to Congress and
24	the Comptroller General of the United States a plan
25	under which the functions of administrative law



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1	judges responsible for hearing cases under title XVIII
2	of the Social Security Act (and related provisions in
3	title XI of such Act) are transferred from the responsi-
4	bility of the Commissioner and the Social Security
5	Administration to the Secretary and the Department
6	of Health and Human Services.
7	(2) GAO EVALUATION.—The Comptroller Gen-
8	eral of the United States shall evaluate the plan and,
9	not later than the date that is 6 months after the date
10	on which the plan is received by the Comptroller Gen-
11	eral, shall submit to Congress a report on such eval-
12	uation.
13	(b) Transfer of Adjudication Authority.—
14	(1) In general.—Not earlier than July 1, 2005,
15	and not later than October 1, 2005, the Commissioner
16	of Social Security and the Secretary shall implement
17	the transition plan under subsection (a) and transfer
18	the administrative law judge functions described in
19	such subsection from the Social Security Administra-
20	tion to the Secretary.
21	(2) Assuring independence of judges.—The
22	Secretary shall assure the independence of adminis-
23	trative law judges performing the administrative law
24	judge functions transferred under paragraph (1) from

the Centers for Medicare & Medicaid Services and its



contractors. In order to assure such independence, the
Secretary shall place such judges in an administra-
tive office that is organizationally and functionally
separate from such Centers. Such judges shall report
to, and be under the general supervision of, the Sec-
retary, but shall not report to, or be subject to super-
vision by, another other officer of the Department.

- (3) Geographic distribution.—The Secretary shall provide for an appropriate geographic distribution of administrative law judges performing the administrative law judge functions transferred under paragraph (1) throughout the United States to ensure timely access to such judges.
- (4) HIRING AUTHORITY.—Subject to the amounts provided in advance in appropriations Act, the Secretary shall have authority to hire administrative law judges to hear such cases, giving priority to those judges with prior experience in handling medicare appeals and in a manner consistent with paragraph (3), and to hire support staff for such judges.
- (5) FINANCING.—Amounts payable under law to the Commissioner for administrative law judges performing the administrative law judge functions transferred under paragraph (1) from the Federal Hospital Insurance Trust Fund and the Federal Supple-



1	mentary Medical Insurance Trust Fund shall become
2	payable to the Secretary for the functions so trans-
3	ferred.
4	(6) Shared resources.—The Secretary shall
5	enter into such arrangements with the Commissioner
6	as may be appropriate with respect to transferred
7	functions of administrative law judges to share office
8	space, support staff, and other resources, with appro-
9	priate reimbursement from the Trust Funds described
10	in paragraph (5).
11	(c) Increased Financial Support.—In addition to
12	any amounts otherwise appropriated, to ensure timely ac-
13	tion on appeals before administrative law judges and the
14	Departmental Appeals Board consistent with section 1869
15	of the Social Security Act (as amended by section 521 of
16	BIPA, 114 Stat. 2763A-534), there are authorized to be ap-
17	propriated (in appropriate part from the Federal Hospital
18	Insurance Trust Fund and the Federal Supplementary
19	Medical Insurance Trust Fund) to the Secretary such sums
20	as are necessary for fiscal year 2005 and each subsequent
21	fiscal year to—
22	(1) increase the number of administrative law
23	judges (and their staffs) under subsection (b)(4);



1	(2) improve education and training opportuni-
2	ties for administrative law judges (and their staffs);
3	and
4	(3) increase the staff of the Departmental Ap-
5	peals Board.
6	(d) Conforming Amendment.—Section
7	1869(f)(2)(A)(i) (42 U.S.C. 1395ff(f)(2)(A)(i)), as added by
8	section 522(a) of BIPA (114 Stat. 2763A-543), is amended
9	by striking "of the Social Security Administration".
10	SEC. 932. PROCESS FOR EXPEDITED ACCESS TO REVIEW.
11	(a) Expedited Access to Judicial Review.—Sec-
12	tion 1869(b) (42 U.S.C. 1395ff(b)) as amended by BIPA,
13	is amended—
14	(1) in paragraph (1)(A), by inserting ", subject
15	to paragraph (2)," before "to judicial review of the
16	Secretary's final decision";
17	(2) in paragraph $(1)(F)$ —
18	(A) by striking clause (ii);
19	(B) by striking "PROCEEDING" and all that
20	follows through "DETERMINATION" and inserting
21	"DETERMINATIONS AND RECONSIDERATIONS";
22	and
23	(C) by redesignating subclauses (I) and (II)
24	as clauses (i) and (ii) and by moving the inden-



1	tation of such subclauses (and the matter that
2	follows) 2 ems to the left; and
3	(3) by adding at the end the following new para-
4	graph:
5	"(2) Expedited access to judicial re-
6	VIEW.—
7	"(A) In general.—The Secretary shall es-
8	tablish a process under which a provider of serv-
9	ices or supplier that furnishes an item or service
10	or an individual entitled to benefits under part
11	A or enrolled under part B, or both, who has
12	filed an appeal under paragraph (1) may obtain
13	access to judicial review when a review panel
14	(described in subparagraph (D)), on its own mo-
15	tion or at the request of the appellant, deter-
16	mines that no entity in the administrative ap-
17	peals process has the authority to decide the
18	question of law or regulation relevant to the mat-
19	ters in controversy and that there is no material
20	issue of fact in dispute. The appellant may make
21	such request only once with respect to a question
22	of law or regulation in a case of an appeal.
23	"(B) Prompt determinations.—If, after
24	or coincident with appropriately filing a request

for an administrative hearing, the appellant re-



1	quests a determination by the appropriate review
2	panel that no review panel has the authority to
3	decide the question of law or regulations relevant
4	to the matters in controversy and that there is
5	no material issue of fact in dispute and if such
6	request is accompanied by the documents and
7	materials as the appropriate review panel shall
8	require for purposes of making such determina-
9	tion, such review panel shall make a determina-
10	tion on the request in writing within 60 days
11	after the date such review panel receives the re-
12	quest and such accompanying documents and
13	materials. Such a determination by such review
14	panel shall be considered a final decision and
15	not subject to review by the Secretary.
16	"(C) Access to Judicial Review.—
17	"(i) In general.—If the appropriate
18	review panel—
19	"(I) determines that there are no
20	material issues of fact in dispute and
21	that the only issue is one of law or reg-
22	ulation that no review panel has the
23	authority to decide; or



1	"(II) fails to make such deter-
2	mination within the period provided
3	$under\ subparagraph\ (B);$
4	then the appellant may bring a civil action
5	as described in this subparagraph.
6	"(ii) Deadline for filing.—Such
7	action shall be filed, in the case described
8	in—
9	"(I) clause (i)(I), within 60 days
10	of date of the determination described
11	in such subparagraph; or
12	"(II) clause (i)(II), within $60$
13	days of the end of the period provided
14	under subparagraph (B) for the deter-
15	mination.
16	"(iii) Venue.—Such action shall be
17	brought in the district court of the United
18	States for the judicial district in which the
19	appellant is located (or, in the case of an
20	action brought jointly by more than one ap-
21	plicant, the judicial district in which the
22	greatest number of applicants are located)
23	or in the district court for the District of
24	Columbia.



1	"(iv) Interest on amounts in con-
2	Troversy.—Where a provider of services or
3	supplier seeks judicial review pursuant to
4	this paragraph, the amount in controversy
5	shall be subject to annual interest beginning
6	on the first day of the first month beginning
7	after the 60-day period as determined pur-
8	suant to clause (ii) and equal to the rate of
9	interest on obligations issued for purchase
10	by the Federal Hospital Insurance Trust
11	Fund and by the Federal Supplementary
12	Medical Insurance Trust Fund for the
13	month in which the civil action authorized
14	under this paragraph is commenced, to be
15	awarded by the reviewing court in favor of
16	the prevailing party. No interest awarded
17	pursuant to the preceding sentence shall be
18	deemed income or cost for the purposes of
19	determining reimbursement due providers of
20	services or suppliers under this Act.
21	"(D) REVIEW PANELS.—For purposes of
22	this subsection, a 'review panel' is a panel con-
23	sisting of 3 members (who shall be administra-
24	tive law judges, members of the Departmental

Appeals Board, or qualified individuals associ-



1	ated with a qualified independent contractor (as
2	defined in subsection $(c)(2)$ or with another
3	independent entity) designated by the Secretary
4	for purposes of making determinations under
5	this paragraph.".
6	(b) Application to Provider Agreement Deter-
7	MINATIONS.—Section 1866(h)(1) (42 U.S.C. 1395cc(h)(1))
8	is amended—
9	(1) by inserting "(A)" after "(h)(1)"; and
10	(2) by adding at the end the following new sub-
11	paragraph:
12	"(B) An institution or agency described in subpara-
13	graph (A) that has filed for a hearing under subparagraph
14	(A) shall have expedited access to judicial review under this
15	subparagraph in the same manner as providers of services,
16	suppliers, and individuals entitled to benefits under part
17	A or enrolled under part B, or both, may obtain expedited
18	access to judicial review under the process established under
19	section 1869(b)(2). Nothing in this subparagraph shall be
20	construed to affect the application of any remedy imposed
21	under section 1819 during the pendency of an appeal under
22	this subparagraph.".
23	(c) Effective Date.—The amendments made by this
24	section shall apply to appeals filed on or after October 1,
25	2004.



1	(d) Expedited Review of Certain Provider
2	AGREEMENT DETERMINATIONS.—
3	(1) TERMINATION AND CERTAIN OTHER IMME-
4	DIATE REMEDIES.—The Secretary shall develop and
5	implement a process to expedite proceedings under
6	sections 1866(h) of the Social Security Act (42 U.S.C.
7	1395cc(h)) in which the remedy of termination of
8	participation, or a remedy described in clause (i) or
9	(iii) of section $1819(h)(2)(B)$ of such Act (42 U.S.C.
10	1395i-3(h)(2)(B)) which is applied on an immediate
11	basis, has been imposed. Under such process priority
12	shall be provided in cases of termination.
13	(2) Increased financial support.—In addi-
14	tion to any amounts otherwise appropriated, to re-
15	duce by 50 percent the average time for administra-
16	tive determinations on appeals under section 1866(h)
17	of the Social Security Act (42 U.S.C. 1395cc(h)),
18	there are authorized to be appropriated (in appro-
19	priate part from the Federal Hospital Insurance
20	Trust Fund and the Federal Supplementary Medical
21	Insurance Trust Fund) to the Secretary such addi-
22	tional sums for fiscal year 2005 and each subsequent
23	fiscal year as may be necessary. The purposes for
24	which such amounts are available include increasing

the number of administrative law judges (and their



1	staffs) and the appellate level staff at the Depart-
2	mental Appeals Board of the Department of Health
3	and Human Services and educating such judges and
4	staffs on long-term care issues.
5	SEC. 933. REVISIONS TO MEDICARE APPEALS PROCESS.
6	(a) Requiring Full and Early Presentation of
7	EVIDENCE.—
8	(1) In General.—Section 1869(b) (42 U.S.C.
9	1395ff(b)), as amended by BIPA and as amended by
10	section 932(a), is further amended by adding at the
11	end the following new paragraph:
12	"(3) Requiring full and early presen-
13	TATION OF EVIDENCE BY PROVIDERS.—A provider of
14	services or supplier may not introduce evidence in
15	any appeal under this section that was not presented
16	at the reconsideration conducted by the qualified
17	independent contractor under subsection (c), unless
18	there is good cause which precluded the introduction
19	of such evidence at or before that reconsideration.".
20	(2) Effective date.—The amendment made by
21	paragraph (1) shall take effect on October 1, 2004.
22	(b) Use of Patients' Medical Records.—Section
23	1869(c)(3)(B)(i) (42 U.S.C. 1395ff(c)(3)(B)(i)), as amended
24	by BIPA, is amended by inserting "(including the medical



