voluntary to mandatory. The change from voluntary to mandatory is necessary to increase the reliability of the data and to make it available on a more timely basis. Section 2605(b)(14) of the Low Income Home Energy Assistance Act, as amended, requires grantees to provide assurance that they will cooperate with the Secretary with respect to data collecting and reporting. This is one of 16 assurances a State's governor or someone specifically designated by the governor, makes as part of each year's LIHEAP application.

To be in full compliance with section 2605(b)(14), grantees must return the completed survey by the due date.

The preliminary estimates collected by the Grantee Survey for the current fiscal year are needed to provide the Administration and Congress with fiscal and case load estimates in time for hearings about LIHEAP appropriations and program performance. Final estimates for the previous fiscal year will be included in the Department's annual LIHEAP Report to Congress and will be posted on the Department's LIHEAP web site for access by grantees and other interested parties.

Respondents: 50 States and the District of Columbia.

Annual Burden Estimates

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Survey	51	1	3.5	178.5
Estimated Total Annual Burden Hours:				178.5

In compliance with the requirements of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: April 4, 2001.

Bob Sargis,

Reports Clearance Officer.

[FR Doc. 01–8765 Filed 4–9–01; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98N–0787]

Parke-Davis Pharmaceutical Research et al.; Withdrawal of Approval of 14 New Drug Applications and 13 Abbreviated New Drug Applications; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of September 25, 1998 (63 FR 51359). The document announced the withdrawal of approval of 14 new drug applications and 13 abbreviated new drug applications (ANDAs). The document inadvertently withdrew approval of ANDA 80–025 for Sulf-10 (sulfacetamide sodium ophthalmic solution, USP) 10% held by Ciba Vision, 11460 Johns Creek Pkwy., Duluth, GA 30097–1556. FDA confirms that approval of ANDA 80–025 is still in effect.

EFFECTIVE DATE: September 25, 1998.

FOR FURTHER INFORMATION CONTACT: Olivia A. Pritzlaff, Center for Drug Evaluation and Research (HFD–7), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–594–2041.

In FR Doc. 98–25713 appearing on page 51359 in the issue of Friday, September 25, 1998, the following correction is made: On page 51360, in the table, the entry for ANDA 80–025 is removed.

Dated: March 26, 2001.

Janet Woodcock,

Director, Center for Drug Evaluation and Research.

[FR Doc. 01–8718 Filed 4–9–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[Document Identifier: HCFA–R–0209 and HCFA–R–0245]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Health Care Financing Administration, DHHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

1. *Type of Information Collection Request:* Extension of a currently