1	records of the individual involved)" after "clinical experi-
2	ence".
3	(c) Notice Requirements for Medicare Ap-
4	PEALS.—
5	(1) Initial determinations and redeter-
6	MINATIONS.—Section 1869(a) (42 U.S.C. 1395ff(a)),
7	as amended by BIPA, is amended by adding at the
8	end the following new paragraphs:
9	"(4) Requirements of notice of determina-
10	Tions.—With respect to an initial determination in-
11	sofar as it results in a denial of a claim for benefits—
12	"(A) the written notice on the determina-
13	tion shall include—
14	"(i) the reasons for the determination,
15	including whether a local medical review
16	policy or a local coverage determination
17	$was\ used;$
18	"(ii) the procedures for obtaining addi-
19	tional information concerning the deter-
20	mination, including the information de-
21	scribed in subparagraph (B); and
22	"(iii) notification of the right to seek a
23	redetermination or otherwise appeal the de-
24	tormination and instructions on how to ini



1	tiate such a redetermination under this sec-
2	tion; and
3	"(B) the person provided such notice may
4	obtain, upon request, the specific provision of the
5	policy, manual, or regulation used in making
6	the determination.
7	"(5) Requirements of notice of redeter-
8	MINATIONS.—With respect to a redetermination inso-
9	far as it results in a denial of a claim for benefits—
10	"(A) the written notice on the redetermina-
11	tion shall include—
12	"(i) the specific reasons for the redeter-
13	mination;
14	"(ii) as appropriate, a summary of the
15	clinical or scientific evidence used in mak-
16	ing the redetermination;
17	"(iii) a description of the procedures
18	for obtaining additional information con-
19	cerning the redetermination; and
20	"(iv) notification of the right to appeal
21	the redetermination and instructions on
22	how to initiate such an appeal under this
23	section;
24	"(B) such written notice shall be provided
25	in printed form and written in a manner cal-



1	culated to be understood by the individual enti-
2	tled to benefits under part A or enrolled under
3	part B, or both; and
4	"(C) the person provided such notice may
5	obtain, upon request, information on the specific
6	provision of the policy, manual, or regulation
7	used in making the redetermination.".
8	(2) Reconsiderations.—Section $1869(c)(3)(E)$
9	(42 U.S.C. $1395ff(c)(3)(E)$ ), as amended by BIPA, is
10	amended—
11	(A) by inserting "be written in a manner
12	calculated to be understood by the individual en-
13	titled to benefits under part A or enrolled under
14	part B, or both, and shall include (to the extent
15	appropriate)" after "in writing,"; and
16	(B) by inserting "and a notification of the
17	right to appeal such determination and instruc-
18	tions on how to initiate such appeal under this
19	section" after "such decision,".
20	(3) Appeals.—Section 1869(d) (42 U.S.C.
21	1395ff(d)), as amended by BIPA, is amended—
22	(A) in the heading, by inserting "; Notice"
23	after "Secretary"; and
24	(B) by adding at the end the following new
25	paragraph:



1	"(4) Notice.—Notice of the decision of an ad-
2	ministrative law judge shall be in writing in a man-
3	ner calculated to be understood by the individual en-
4	titled to benefits under part A or enrolled under part
5	B, or both, and shall include—
6	"(A) the specific reasons for the determina-
7	tion (including, to the extent appropriate, a
8	summary of the clinical or scientific evidence
9	used in making the determination);
10	"(B) the procedures for obtaining addi-
11	tional information concerning the decision; and
12	"(C) notification of the right to appeal the
13	decision and instructions on how to initiate such
14	an appeal under this section.".
15	(4) Submission of record for appeal.—Sec-
16	tion $1869(c)(3)(J)(i)$ (42 U.S.C. $1395ff(c)(3)(J)(i)$ ) by
17	striking "prepare" and inserting "submit" and by
18	striking "with respect to" and all that follows through
19	"and relevant policies".
20	(d) Qualified Independent Contractors.—
21	(1) Eligibility requirements of qualified
22	INDEPENDENT CONTRACTORS.—Section 1869(c)(3) (42
23	U.S.C. 1395ff(c)(3)), as amended by BIPA, is
24	amended—



1	(A) in subparagraph (A), by striking "suffi-
2	cient training and expertise in medical science
3	and legal matters" and inserting "sufficient
4	medical, legal, and other expertise (including
5	knowledge of the program under this title) and
6	sufficient staffing"; and
7	(B) by adding at the end the following new
8	subparagraph:
9	"(K) Independence requirements.—
10	"(i) In general.—Subject to clause
11	(ii), a qualified independent contractor
12	shall not conduct any activities in a case
13	unless the entity—
14	"(I) is not a related party (as de-
15	fined in subsection $(g)(5)$ ;
16	"(II) does not have a material fa-
17	milial, financial, or professional rela-
18	tionship with such a party in relation
19	to such case; and
20	"(III) does not otherwise have a
21	conflict of interest with such a party.
22	"(ii) Exception for reasonable
23	Compensation.—Nothing in clause (i) shall
24	be construed to prohibit receipt by a quali-
25	fied independent contractor of compensation



1	from the Secretary for the conduct of activi-
2	ties under this section if the compensation
3	is provided consistent with clause (iii).
4	"(iii) Limitations on entity com-
5	PENSATION.—Compensation provided by the
6	Secretary to a qualified independent con-
7	tractor in connection with reviews under
8	this section shall not be contingent on any
9	decision rendered by the contractor or by
10	any reviewing professional.".
11	(2) Eligibility requirements for review-
12	ERS.—Section 1869 (42 U.S.C. 1395ff), as amended
13	by BIPA, is amended—
14	(A) by amending subsection $(c)(3)(D)$ to
15	read as follows:
16	"(D) Qualifications for reviewers.—
17	The requirements of subsection (g) shall be met
18	(relating to qualifications of reviewing profes-
19	sionals)."; and
20	(B) by adding at the end the following new
21	subsection:
22	"(g) Qualifications of Reviewers.—
23	"(1) In general.—In reviewing determinations
24	under this section, a qualified independent contractor
25	shall assure that—



1	"(A) each individual conducting a review
2	shall meet the qualifications of paragraph (2);
3	"(B) compensation provided by the con-
4	tractor to each such reviewer is consistent with
5	paragraph (3); and
6	"(C) in the case of a review by a panel de-
7	scribed in subsection $(c)(3)(B)$ composed of phy-
8	sicians or other health care professionals (each in
9	this subsection referred to as a 'reviewing profes-
10	sional'), a reviewing professional meets the
11	qualifications described in paragraph (4) and,
12	where a claim is regarding the furnishing of
13	treatment by a physician (allopathic or osteo-
14	pathic) or the provision of items or services by
15	a physician (allopathic or osteopathic), a review-
16	ing professional shall be a physician (allopathic
17	$or\ osteopathic).$
18	"(2) Independence.—
19	"(A) In general.—Subject to subpara-
20	graph (B), each individual conducting a review
21	in a case shall—
22	"(i) not be a related party (as defined
23	in paragraph (5));



1	"(ii) not have a material familial, fi-
2	nancial, or professional relationship with
3	such a party in the case under review; and
4	"(iii) not otherwise have a conflict of
5	interest with such a party.
6	"(B) Exception.—Nothing in subpara-
7	graph (A) shall be construed to—
8	"(i) prohibit an individual, solely on
9	the basis of a participation agreement with
10	a fiscal intermediary, carrier, or other con-
11	tractor, from serving as a reviewing profes-
12	sional if—
13	"(I) the individual is not involved
14	in the provision of items or services in
15	the case under review;
16	"(II) the fact of such an agree-
17	ment is disclosed to the Secretary and
18	the individual entitled to benefits
19	under part A or enrolled under part B,
20	or both, (or authorized representative)
21	and neither party objects; and
22	"(III) the individual is not an
23	employee of the intermediary, carrier,
24	or contractor and does not provide
25	services exclusively or primarily to or



1	on behalf of such intermediary, carrier,
2	$or\ contractor;$
3	"(ii) prohibit an individual who has
4	staff privileges at the institution where the
5	treatment involved takes place from serving
6	as a reviewer merely on the basis of having
7	such staff privileges if the existence of such
8	privileges is disclosed to the Secretary and
9	such individual (or authorized representa-
10	tive), and neither party objects; or
11	"(iii) prohibit receipt of compensation
12	by a reviewing professional from a con-
13	tractor if the compensation is provided con-
14	sistent with paragraph (3).
15	For purposes of this paragraph, the term 'par-
16	ticipation agreement' means an agreement relat-
17	ing to the provision of health care services by the
18	individual and does not include the provision of
19	services as a reviewer under this subsection.
20	"(3) Limitations on reviewer compensa-
21	TION.—Compensation provided by a qualified inde-
22	pendent contractor to a reviewer in connection with
23	a review under this section shall not be contingent on
24	the decision rendered by the reviewer.



1	"(4) Licensure and expertise.—Each review-
2	ing professional shall be—
3	"(A) a physician (allopathic or osteopathic)
4	who is appropriately credentialed or licensed in
5	one or more States to deliver health care services
6	and has medical expertise in the field of practice
7	that is appropriate for the items or services at
8	$issue;\ or$
9	"(B) a health care professional who is le-
10	gally authorized in one or more States (in ac-
11	cordance with State law or the State regulatory
12	mechanism provided by State law) to furnish the
13	health care items or services at issue and has
14	medical expertise in the field of practice that is
15	appropriate for such items or services.
16	"(5) Related party defined.—For purposes
17	of this section, the term 'related party' means, with
18	respect to a case under this title involving a specific
19	individual entitled to benefits under part A or en-
20	rolled under part B, or both, any of the following:
21	"(A) The Secretary, the medicare adminis-
22	trative contractor involved, or any fiduciary, of-
23	ficer, director, or employee of the Department of
24	Health and Human Services, or of such con-



25

tractor.

1	"(B) The individual (or authorized rep-
2	resentative).
3	"(C) The health care professional that pro-
4	vides the items or services involved in the case.
5	"(D) The institution at which the items or
6	services (or treatment) involved in the case are
7	provided.
8	"(E) The manufacturer of any drug or
9	other item that is included in the items or serv-
10	ices involved in the case.
11	"(F) Any other party determined under any
12	regulations to have a substantial interest in the
13	case involved.".
14	(3) Reducing minimum number of qualified
15	INDEPENDENT CONTRACTORS.—Section 1869(c)(4) (42
16	U.S.C. 1395ff(c)(4)) is amended by striking "not
17	fewer than 12 qualified independent contractors under
18	this subsection" and inserting "with a sufficient num-
19	ber of qualified independent contractors (but not
20	fewer than 4 such contractors) to conduct reconsider-
21	ations consistent with the timeframes applicable
22	under this subsection".
23	(4) Effective date.—The amendments made

by paragraphs (1) and (2) shall be effective as if in-



1	cluded in the enactment of the respective provisions of
2	subtitle C of title V of BIPA, (114 Stat. 2763A-534).
3	(5) Transition.—In applying section 1869(g) of
4	the Social Security Act (as added by paragraph (2)),
5	any reference to a medicare administrative contractor
6	shall be deemed to include a reference to a fiscal
7	intermediary under section 1816 of the Social Secu-
8	rity Act (42 U.S.C. 1395h) and a carrier under sec-
9	tion 1842 of such Act (42 U.S.C. 1395u).
10	SEC. 934. PREPAYMENT REVIEW.
11	(a) In General.—Section 1874A, as added by section
12	911(a)(1) and as amended by sections 912(b), 921(b)(1),
13	and 921(c)(1), is further amended by adding at the end the
14	following new subsection:
15	"(h) Conduct of Prepayment Review.—
16	"(1) Conduct of random prepayment re-
17	VIEW.—
18	"(A) In General.—A medicare adminis-
19	trative contractor may conduct random prepay-
20	ment review only to develop a contractor-wide or
21	program-wide claims payment error rates or
22	under such additional circumstances as may be
23	provided under regulations, developed in con-
24	sultation with providers of services and sup-
25	pliers.



1	"(B) Use of standard protocols when
2	CONDUCTING PREPAYMENT REVIEWS.—When a
3	medicare administrative contractor conducts a
4	random prepayment review, the contractor may
5	conduct such review only in accordance with a
6	standard protocol for random prepayment audits
7	developed by the Secretary.
8	"(C) Construction.—Nothing in this
9	paragraph shall be construed as preventing the
10	denial of payments for claims actually reviewed
11	under a random prepayment review.
12	"(D) Random prepayment review.—For
13	purposes of this subsection, the term 'random
14	prepayment review' means a demand for the
15	production of records or documentation absent
16	cause with respect to a claim.
17	"(2) Limitations on non-random prepayment
18	REVIEW.—
19	"(A) Limitations on initiation of non-
20	RANDOM PREPAYMENT REVIEW.—A medicare ad-
21	ministrative contractor may not initiate non-
22	random prepayment review of a provider of serv-
23	ices or supplier based on the initial identifica-
24	tion by that provider of services or supplier of

an improper billing practice unless there is a



1	likelihood of sustained or high level of payment
2	error (as defined in subsection $(i)(3)(A)$ ).
3	"(B) TERMINATION OF NON-RANDOM PRE-
4	PAYMENT REVIEW.—The Secretary shall issue
5	regulations relating to the termination, includ-
6	ing termination dates, of non-random prepay-
7	ment review. Such regulations may vary such a
8	termination date based upon the differences in
9	the circumstances triggering prepayment re-
10	view.".
11	(b) Effective Date.—
12	(1) In general.—Except as provided in this
13	subsection, the amendment made by subsection (a)
14	shall take effect 1 year after the date of the enactment
15	$of\ this\ Act.$
16	(2) Deadline for promulgation of certain
17	REGULATIONS.—The Secretary shall first issue regula-
18	tions under section 1874A(h) of the Social Security
19	Act, as added by subsection (a), by not later than 1
20	year after the date of the enactment of this Act.
21	(3) Application of standard protocols for
22	RANDOM PREPAYMENT REVIEW.—Section
23	1874A(h)(1)(B) of the Social Security Act, as added
24	by subsection (a), shall apply to random prepayment

 $reviews \ \ conducted \ \ on \ \ or \ \ after \ such \ \ date \ \ (not \ \ later$ 



1	than 1 year after the date of the enactment of this
2	Act) as the Secretary shall specify.
3	(c) Application to Fiscal Intermediaries and
4	Carriers.—The provisions of section 1874A(h) of the So-
5	cial Security Act, as added by subsection (a), shall apply
6	to each fiscal intermediary under section 1816 of the Social
7	Security Act (42 U.S.C. 1395h) and each carrier under sec-
8	tion 1842 of such Act (42 U.S.C. 1395u) in the same man-
9	ner as they apply to medicare administrative contractors
10	under such provisions.
11	SEC. 935. RECOVERY OF OVERPAYMENTS.
12	(a) In General.—Section 1893 (42 U.S.C. 1395ddd)
13	is amended by adding at the end the following new sub-
14	section:
15	"(f) Recovery of Overpayments.—
16	"(1) Use of repayment plans.—
17	"(A) In General.—If the repayment, with-
18	in 30 days by a provider of services or supplier,
19	of an overpayment under this title would con-
20	stitute a hardship (as defined in subparagraph
21	(B)), subject to subparagraph (C), upon request
22	of the provider of services or supplier the Sec-
23	retary shall enter into a plan with the provider
24	of services or supplier for the repayment
25	(through offset or otherwise) of such overpayment



1	over a period of at least 6 months but not longer
2	than 3 years (or not longer than 5 years in the
3	case of extreme hardship, as determined by the
4	Secretary). Interest shall accrue on the balance
5	through the period of repayment. Such plan shall
6	meet terms and conditions determined to be ap-
7	propriate by the Secretary.
8	"(B) Hardship.—
9	"(i) In general.—For purposes of
10	subparagraph (A), the repayment of an
11	overpayment (or overpayments) within 30
12	days is deemed to constitute a hardship if—
13	"(I) in the case of a provider of
14	services that files cost reports, the ag-
15	gregate amount of the overpayments
16	exceeds 10 percent of the amount paid
17	under this title to the provider of serv-
18	ices for the cost reporting period cov-
19	ered by the most recently submitted
20	cost report; or
21	"(II) in the case of another pro-
22	vider of services or supplier, the aggre-
23	gate amount of the overpayments ex-
24	ceeds 10 percent of the amount paid

under this title to the provider of serv-



1	ices or supplier for the previous cal-
2	endar year.
3	"(ii) Rule of application.—The
4	Secretary shall establish rules for the appli-
5	cation of this subparagraph in the case of a
6	provider of services or supplier that was not
7	paid under this title during the previous
8	year or was paid under this title only dur-
9	ing a portion of that year.
10	"(iii) Treatment of previous over-
11	PAYMENTS.—If a provider of services or
12	supplier has entered into a repayment plan
13	under subparagraph (A) with respect to a
14	specific overpayment amount, such payment
15	amount under the repayment plan shall not
16	be taken into account under clause (i) with
17	respect to subsequent overpayment amounts.
18	"(C) Exceptions.—Subparagraph (A)
19	shall not apply if—
20	"(i) the Secretary has reason to suspect
21	that the provider of services or supplier
22	may file for bankruptcy or otherwise cease
23	to do business or discontinue participation
24	in the program under this title; or



1	"(ii) there is an indication of fraud or
2	abuse committed against the program.
3	"(D) Immediate collection if violation
4	OF REPAYMENT PLAN.—If a provider of services
5	or supplier fails to make a payment in accord-
6	ance with a repayment plan under this para-
7	graph, the Secretary may immediately seek to
8	offset or otherwise recover the total balance out-
9	standing (including applicable interest) under
10	the repayment plan.
11	"(E) Relation to no fault provision.—
12	Nothing in this paragraph shall be construed as
13	affecting the application of section 1870(c) (re-
14	lating to no adjustment in the cases of certain
15	overpayments).
16	"(2) Limitation on recoupment.—
17	"(A) In general.—In the case of a pro-
18	vider of services or supplier that is determined to
19	have received an overpayment under this title
20	and that seeks a reconsideration by a qualified
21	independent contractor on such determination
22	under section 1869(b)(1), the Secretary may not
23	take any action (or authorize any other person,
24	including any medicare contractor, as defined in

subparagraph (C)) to recoup the overpayment



1	until the date the decision on the reconsideration
2	has been rendered. If the provisions of section
3	1869(b)(1) (providing for such a reconsideration
4	by a qualified independent contractor) are not in
5	effect, in applying the previous sentence any ref-
6	erence to such a reconsideration shall be treated
7	as a reference to a redetermination by the fiscal
8	intermediary or carrier involved.
9	"(B) Collection with interest.—Inso-
10	far as the determination on such appeal is
11	against the provider of services or supplier, in-
12	terest on the overpayment shall accrue on and
13	after the date of the original notice of overpay-
14	ment. Insofar as such determination against the
15	provider of services or supplier is later reversed,
16	the Secretary shall provide for repayment of the
17	amount recouped plus interest at the same rate
18	as would apply under the previous sentence for
19	the period in which the amount was recouped.
20	"(C) Medicare contractor defined.—
21	For purposes of this subsection, the term 'medi-
22	care contractor' has the meaning given such term
23	in section $1889(g)$ .
24	"(3) Limitation on use of extrapolation.—

 $A\ medicare\ contractor\ may\ not\ use\ extrapolation\ to$ 



1	determine overpayment amounts to be recovered by
2	recoupment, offset, or otherwise unless—
3	"(A) there is a sustained or high level of
4	payment error (as defined by the Secretary by
5	regulation); or
6	"(B) documented educational intervention
7	has failed to correct the payment error (as deter-
8	mined by the Secretary).
9	"(4) Provision of supporting documenta-
10	TION.—In the case of a provider of services or sup-
11	plier with respect to which amounts were previously
12	overpaid, a medicare contractor may request the peri-
13	odic production of records or supporting documenta-
14	tion for a limited sample of submitted claims to en-
15	sure that the previous practice is not continuing.
16	"(5) Consent settlement reforms.—
17	"(A) In general.—The Secretary may use
18	a consent settlement (as defined in subparagraph
19	(D)) to settle a projected overpayment.
20	"(B) Opportunity to submit additional
21	INFORMATION BEFORE CONSENT SETTLEMENT
22	OFFER.—Before offering a provider of services or
23	supplier a consent settlement, the Secretary
24	shall—



1	"(i) communicate to the provider of
2	services or supplier—
3	"(I) that, based on a review of the
4	medical records requested by the Sec-
5	retary, a preliminary evaluation of
6	those records indicates that there would
7	be an overpayment;
8	"(II) the nature of the problems
9	identified in such evaluation; and
10	"(III) the steps that the provider
11	of services or supplier should take to
12	address the problems; and
13	"(ii) provide for a 45-day period dur-
14	ing which the provider of services or sup-
15	plier may furnish additional information
16	concerning the medical records for the
17	claims that had been reviewed.
18	"(C) Consent settlement offer.—The
19	Secretary shall review any additional informa-
20	tion furnished by the provider of services or sup-
21	plier under subparagraph (B)(ii). Taking into
22	consideration such information, the Secretary
23	shall determine if there still appears to be an
24	overpayment. If so, the Secretary—



1	"(i) shall provide notice of such deter-
2	mination to the provider of services or sup-
3	plier, including an explanation of the rea-
4	son for such determination; and
5	"(ii) in order to resolve the overpay-
6	ment, may offer the provider of services or
7	supplier—
8	"(I) the opportunity for a statis-
9	tically valid random sample; or
10	"(II) a consent settlement.
11	The opportunity provided under clause $(ii)(I)$
12	does not waive any appeal rights with respect to
13	the alleged overpayment involved.
14	"(D) Consent settlement defined.—
15	For purposes of this paragraph, the term 'con-
16	sent settlement' means an agreement between the
17	Secretary and a provider of services or supplier
18	whereby both parties agree to settle a projected
19	overpayment based on less than a statistically
20	valid sample of claims and the provider of serv-
21	ices or supplier agrees not to appeal the claims
22	involved.
23	"(6) Notice of over-utilization of codes.—
24	The Secretary shall establish, in consultation with or-
25	ganizations representing the classes of providers of



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1	services and suppliers, a process under which the Sec-
2	retary provides for notice to classes of providers of
3	services and suppliers served by the contractor in
4	cases in which the contractor has identified that par-
5	ticular billing codes may be overutilized by that class
6	of providers of services or suppliers under the pro-
7	grams under this title (or provisions of title XI inso-
8	far as they relate to such programs).
9	"(7) Payment audits.—
10	"(A) Written notice for post-payment
11	AUDITS.—Subject to subparagraph (C), if a
12	medicare contractor decides to conduct a post-
13	payment audit of a provider of services or sup-
14	plier under this title, the contractor shall provide
15	the provider of services or supplier with written
16	notice (which may be in electronic form) of the
17	intent to conduct such an audit.
18	"(B) Explanation of findings for all
19	AUDITS.—Subject to subparagraph (C), if a
20	medicare contractor audits a provider of services
21	or supplier under this title, the contractor
22	shall—
23	"(i) give the provider of services or
24	supplier a full review and explanation of

the findings of the audit in a manner that



1	is understandable to the provider of services
2	or supplier and permits the development of
3	an appropriate corrective action plan;
4	"(ii) inform the provider of services or
5	supplier of the appeal rights under this title
6	as well as consent settlement options (which
7	are at the discretion of the Secretary);
8	"(iii) give the provider of services or
9	supplier an opportunity to provide addi-
10	tional information to the contractor; and
11	"(iv) take into account information
12	provided, on a timely basis, by the provider
13	of services or supplier under clause (iii).
14	"(C) Exception.—Subparagraphs (A) and
15	(B) shall not apply if the provision of notice or
16	findings would compromise pending law enforce-
17	ment activities, whether civil or criminal, or re-
18	veal findings of law enforcement-related audits.
19	"(8) Standard methodology for probe sam-
20	PLING.—The Secretary shall establish a standard
21	methodology for medicare contractors to use in select-
22	ing a sample of claims for review in the case of an
23	abnormal billing pattern.".
24	(b) Effective Dates and Deadlines.—



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1	(1) USE OF REPAYMENT PLANS.—Section
2	1893(f)(1) of the Social Security Act, as added by
3	subsection (a), shall apply to requests for repayment
4	plans made after the date of the enactment of this Act.
5	(2) Limitation on recoupment.—Section
6	1893(f)(2) of the Social Security Act, as added by
7	subsection (a), shall apply to actions taken after the
8	date of the enactment of this Act.
9	(3) USE OF EXTRAPOLATION.—Section
10	1893(f)(3) of the Social Security Act, as added by
11	subsection (a), shall apply to statistically valid ran-
12	dom samples initiated after the date that is 1 year
13	after the date of the enactment of this Act.
14	(4) Provision of supporting documenta-
15	TION.—Section 1893(f)(4) of the Social Security Act,
16	as added by subsection (a), shall take effect on the
17	date of the enactment of this Act.
18	(5) Consent settlement.—Section 1893(f)(5)
19	of the Social Security Act, as added by subsection (a),
20	shall apply to consent settlements entered into after
21	the date of the enactment of this Act.
22	(6) Notice of overutilization.—Not later
23	than 1 year after the date of the enactment of this
24	Act, the Secretary shall first establish the process for

 $notice\ of\ over utilization\ of\ billing\ codes\ under\ section$ 



1	1893A(f)(6) of the Social Security Act, as added by
2	subsection (a).
3	(7) Payment audits.—Section $1893A(f)$ (7) of
4	the Social Security Act, as added by subsection (a),
5	shall apply to audits initiated after the date of the
6	enactment of this Act.
7	(8) Standard for abnormal billing pat-
8	TERNS.—Not later than 1 year after the date of the
9	enactment of this Act, the Secretary shall first estab-
10	lish a standard methodology for selection of sample
11	claims for abnormal billing patterns under section
12	1893(f)(8) of the Social Security Act, as added by
13	subsection (a).
14	SEC. 936. PROVIDER ENROLLMENT PROCESS; RIGHT OF AP-
15	PEAL.
16	(a) In General.—Section 1866 (42 U.S.C. 1395cc)
17	is amended—
18	(1) by adding at the end of the heading the fol-
19	lowing: "; ENROLLMENT PROCESSES"; and
20	(2) by adding at the end the following new sub-
21	section:
22	"(j) Enrollment Process for Providers of Serv-
23	ICES AND SUPPLIERS.—
24	"(1) Enrollment process.—



1	"(A) In general.—The Secretary shall es-
2	tablish by regulation a process for the enrollment
3	of providers of services and suppliers under this
4	title.
5	"(B) Deadlines.—The Secretary shall es-
6	tablish by regulation procedures under which
7	there are deadlines for actions on applications
8	for enrollment (and, if applicable, renewal of en-
9	rollment). The Secretary shall monitor the per-
10	formance of medicare administrative contractors
11	in meeting the deadlines established under this
12	subparagraph.
13	"(C) Consultation before changing
14	PROVIDER ENROLLMENT FORMS.—The Secretary
15	shall consult with providers of services and sup-
16	pliers before making changes in the provider en-
17	rollment forms required of such providers and
18	suppliers to be eligible to submit claims for
19	which payment may be made under this title.
20	"(2) Hearing rights in cases of denial or
21	NON-RENEWAL.—A provider of services or supplier
22	whose application to enroll (or, if applicable, to renew
23	enrollment) under this title is denied may have a
24	hearing and judicial review of such denial under the

procedures that apply under subsection (h)(1)(A) to a



1	provider of services that is dissatisfied with a deter-
2	mination by the Secretary.".
3	(b) Effective Dates.—
4	(1) Enrollment process.—The Secretary shall
5	provide for the establishment of the enrollment process
6	under section 1866(j)(1) of the Social Security Act, as
7	added by subsection (a)(2), within 6 months after the
8	date of the enactment of this Act.
9	(2) Consultation.—Section $1866(j)(1)(C)$ of
10	the Social Security Act, as added by subsection
11	(a)(2), shall apply with respect to changes in provider
12	enrollment forms made on or after January 1, 2004.
13	(3) Hearing rights.—Section 1866(j)(2) of the
14	Social Security Act, as added by subsection (a)(2),
15	shall apply to denials occurring on or after such date
16	(not later than 1 year after the date of the enactment
17	of this Act) as the Secretary specifies.
18	SEC. 937. PROCESS FOR CORRECTION OF MINOR ERRORS
19	AND OMISSIONS WITHOUT PURSUING AP-
20	PEALS PROCESS.
21	(a) Claims.—The Secretary shall develop, in consulta-
22	tion with appropriate medicare contractors (as defined in
23	section 1889(g) of the Social Security Act, as inserted by
24	section 301(a)(1)) and representatives of providers of serv-
25	ices and suppliers, a process whereby, in the case of minor



1	errors or omissions (as defined by the Secretary) that are
2	detected in the submission of claims under the programs
3	under title XVIII of such Act, a provider of services or sup-
4	plier is given an opportunity to correct such an error or
5	omission without the need to initiate an appeal. Such proc-
6	ess shall include the ability to resubmit corrected claims.
7	(b) Permitting Use of Corrected and Supple-
8	MENTARY DATA.—
9	(1) In General.—Section $1886(d)(10)(D)(vi)$
10	(42 U.S.C. $1395ww(d)(10)(D)(vi)$ ) is amended by
11	adding after subclause (II) at the end the following:
12	"Notwithstanding subclause (I), a hospital may submit,
13	and the Secretary may accept upon verification, data that
14	corrects or supplements the data described in such subclause
15	without regard to whether the corrected or supplementary
16	data relate to a cost report that has been settled.".
17	(2) Effective date.—The amendment made by
18	paragraph (1) shall apply to fiscal years beginning
19	with fiscal year 2004.
20	(3) Submittal and resubmittal of applica-
21	TIONS PERMITTED FOR FISCAL YEAR 2004.—
22	(A) In General.—Notwithstanding any
23	other provision of law, a hospital may submit
24	(or resubmit) an application for a change de-
25	scribed in section $1886(d)(10)(C)(i)(II)$ of the



1		Social Security Act for fiscal year 2004 if the
2		hospital demonstrates on a timely basis to the
3		satisfaction of the Secretary that the use of cor-
4		rected or supplementary data under the amend-
5		ment made by paragraph (1) would materially
6		affect the approval of such an application.
7		(B) Application of budget neu-
8		TRALITY.—If one or more hospital's applications
9		are approved as a result of paragraph (1) and
10		subparagraph (A) for fiscal year 2004, the Sec-
11		retary shall make a proportional adjustment in
12		the standardized amounts determined under sec-
13		tion $1886(d)(3)$ of the Social Security Act (42)
14		$U.S.C.\ 1395ww(d)(3))$ for fiscal year 2004 to as-
15		sure that approval of such applications does not
16		result in aggregate payments under section
17		1886(d) of such Act that are greater or less than
18		those that would otherwise be made if paragraph
19		(1) and subparagraph (A) did not apply.
20	SEC. 938.	PRIOR DETERMINATION PROCESS FOR CERTAIN
21		ITEMS AND SERVICES; ADVANCE BENE-
22		FICIARY NOTICES.
23	(a) I	N GENERAL.—Section 1869 (42 U.S.C. 1395ff(b)),
24	as amend	ed by sections 521 and 522 of BIPA and section



1	933(d)(2)(B), is further amended by adding at the end the
2	following new subsection:
3	"(h) Prior Determination Process for Certain
4	Items and Services.—
5	"(1) Establishment of process.—
6	"(A) In general.—With respect to a medi-
7	care administrative contractor that has a con-
8	tract under section 1874A that provides for mak-
9	ing payments under this title with respect to eli-
10	gible items and services described in subpara-
11	graph (C), the Secretary shall establish a prior
12	determination process that meets the require-
13	ments of this subsection and that shall be applied
14	by such contractor in the case of eligible request-
15	ers.
16	"(B) Eligible requester.—For purposes
17	of this subsection, each of the following shall be
18	an eligible requester:
19	"(i) A physician, but only with respect
20	to eligible items and services for which the
21	physician may be paid directly.
22	"(ii) An individual entitled to benefits
23	under this title, but only with respect to an
24	item or service for which the individual re-
25	ceives, from the physician who may be paid



1	directly for the item or service, an advance
2	beneficiary notice under section 1879(a)
3	that payment may not be made (or may no
4	longer be made) for the item or service
5	under this title.
6	"(C) Eligible items and services.—For
7	purposes of this subsection and subject to para-
8	graph (2), eligible items and services are items
9	and services which are physicians' services (as
10	defined in paragraph (4)(A) of section 1848(f)
11	for purposes of calculating the sustainable
12	growth rate under such section).
13	"(2) Secretarial flexibility.—The Secretary
14	shall establish by regulation reasonable limits on the
15	categories of eligible items and services for which a
16	prior determination of coverage may be requested
17	under this subsection. In establishing such limits, the
18	Secretary may consider the dollar amount involved
19	with respect to the item or service, administrative
20	costs and burdens, and other relevant factors.
21	"(3) Request for prior determination.—
22	"(A) In general.—Subject to paragraph
23	(2), under the process established under this sub-
24	section an eligible requester may submit to the

contractor a request for a determination, before



1	the furnishing of an eligible item or service in-
2	volved as to whether the item or service is cov-
3	ered under this title consistent with the applica-
4	ble requirements of section 1862(a)(1)(A) (relat-
5	ing to medical necessity).
6	"(B) Accompanying documentation.—
7	The Secretary may require that the request be
8	accompanied by a description of the item or
9	service, supporting documentation relating to the
10	medical necessity for the item or service, and
11	any other appropriate documentation. In the
12	case of a request submitted by an eligible re-
13	quester who is described in paragraph (1)(B)(ii),
14	the Secretary may require that the request also
15	be accompanied by a copy of the advance bene-
16	ficiary notice involved.
17	"(4) Response to request.—
18	"(A) In general.—Under such process, the
19	contractor shall provide the eligible requester
20	with written notice of a determination as to
21	whether—
22	"(i) the item or service is so covered;
23	"(ii) the item or service is not so cov-
24	$ered;\ or$



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1	"(iii) the contractor lacks sufficient in-
2	formation to make a coverage determina-
3	tion.
4	If the contractor makes the determination de-
5	scribed in clause (iii), the contractor shall in-
6	clude in the notice a description of the addi-
7	tional information required to make the coverage
8	determination.
9	"(B) Deadline to respond.—Such notice
10	shall be provided within the same time period as
11	the time period applicable to the contractor pro-
12	viding notice of initial determinations on a
13	claim for benefits under subsection $(a)(2)(A)$ .
14	"(C) Informing beneficiary in case of
15	PHYSICIAN REQUEST.—In the case of a request
16	in which an eligible requester is not the indi-
17	$vidual\ described\ in\ paragraph\ (1)(B)(ii),\ the$
18	process shall provide that the individual to
19	whom the item or service is proposed to be fur-
20	nished shall be informed of any determination
21	described in clause (ii) (relating to a determina-
22	tion of non-coverage) and the right (referred to
23	in paragraph (6)(B)) to obtain the item or serv-
24	ice and have a claim submitted for the item or



25

service.

1	"(5) Effect of Determinations.—
2	"(A) Binding nature of positive deter-
3	MINATION.—If the contractor makes the deter-
4	mination described in paragraph (4)(A)(i), such
5	determination shall be binding on the contractor
6	in the absence of fraud or evidence of misrepre-
7	sentation of facts presented to the contractor.
8	"(B) Notice and right to redetermina-
9	TION IN CASE OF A DENIAL.—
10	"(i) In General.—If the contractor
11	makes the determination described in para-
12	graph (4)(A)(ii)—
13	"(I) the eligible requester has the
14	right to a redetermination by the con-
15	tractor on the determination that the
16	item or service is not so covered; and
17	"(II) the contractor shall include
18	in notice under paragraph (4)(A) a
19	brief explanation of the basis for the
20	determination, including on what na-
21	tional or local coverage or noncoverage
22	determination (if any) the determina-
23	tion is based, and the right to such a
24	redetermination.



1	"(ii) Deadline for redetermina-
2	TIONS.—The contractor shall complete and
3	provide notice of such redetermination with-
4	in the same time period as the time period
5	applicable to the contractor providing notice
6	of redeterminations relating to a claim for
7	benefits under subsection $(a)(3)(C)(ii)$ .
8	"(6) Limitation on further review.—
9	"(A) In General.—Contractor determina-
10	tions described in paragraph $(4)(A)(ii)$ or
11	(4)(A)(iii) (and redeterminations made under
12	paragraph $(5)(B)$ , relating to pre-service claims
13	are not subject to further administrative appear
14	or judicial review under this section or other-
15	wise.
16	"(B) Decision not to seek prior deter-
17	MINATION OR NEGATIVE DETERMINATION DOES
18	NOT IMPACT RIGHT TO OBTAIN SERVICES, SEEK
19	REIMBURSEMENT, OR APPEAL RIGHTS.—Nothing
20	in this subsection shall be construed as affecting
21	the right of an individual who—
22	"(i) decides not to seek a prior deter-
23	mination under this subsection with respect
24	to items or services; or



1	"(ii) seeks such a determination and
2	has received a determination described in
3	paragraph (4)(A)(ii),
4	from receiving (and submitting a claim for) such
5	items services and from obtaining administrative
6	or judicial review respecting such claim under
7	the other applicable provisions of this section.
8	Failure to seek a prior determination under this
9	subsection with respect to items and services
10	shall not be taken into account in such adminis-
11	trative or judicial review.
12	"(C) No prior determination after re-
13	CEIPT OF SERVICES.—Once an individual is pro-
14	vided items and services, there shall be no prior
15	determination under this subsection with respect
16	to such items or services.".
17	(b) Effective Date; Transition.—
18	(1) Effective date.—The Secretary shall es-
19	tablish the prior determination process under the
20	amendment made by subsection (a) in such a manner
21	as to provide for the acceptance of requests for deter-
22	minations under such process filed not later than 18
23	months after the date of the enactment of this Act.
24	(2) Transition.—During the period in which

the amendment made by subsection (a) has become ef-



1	fective but contracts are not provided under section
2	1874A of the Social Security Act with medicare ad-
3	ministrative contractors, any reference in section
4	1869(g) of such Act (as added by such amendment) to
5	such a contractor is deemed a reference to a fiscal
6	intermediary or carrier with an agreement under sec-
7	tion 1816, or contract under section 1842, respec-
8	tively, of such Act.
9	(3) Limitation on application to sgr.—For
10	purposes of applying section $1848(f)(2)(D)$ of the So-
11	cial Security Act (42 U.S.C. $1395w-4(f)(2)(D)$ ), the
12	amendment made by subsection (a) shall not be con-
13	sidered to be a change in law or regulation.
14	(c) Provisions Relating to Advance Beneficiary
15	Notices; Report on Prior Determination Process.—
16	(1) Data collection.—The Secretary shall es-
17	tablish a process for the collection of information on
18	the instances in which an advance beneficiary notice
19	(as defined in paragraph (5)) has been provided and
20	on instances in which a beneficiary indicates on such
21	a notice that the beneficiary does not intend to seek
22	to have the item or service that is the subject of the
23	$notice\ furnished.$
24	(2) Outreach and education.—The Secretary

shall establish a program of outreach and education



1	for beneficiaries and providers of services and other
2	persons on the appropriate use of advance beneficiary
3	notices and coverage policies under the medicare pro-
4	gram.
5	(3) GAO REPORT REPORT ON USE OF ADVANCE
6	BENEFICIARY NOTICES.—Not later than 18 months
7	after the date on which section 1869(g) of the Social
8	Security Act (as added by subsection (a)) takes effect,
9	the Comptroller General of the United States shall
10	submit to Congress a report on the use of advance
11	beneficiary notices under title XVIII of such Act
12	Such report shall include information concerning the
13	providers of services and other persons that have pro-
14	vided such notices and the response of beneficiaries to
15	such notices.
16	(4) GAO REPORT ON USE OF PRIOR DETERMINA-
17	TION PROCESS.—Not later than 18 months after the
18	date on which section 1869(g) of the Social Security
19	Act (as added by subsection (a)) takes effect, the
20	Comptroller General of the United States shall submit
21	to Congress a report on the use of the prior deter-
22	mination process under such section. Such report
23	shall include—
24	(A) information concerning the types of



1	been sought, determinations made under the
2	process, and changes in receipt of services result-
3	ing from the application of such process; and
4	(B) an evaluation of whether the process
5	was useful for physicians (and other suppliers)
6	and beneficiaries, whether it was timely, and
7	whether the amount of information required was
8	burdensome to physicians and beneficiaries.
9	(5) Advance beneficiary notice defined.—
10	In this subsection, the term "advance beneficiary no-
11	tice" means a written notice provided under section
12	1879(a) of the Social Security Act (42 U.S.C.
13	1395pp(a)) to an individual entitled to benefits under
14	part A or B of title XVIII of such Act before items
15	or services are furnished under such part in cases
16	where a provider of services or other person that
17	would furnish the item or service believes that pay-
18	ment will not be made for some or all of such items

or services under such title.



1	Subtitle V—Miscellaneous
2	Provisions
3	SEC. 941. POLICY DEVELOPMENT REGARDING EVALUATION
4	AND MANAGEMENT (E & M) DOCUMENTATION
5	GUIDELINES.
6	(a) In General.—The Secretary may not implement
7	any new documentation guidelines for, or clinical examples
8	of, evaluation and management physician services under
9	the title XVIII of the Social Security Act on or after the
10	date of the enactment of this Act unless the Secretary—
11	(1) has developed the guidelines in collaboration
12	with practicing physicians (including both generalists
13	and specialists) and provided for an assessment of the
14	proposed guidelines by the physician community;
15	(2) has established a plan that contains specific
16	goals, including a schedule, for improving the use of
17	such guidelines;
18	(3) has conducted appropriate and representative
19	pilot projects under subsection (b) to test modifica-
20	tions to the evaluation and management documenta-
21	tion guidelines;
22	(4) finds that the objectives described in sub-
23	section (c) will be met in the implementation of such
24	guidelines; and



1	(5) has established, and is implementing, a pro-
2	gram to educate physicians on the use of such guide-
3	lines and that includes appropriate outreach.
4	The Secretary shall make changes to the manner in which
5	existing evaluation and management documentation guide-
6	lines are implemented to reduce paperwork burdens on phy-
7	sicians.
8	(b) Pilot Projects to Test Evaluation and Man-
9	AGEMENT DOCUMENTATION GUIDELINES.—
10	(1) In general.—The Secretary shall conduct
11	under this subsection appropriate and representative
12	pilot projects to test new evaluation and management
13	documentation guidelines referred to in subsection
14	(a).
15	(2) Length and consultation.—Each pilot
16	project under this subsection shall—
17	(A) be voluntary;
18	(B) be of sufficient length as determined by
19	the Secretary to allow for preparatory physician
20	and medicare contractor education, analysis,
21	and use and assessment of potential evaluation
22	and management guidelines; and
23	(C) be conducted, in development and
24	throughout the planning and operational stages
25	of the project, in consultation with practicing



1	physicians (including both generalists and spe-
2	cialists).
3	(3) Range of pilot projects.—Of the pilot
4	projects conducted under this subsection—
5	(A) at least one shall focus on a peer review
6	method by physicians (not employed by a medi-
7	care contractor) which evaluates medical record
8	information for claims submitted by physicians
9	identified as statistical outliers relative to defini-
10	tions published in the Current Procedures Ter-
11	minology (CPT) code book of the American Med-
12	$ical\ Association;$
13	(B) at least one shall focus on an alter-
14	native method to detailed guidelines based on
15	physician documentation of face to face encoun-
16	ter time with a patient;
17	(C) at least one shall be conducted for serv-
18	ices furnished in a rural area and at least one
19	for services furnished outside such an area; and
20	(D) at least one shall be conducted in a set-
21	ting where physicians bill under physicians
22	services in teaching settings and at least one
23	shall be conducted in a setting other than a

teaching setting.



1	(4) Banning of targeting of pilot project
2	Participants.—Data collected under this subsection
3	shall not be used as the basis for overpayment de-
4	mands or post-payment audits. Such limitation ap-
5	plies only to claims filed as part of the pilot project
6	and lasts only for the duration of the pilot project
7	and only as long as the provider is a participant in
8	the pilot project.
9	(5) Study of impact.—Each pilot project shall
10	examine the effect of the new evaluation and manage-
11	ment documentation guidelines on—
12	(A) different types of physician practices,
13	including those with fewer than 10 full-time-
14	equivalent employees (including physicians); and
15	(B) the costs of physician compliance, in-
16	cluding education, implementation, auditing,
17	and monitoring.
18	(6) Periodic Reports.—The Secretary shall
19	submit to Congress periodic reports on the pilot
20	projects under this subsection.
21	(c) Objectives for Evaluation and Management
22	Guidelines.—The objectives for modified evaluation and
23	management documentation guidelines developed by the
24	Secretary shall be to—



1	(1) identify clinically relevant documentation
2	needed to code accurately and assess coding levels ac-
3	curately;
4	(2) decrease the level of non-clinically pertinent
5	and burdensome documentation time and content in
6	the physician's medical record;
7	(3) increase accuracy by reviewers; and
8	(4) educate both physicians and reviewers.
9	(d) Study of Simpler, Alternative Systems of
10	Documentation for Physician Claims.—
11	(1) Study.—The Secretary shall carry out a
12	study of the matters described in paragraph (2).
13	(2) Matters described.—The matters referred
14	to in paragraph (1) are—
15	(A) the development of a simpler, alter-
16	native system of requirements for documentation
17	accompanying claims for evaluation and man-
18	agement physician services for which payment is
19	made under title XVIII of the Social Security
20	Act; and
21	(B) consideration of systems other than cur-
22	rent coding and documentation requirements for
23	payment for such physician services.
24	(3) Consultation with practicing physi-
25	CIANS.—In designing and carrying out the study



1	under paragraph (1), the Secretary shall consult with
2	practicing physicians, including physicians who are
3	part of group practices and including both generalists
4	and specialists.
5	(4) Application of Hipaa Uniform coding re-
6	QUIREMENTS.—In developing an alternative system
7	under paragraph (2), the Secretary shall consider re-
8	quirements of administrative simplification under
9	part C of title XI of the Social Security Act.
10	(5) Report to congress.—(A) Not later than
11	October 1, 2005, the Secretary shall submit to Con-
12	gress a report on the results of the study conducted
13	under paragraph (1).
14	(B) The Medicare Payment Advisory Commis-
15	sion shall conduct an analysis of the results of the
16	study included in the report under subparagraph (A)
17	and shall submit a report on such analysis to Con-
18	gress.
19	(e) Study on Appropriate Coding of Certain Ex-
20	TENDED OFFICE VISITS.—The Secretary shall conduct a
21	study of the appropriateness of coding in cases of extended
22	office visits in which there is no diagnosis made. Not later
23	than October 1, 2005, the Secretary shall submit a report
24	to Congress on such study and shall include recommenda-

25 tions on how to code appropriately for such visits in a man-



1	ner that takes into account the amount of time the physi-
2	cian spent with the patient.
3	(f) Definitions.—In this section—
4	(1) the term "rural area" has the meaning given
5	that term in section $1886(d)(2)(D)$ of the Social Secu-
6	$rity\ Act,\ 42\ U.S.C.\ 1395ww(d)(2)(D);\ and$
7	(2) the term "teaching settings" are those set-
8	tings described in section 415.150 of title 42, Code of
9	Federal Regulations.
10	SEC. 942. IMPROVEMENT IN OVERSIGHT OF TECHNOLOGY
11	AND COVERAGE.
12	(a) Council for Technology and Innovation.—
13	Section 1868 (42 U.S.C. 1395ee), as amended by section
14	921(a), is amended by adding at the end the following new
15	subsection:
16	"(c) Council for Technology and Innovation.—
17	"(1) Establishment.—The Secretary shall es-
18	tablish a Council for Technology and Innovation
19	within the Centers for Medicare & Medicaid Services
20	(in this section referred to as 'CMS').
21	"(2) Composition.—The Council shall be com-
22	posed of senior CMS staff and clinicians and shall be
23	chaired by the Executive Coordinator for Technology
24	and Innovation (appointed or designated under para-
25	graph(4)).



1	"(3) Duties.—The Council shall coordinate the
2	activities of coverage, coding, and payment processes
3	under this title with respect to new technologies and
4	procedures, including new drug therapies, and shall
5	coordinate the exchange of information on new tech-
6	nologies between CMS and other entities that make
7	similar decisions.
8	"(4) Executive coordinator for tech-
9	NOLOGY AND INNOVATION.—The Secretary shall ap-
10	point (or designate) a noncareer appointee (as defined
11	in section 3132(a)(7) of title 5, United States Code,
12	who shall serve as the Executive Coordinator for Tech-
13	nology and Innovation. Such executive coordinator
14	shall report to the Administrator of CMS, shall chair
15	the Council, shall oversee the execution of its duties,
16	and shall serve as a single point of contact for outside
17	groups and entities regarding the coverage, coding,
18	and payment processes under this title.".
19	(b) Methods for Determining Payment Basis
20	FOR NEW LAB TESTS.—Section 1833(h) (42 U.S.C.
21	1395l(h)) is amended by adding at the end the following.
22	"(8)(A) The Secretary shall establish by regulation
23	procedures for determining the basis for, and amount of
24	payment under this subsection for any clinical diagnostic

25 laboratory test with respect to which a new or substantially



1	revised HCPCS code is assigned on or after January 1,
2	2005 (in this paragraph referred to as 'new tests').
3	"(B) Determinations under subparagraph (A) shall be
4	made only after the Secretary—
5	"(i) makes available to the public (through an
6	Internet site and other appropriate mechanisms) a
7	list that includes any such test for which establish-
8	ment of a payment amount under this subsection is
9	being considered for a year;
10	"(ii) on the same day such list is made avail-
11	able, causes to have published in the Federal Register
12	notice of a meeting to receive comments and rec-
13	ommendations (and data on which recommendations
14	are based) from the public on the appropriate basis
15	under this subsection for establishing payment
16	amounts for the tests on such list;
17	"(iii) not less than 30 days after publication of
18	such notice convenes a meeting, that includes rep-
19	resentatives of officials of the Centers for Medicare &
20	Medicaid Services involved in determining payment
21	amounts, to receive such comments and recommenda-
22	tions (and data on which the recommendations are
23	based);
24	"(iv) taking into account the comments and rec-

ommendations (and accompanying data) received at



1	such meeting, develops and makes available to the
2	public (through an Internet site and other appro-
3	priate mechanisms) a list of proposed determinations
4	with respect to the appropriate basis for establishing
5	a payment amount under this subsection for each
6	such code, together with an explanation of the reasons
7	for each such determination, the data on which the
8	determinations are based, and a request for public
9	written comments on the proposed determination; and
10	"(v) taking into account the comments received
11	during the public comment period, develops and
12	makes available to the public (through an Internet
13	site and other appropriate mechanisms) a list of final
14	determinations of the payment amounts for such tests
15	under this subsection, together with the rationale for
16	each such determination, the data on which the deter-
17	minations are based, and responses to comments and
18	suggestions received from the public.
19	"(C) Under the procedures established pursuant to sub-
20	paragraph (A), the Secretary shall—
21	"(i) set forth the criteria for making determina-
22	tions under subparagraph (A); and
23	"(ii) make available to the public the data (other
24	than proprietary data) considered in making such de-
25	terminations.



1	"(D) The Secretary may convene such further public
2	meetings to receive public comments on payment amounts
3	for new tests under this subsection as the Secretary deems
4	appropriate.
5	"(E) For purposes of this paragraph:
6	"(i) The term 'HCPCS' refers to the Health Care
7	Procedure Coding System.
8	"(ii) A code shall be considered to be 'substan-
9	tially revised' if there is a substantive change to the
10	definition of the test or procedure to which the code
11	applies (such as a new analyte or a new methodology
12	for measuring an existing analyte-specific test).".
13	(c) GAO STUDY ON IMPROVEMENTS IN EXTERNAL
14	Data Collection for Use in the Medicare Inpatient
15	Payment System.—
16	(1) STUDY.—The Comptroller General of the
17	United States shall conduct a study that analyzes
18	which external data can be collected in a shorter time
19	frame by the Centers for Medicare & Medicaid Serv-
20	ices for use in computing payments for inpatient hos-
21	pital services. The study may include an evaluation
22	of the feasibility and appropriateness of using of
23	quarterly samples or special surveys or any other
24	methods. The study shall include an analysis of
25	whether other executive agencies, such as the Bureau



1	of Labor Statistics in the Department of Commerce,
2	are best suited to collect this information.
3	(2) Report.—By not later than October 1,
4	2004, the Comptroller General shall submit a report
5	to Congress on the study under paragraph (1).
6	(d) Process for Adoption of ICD Codes as Data
7	Standard.—Section 1172(f) (42 U.S.C. 1320d-1(f)) is
8	amended by inserting after the first sentence the following:
9	"Notwithstanding the preceding sentence, if the National
10	Committee on Vital and Health Statistics has not made a
11	recommendation to the Secretary before the date of the en-
12	actment of this sentence, with respect to the adoption of the
13	International Classification of Diseases, 10th Revision,
14	Procedure Coding System (ICD-10-PCS') and the Inter-
15	national Classification of Diseases, 10th Revision, Clinical
16	Modification ('ICD-10-CM') as a standard under this part
17	for the reporting of diagnoses, the Secretary may adopt
18	ICD-10-PCS and ICD-10-CM as such a standard on or
19	after 1 year after such date without receiving such a rec-
20	ommendation.".
21	SEC. 943. TREATMENT OF HOSPITALS FOR CERTAIN SERV-
22	ICES UNDER MEDICARE SECONDARY PAYOR
23	(MSP) PROVISIONS.
24	(a) In General.—The Secretary shall not require a
25	hospital (including a critical access hospital) to ask ques-



- 1 tions (or obtain information) relating to the application of
- 2 section 1862(b) of the Social Security Act (relating to medi-
- 3 care secondary payor provisions) in the case of reference
- 4 laboratory services described in subsection (b), if the Sec-
- 5 retary does not impose such requirement in the case of such
- 6 services furnished by an independent laboratory.
- 7 (b) Reference Laboratory Services De-
- 8 SCRIBED.—Reference laboratory services described in this
- 9 subsection are clinical laboratory diagnostic tests (or the
- 10 interpretation of such tests, or both) furnished without a
- 11 face-to-face encounter between the individual entitled to
- 12 benefits under part A or enrolled under part B, or both,
- 13 and the hospital involved and in which the hospital submits
- 14 a claim only for such test or interpretation.
- 15 SEC. 944. EMTALA IMPROVEMENTS.
- 16 (a) Payment for EMTALA-Mandated Screening
- 17 AND STABILIZATION SERVICES.—
- 18 (1) In General.—Section 1862 (42 U.S.C.
- 19 1395y) is amended by inserting after subsection (c)
- 20 the following new subsection:
- 21 "(d) For purposes of subsection (a)(1)(A), in the case
- 22 of any item or service that is required to be provided pursu-
- 23 ant to section 1867 to an individual who is entitled to bene-
- 24 fits under this title, determinations as to whether the item
- 25 or service is reasonable and necessary shall be made on the



- 1 basis of the information available to the treating physician
- 2 or practitioner (including the patient's presenting symp-
- 3 toms or complaint) at the time the item or service was or-
- 4 dered or furnished by the physician or practitioner (and
- 5 not on the patient's principal diagnosis). When making
- 6 such determinations with respect to such an item or service,
- 7 the Secretary shall not consider the frequency with which
- 8 the item or service was provided to the patient before or
- 9 after the time of the admission or visit.".
- 10 (2) Effective date.—The amendment made by
- 11 paragraph (1) shall apply to items and services fur-
- 12 nished on or after January 1, 2004.
- 13 (b) Notification of Providers When EMTALA In-
- 14 VESTIGATION CLOSED.—Section 1867(d) (42 U.S.C. 42
- 15 U.S.C. 1395dd(d)) is amended by adding at the end the
- 16 following new paragraph:
- 17 "(4) Notice upon closing an investiga-
- 18 Tion.—The Secretary shall establish a procedure to
- 19 notify hospitals and physicians when an investigation
- 20 under this section is closed.".
- 21 (c) Prior Review by Peer Review Organizations
- 22 IN EMTALA CASES INVOLVING TERMINATION OF PARTICI-
- 23 PATION.—
- 24 (1) In General.—Section 1867(d)(3) (42 U.S.C.
- 25 1395dd(d)(3)) is amended—



1	(A) in the first sentence, by inserting "or in
2	terminating a hospital's participation under this
3	title" after "in imposing sanctions under para-
4	graph (1)"; and
5	(B) by adding at the end the following new
6	sentences: "Except in the case in which a delay
7	would jeopardize the health or safety of individ-
8	uals, the Secretary shall also request such a re-
9	view before making a compliance determination
10	as part of the process of terminating a hospital's
11	participation under this title for violations re-
12	lated to the appropriateness of a medical screen-
13	ing examination, stabilizing treatment, or an
14	appropriate transfer as required by this section,
15	and shall provide a period of 5 days for such re-
16	view. The Secretary shall provide a copy of the
17	organization's report to the hospital or physician
18	consistent with confidentiality requirements im-
19	posed on the organization under such part B.".
20	(2) Effective date.—The amendments made
21	by paragraph (1) shall apply to terminations of par-
22	ticipation initiated on or after the date of the enact-
23	ment of this Act.



1	(d) Modification of Requirment for Medical
2	Screening Examinations for Patients Not Request-
3	ING EMERGENCY DEPARTMENT SERVICES.—
4	(1) In general.—Section 1867(a) (42 U.S.C.
5	1395dd(a)) is amended—
6	(A) by designating all that follows "(a)
7	Medical Screening Requirement.—" as
8	paragraph (1) with the heading "In GENERAL.—
9	";
10	(B) by aligning such paragraph with the
11	paragraph added by paragraph (3); and
12	(C) by adding at the end the following new
13	paragraph:
14	"(2) Exception for certain cases.—The re-
15	quirement for an appropriate medical screening ex-
16	amination under paragraph (1) shall not apply in
17	the case of an individual who comes to the emergency
18	department and does not request examination or
19	treatment for an emergency medical condition (such
20	as a request solely for prescription refills, blood pres-
21	sure screening, and non-emergency laboratory and di-
22	agnostic tests).".
23	(2) Effective date.—The amendments made
24	by paragraph (1) shall apply to terminations of par-



1	ticipation initiated on or after the date of the enact-
2	ment of this Act.
3	SEC. 945. EMERGENCY MEDICAL TREATMENT AND ACTIVE
4	LABOR ACT (EMTALA) TECHNICAL ADVISORY
5	GROUP.
6	(a) Establishment.—The Secretary shall establish a
7	Technical Advisory Group (in this section referred to as the
8	"Advisory Group") to review issues related to the Emer-
9	gency Medical Treatment and Labor Act (EMTALA) and
10	its implementation. In this section, the term "EMTALA"
11	refers to the provisions of section 1867 of the Social Security
12	Act (42 U.S.C. 1395dd).
13	(b) Membership.—The Advisory Group shall be com-
14	posed of 19 members, including the Administrator of the
15	Centers for Medicare & Medicaid Services and the Inspector
16	General of the Department of Health and Human Services
17	and of which—
18	(1) 4 shall be representatives of hospitals, includ-
19	ing at least one public hospital, that have experience
20	with the application of EMTALA and at least 2 of
21	which have not been cited for EMTALA violations;
22	(2) 7 shall be practicing physicians drawn from
23	the fields of emergency medicine, cardiology or
24	cardiothoracic surgery, orthopedic surgery, neuro-
25	surgery, pediatrics or a pediatric subspecialty, obstet-



1	rics-gynecology, and psychiatry, with not more than
2	one physician from any particular field;
3	(3) 2 shall represent patients;
4	(4) 2 shall be staff involved in EMTALA inves-
5	tigations from different regional offices of the Centers
6	for Medicare & Medicaid Services; and
7	(5) 1 shall be from a State survey office involved
8	in EMTALA investigations and 1 shall be from a
9	peer review organization, both of whom shall be from
10	areas other than the regions represented under para-
11	graph (4).
12	In selecting members described in paragraphs (1) through
13	(3), the Secretary shall consider qualified individuals nomi-
14	nated by organizations representing providers and patients.
15	(c) General Responsibilities.—The Advisory
16	Group—
17	(1) shall review EMTALA regulations;
18	(2) may provide advice and recommendations to
19	the Secretary with respect to those regulations and
20	their application to hospitals and physicians;
21	(3) shall solicit comments and recommendations
22	from hospitals, physicians, and the public regarding
23	the implementation of such regulations; and



1	(4) may disseminate information on the applica-
2	tion of such regulations to hospitals, physicians, and
3	$the \ public.$
4	(d) Administrative Matters.—
5	(1) Chairperson.—The members of the Advi-
6	sory Group shall elect a member to serve as chair-
7	person of the Advisory Group for the life of the Advi-
8	sory Group.
9	(2) Meetings.—The Advisory Group shall first
10	meet at the direction of the Secretary. The Advisory
11	Group shall then meet twice per year and at such
12	other times as the Advisory Group may provide.
13	(e) Termination.—The Advisory Group shall termi-
14	nate 30 months after the date of its first meeting.
15	(f) Waiver of Administrative Limitation.—The
16	Secretary shall establish the Advisory Group notwith-
17	standing any limitation that may apply to the number of
18	advisory committees that may be established (within the
19	Department of Health and Human Services or otherwise).
20	SEC. 946. AUTHORIZING USE OF ARRANGEMENTS TO PRO-
21	VIDE CORE HOSPICE SERVICES IN CERTAIN
22	CIRCUMSTANCES.
23	(a) In General.—Section 1861(dd)(5) (42 U.S.C.
24	1395x(dd)(5)) is amended by adding at the end the fol-
25	lowing:



- 1 "(D) In extraordinary, exigent, or other non-routine
- 2 circumstances, such as unanticipated periods of high pa-
- 3 tient loads, staffing shortages due to illness or other events,
- 4 or temporary travel of a patient outside a hospice pro-
- 5 gram's service area, a hospice program may enter into ar-
- 6 rangements with another hospice program for the provision
- 7 by that other program of services described in paragraph
- 8 (2)(A)(ii)(I). The provisions of paragraph (2)(A)(ii)(II)
- 9 shall apply with respect to the services provided under such
- 10 arrangements.
- 11 "(E) A hospice program may provide services de-
- 12 scribed in paragraph (1)(A) other than directly by the pro-
- 13 gram if the services are highly specialized services of a reg-
- 14 istered professional nurse and are provided non-routinely
- 15 and so infrequently so that the provision of such services
- 16 directly would be impracticable and prohibitively expen-
- 17 *sive.*".
- 18 (b) Conforming Payment Provision.—Section
- 19 1814(i) (42 U.S.C. 1395f(i)) is amended by adding at the
- 20 end the following new paragraph:
- 21 "(4) In the case of hospice care provided by a hospice
- 22 program under arrangements under section 1861(dd)(5)(D)
- 23 made by another hospice program, the hospice program that
- 24 made the arrangements shall bill and be paid for the hospice
- 25 care.".



1	(c) Effective Date.—The amendments made by this
2	section shall apply to hospice care provided on or after the
3	date of the enactment of this Act.
4	SEC. 947. APPLICATION OF OSHA BLOODBORNE PATHO-
5	GENS STANDARD TO CERTAIN HOSPITALS.
6	(a) In General.—Section 1866 (42 U.S.C. 1395cc)
7	is amended—
8	(1) in subsection $(a)(1)$ —
9	(A) in subparagraph (R), by striking "and"
10	at the end;
11	(B) in subparagraph (S), by striking the
12	period at the end and inserting ", and"; and
13	(C) by inserting after subparagraph (S) the
14	following new subparagraph:
15	"(T) in the case of hospitals that are not other-
16	wise subject to the Occupational Safety and Health
17	Act of 1970, to comply with the Bloodborne Pathogens
18	standard under section 1910.1030 of title 29 of the
19	Code of Federal Regulations (or as subsequently redes-
20	ignated)."; and
21	(2) by adding at the end of subsection (b) the fol-
22	lowing new paragraph:
23	"(4)(A) A hospital that fails to comply with the re-
24	quirement of $subsection$ $(a)(1)(T)$ $(relating$ to the
25	Bloodborne Pathogens standard) is subject to a civil money



- 1 penalty in an amount described in subparagraph (B), but
- 2 is not subject to termination of an agreement under this
- 3 section.
- 4 "(B) The amount referred to in subparagraph (A) is
- 5 an amount that is similar to the amount of civil penalties
- 6 that may be imposed under section 17 of the Occupational
- 7 Safety and Health Act of 1970 for a violation of the
- 8 Bloodborne Pathogens standard referred to in subsection
- 9 (a)(1)(T) by a hospital that is subject to the provisions of
- 10 such Act.
- 11 "(C) A civil money penalty under this paragraph shall
- 12 be imposed and collected in the same manner as civil money
- 13 penalties under subsection (a) of section 1128A are imposed
- 14 and collected under that section.".
- 15 (b) Effective Date.—The amendments made by this
- 16 subsection (a) shall apply to hospitals as of July 1, 2004.
- 17 SEC. 948. BIPA-RELATED TECHNICAL AMENDMENTS AND
- 18 *CORRECTIONS*.
- 19 (a) Technical Amendments Relating to Advisory
- 20 Committee under BIPA Section 522.—(1) Subsection
- 21 (i) of section 1114 (42 U.S.C. 1314)—
- 22 (A) is transferred to section 1862 and added at
- 23 the end of such section; and
- 24 (B) is redesignated as subsection (j).
- 25 (2) Section 1862 (42 U.S.C. 1395y) is amended—



1	(A) in the last sentence of subsection (a), by
2	striking "established under section 1114(f)"; and
3	(B) in subsection (j), as so transferred and
4	redesignated—
5	(i) by striking "under subsection (f)"; and
6	(ii) by striking "section 1862(a)(1)" and
7	inserting "subsection $(a)(1)$ ".
8	(b) Terminology Corrections.—(1) Section
9	1869(c)(3)(I)(ii) (42 U.S.C. 1395ff(c)(3)(I)(ii)), as amend-
10	ed by section 521 of BIPA, is amended—
11	(A) in subclause (III), by striking "policy" and
12	inserting "determination"; and
13	(B) in subclause (IV), by striking "medical re-
14	view policies" and inserting "coverage determina-
15	tions".
16	(2) Section $1852(a)(2)(C)$ (42 U.S.C. $1395w$ -
17	22(a)(2)(C)) is amended by striking "policy" and "POLICY"
18	and inserting "determination" each place it appears and
19	"DETERMINATION", respectively.
20	(c) Reference Corrections.—Section 1869(f)(4)
21	(42 U.S.C. 1395ff(f)(4)), as added by section 522 of BIPA,
22	is amended—
23	(1) in subparagraph (A)(iv), by striking "sub-
24	clause (I), (II), or (III)" and inserting "clause (i),
25	(ii), or (iii)":



1	(2) in subparagraph (B), by striking "clause
2	(i)(IV)" and "clause (i)(III)" and inserting "sub-
3	paragraph (A)(iv)" and "subparagraph (A)(iii)", re-
4	spectively; and
5	(3) in subparagraph (C), by striking "clause
6	(i)", "subclause (IV)" and "subparagraph (A)" and
7	inserting "subparagraph (A)", "clause (iv)" and
8	"paragraph (1)(A)", respectively each place it ap-
9	pears.
10	(d) Other Corrections.—Effective as if included in
11	the enactment of section 521(c) of BIPA, section 1154(e)
12	(42 U.S.C. 1320c-3(e)) is amended by striking paragraph
13	(5).
14	(e) Effective Date.—Except as otherwise provided,
15	the amendments made by this section shall be effective as
16	if included in the enactment of BIPA.
17	SEC. 949. CONFORMING AUTHORITY TO WAIVE A PROGRAM
18	EXCLUSION.
19	The first sentence of section $1128(c)(3)(B)$ (42 U.S.C.
20	1320a-7(c)(3)(B)) is amended to read as follows: "Subject
21	to subparagraph (G), in the case of an exclusion under sub-
22	section (a), the minimum period of exclusion shall be not
23	less than five years, except that, upon the request of the ad-
24	ministrator of a Federal health care program (as defined
25	in section 1128B(f)) who determines that the exclusion



- 1 would impose a hardship on individuals entitled to benefits
- 2 under part A of title XVIII or enrolled under part B of
- 3 such title, or both, the Secretary may waive the exclusion
- 4 under subsection (a)(1), (a)(3), or (a)(4) with respect to
- 5 that program in the case of an individual or entity that
- 6 is the sole community physician or sole source of essential
- 7 specialized services in a community.".
- 8 SEC. 950. TREATMENT OF CERTAIN DENTAL CLAIMS.
- 9 (a) In General.—Section 1862 (42 U.S.C. 1395y) is
- 10 amended by adding after subsection (g) the following new
- 11 subsection:
- 12 "(h)(1) Subject to paragraph (2), a group health plan
- 13 (as defined in subsection (a)(1)(A)(v)) providing supple-
- 14 mental or secondary coverage to individuals also entitled
- 15 to services under this title shall not require a medicare
- 16 claims determination under this title for dental benefits spe-
- 17 cifically excluded under subsection (a)(12) as a condition
- 18 of making a claims determination for such benefits under
- 19 the group health plan.
- 20 "(2) A group health plan may require a claims deter-
- 21 mination under this title in cases involving or appearing
- 22 to involve inpatient dental hospital services or dental serv-
- 23 ices expressly covered under this title pursuant to actions
- 24 taken by the Secretary.".



- 1 (b) Effective Date.—The amendment made by sub-
- 2 section (a) shall take effect on the date that is 60 days after
- 3 the date of the enactment of this Act.
- 4 SEC. 951. FURNISHING HOSPITALS WITH INFORMATION TO
- 5 COMPUTE DSH FORMULA.
- 6 Beginning not later than 1 year after the date of the
- 7 enactment of this Act, the Secretary shall furnish to sub-
- 8 section (d) hospitals (as defined in section 1886(d)(1)(B)
- 9 of the Social Security Act, 42 U.S.C. 1395ww(d)(1)(B)) the
- 10 data necessary for such hospitals to compute the number
- 11 of patient days described in subclause (II) of section
- 12 1886(d)(5)(F)(vi) of the Social Security Act (42 U.S.C.
- 13 1395ww(d)(5)(F)(vi)) used in computing the dispropor-
- 14 tionate patient percentage under such section for that hos-
- 15 pital. Such data shall also be furnished to other hospitals
- 16 which would qualify for additional payments under part
- 17 A of title XVIII of the Social Security Act on the basis of
- 18 such data.
- 19 SEC. 952. REVISIONS TO REASSIGNMENT PROVISIONS.
- 20 (a) IN GENERAL.—Section 1842(b)(6)(A) (42 U.S.C.
- 21 1395u(b)(6)(A)) is amended by striking "or (ii) (where the
- 22 service was provided in a hospital, critical access hospital,
- 23 clinic, or other facility) to the facility in which the service
- 24 was provided if there is a contractual arrangement between
- 25 such physician or other person and such facility under



- 3 4 5 6 8 9 10 13 14 15 16 17 18 TION.
  - which such facility submits the bill for such service," and
  - inserting "or (ii) where the service was provided under a
  - contractual arrangement between such physician or other
  - person and an entity (as defined by the Secretary), to the
  - entity if, under the contractual arrangement, the entity sub-
  - mits the bill for the service and the contractual arrangement
  - meets such other program integrity and other safeguards
  - as the Secretary may determine to be appropriate,".
  - (b) Conforming Amendment.—The second sentence
- of section 1842(b)(6) (42 U.S.C. 1395u(b)(6)) is amended
- by striking "except to an employer or facility" and insert-
- ing "except to an employer, entity, or other person".
- (c) Effective Date.—The amendments made by sec-
- tion shall apply to payments made on or after the date that
- is one year after the date of the enactment of this Act.
- SEC. 953. OTHER PROVISIONS.
- (a) GAO REPORTS ON THE PHYSICIAN COMPENSA-
- 19 Sustainable Growth Rate and
- 20 DATES.—Not later than 6 months after the date of the
- 21 enactment of this Act, the Comptroller General of the
- 22 United States shall submit to Congress a report on
- 23 the appropriateness of the updates in the conversion
- 24 factor under subsection (d)(3) of section 1848 of the
- 25 Social Security Act (42 U.S.C. 1395w-4), including



1	the appropriateness of the sustainable growth rate for-
2	mula under subsection (f) of such section for 2002
3	and succeeding years. Such report shall examine the
4	stability and predictability of such updates and rate
5	and alternatives for the use of such rate in the up-
6	dates.
7	(2) Physician compensation generally.—Not
8	later than 12 months after the date of the enactment
9	of this Act, the Comptroller General shall submit to
10	Congress a report on all aspects of physician com-
11	pensation for services furnished under title XVIII of
12	the Social Security Act, and how those aspects inter-
13	act and the effect on appropriate compensation for
14	physician services. Such report shall review alter-
15	natives for the physician fee schedule under section
16	1848 of such title (42 U.S.C. 1395w-4).
17	(b) Annual Publication of List of National Cov-
18	ERAGE DETERMINATIONS.—The Secretary shall provide, in
19	an appropriate annual publication available to the public,
20	a list of national coverage determinations made under title
21	XVIII of the Social Security Act in the previous year and
22	information on how to get more information with respect
23	to such determinations.
24	(c) GAO REPORT ON FLEXIBILITY IN APPLYING HOME

25 Health Conditions of Participation to Patients



1	Who are Not Medicare Beneficiaries.—Not later than
2	6 months after the date of the enactment of this Act, the
3	Comptroller General of the United States shall submit to
4	Congress a report on the implications if there were flexi-
5	bility in the application of the medicare conditions of par-
6	ticipation for home health agencies with respect to groups
7	or types of patients who are not medicare beneficiaries. The
8	report shall include an analysis of the potential impact of
9	such flexible application on clinical operations and the re-
10	cipients of such services and an analysis of methods for
11	monitoring the quality of care provided to such recipients.
12	(d) OIG REPORT ON NOTICES RELATING TO USE OF
13	Hospital Lifetime Reserve Days.—Not later than 1
14	year after the date of the enactment of this Act, the Inspec-
15	tor General of the Department of Health and Human Serv-
16	ices shall submit a report to Congress on—
17	(1) the extent to which hospitals provide notice
18	to medicare beneficiaries in accordance with applica-
19	ble requirements before they use the 60 lifetime reserve
20	days described in section 1812(a)(1) of the Social Se-
21	$curity \ Act \ (42 \ U.S.C. \ 1395d(a)(1)); \ and$
22	(2) the appropriateness and feasibility of hos-
23	pitals providing a notice to such beneficiaries before

they completely exhaust such lifetime reserve days.



1	SEC. 954. TEMPORARY SUSPENSION OF OASIS REQUIRE-
2	MENT FOR COLLECTION OF DATA ON NON-
3	MEDICARE AND NON-MEDICAID PATIENTS.
4	(a) In General.—During the period described in sub-
5	section (b), the Secretary may not require, under section
6	4602(e) of the Balanced Budget Act of 1997 or otherwise
7	under OASIS, a home health agency to gather or submit
8	information that relates to an individual who is not eligible
9	for benefits under either title XVIII or title XIX of the So-
10	cial Security Act (such information in this section referred
11	to as "non-medicare/medicaid OASIS information").
12	(b) Period of Suspension.—The period described in
13	this subsection—
14	(1) begins on the date of the enactment of this
15	Act; and
16	(2) ends on the last day of the 2nd month begin-
17	ning after the date as of which the Secretary has pub-
18	lished final regulations regarding the collection and
19	use by the Centers for Medicare & Medicaid Services
20	of non-medicare/medicaid OASIS information fol-
21	lowing the submission of the report required under
22	subsection (c).
23	(c) Report.—
24	(1) Study.—The Secretary shall conduct a study
25	on how non-medicare/medicaid OASIS information is



1	and can be used by large home health agencies. Such
2	study shall examine—
3	(A) whether there are unique benefits from
4	the analysis of such information that cannot be
5	derived from other information available to, or
6	collected by, such agencies; and
7	(B) the value of collecting such information
8	by small home health agencies compared to the
9	administrative burden related to such collection.
10	In conducting the study the Secretary shall obtain
11	recommendations from quality assessment experts in
12	the use of such information and the necessity of small,
13	as well as large, home health agencies collecting such
14	information.
15	(2) Report.—The Secretary shall submit to
16	Congress a report on the study conducted under para-
17	graph (1) by not later than 18 months after the date
18	of the enactment of this Act.
19	(d) Construction.—Nothing in this section shall be
20	construed as preventing home health agencies from col-
21	lecting non-medicare/medicaid OASIS information for
22	their own use.

