۹	REVIEW GROUP	TYPE	PROGRAM		T NUMBER (IN	ERT	
DEPARTMENT OF		5	ROI	}	PAGES) 4650-06		
HEALTH, EDUCATION, AND WELFARE	IMB		RUI	GPII	4050-08		
PUBLIC HEALTH SERVICE							
	FROM: 12/01/	/69		UGH: 1	1/30/74		
APPLICATION	REQUESTED BUDGET PERIOD						
FOR CONTINUATION GRANT	12.00				1/20/22		
TO BE VERIFIED BY APPLICANT. CHECK INFORMATION IN ITEMS 1 TH	FROM: 12/01/	T, FURNISH			1/30/72 N IN ITEM 13.		
1. TITLE		<del></del> .					
GENETICS OF HUMAN TISSUE AXX	NXXXXX ANTIGE	NS					
24. PRINCIPAL INVESTIGATOR OR PROGRAM DIRECTOR (Name and Address, Street, City, State, Zip Code)	4. APPLICANT OF State, Zip Code)	RGANIZAT	ION (Name	ənd Addı	ress, Street, Cit	у,	
LEDERBERG, JOSHUA	STANFORD	UNIV	ERSITY	,			
DEPT OF GENETICS	STANFORD	), CAL	IFORNI	A 9	4305		
STANFORD UNIVERSITY SCH OF MED							
STANFORD, CALIF 94305 28. DEGREE 12C. SOCIAL SECURITY NO.	5. PHS ACCOUNT	NUMBER					
Ph.D.			45821	0			
20. DEPARTMENT, SERVICE, LABORATORY OR EQUIVALENT	6. TITLE AND AD	DRESS O			SINESS OFFICE		
	OF APPLICAN						
GENETICS							
2E. MAJOR SUBDIVISION	CONTROLLER						
SCHOOL OF MEDICINE	STANFORD UNIVERSITY STANFORD, CALIFORNIA 94305						
3. ORGANIZATIONAL COMPONENT TO RECEIVE CREDIT FOR INSTITUTIONAL GRANT PURPOSES	- STANFURL	I LAL	I FUKNI	A 7	4302		
01 SCHUDL OF MEDICINE							
COMPLETE THE FO							
7. RESEARCH (NVOLVING HUMAN SUBJECTS (See Instructions)	INVENTION CERTIFICATION (See Instructions)					Y	
NO YES APPROVED: DATE	- YES-PREVIO	USLY REPO	RTED		PORTED		
9. PERFORMANCE SITE (S)		·	HONE INFO	RMATIO	N		
	11A. PRINCIPAL I	VESTIGA	TOR	AREA	TELE. NO. &	EXT.	
Genetics Department		R	17514 3 41				
Stanford University School of Medicine Stanford, California	PROGRAM DI			415	321-1200	<u>580</u>	
	118. NAME OF BU. (ITEM 6)	SINESS O	FFICIAL				
	K. D. Crei	ighton		415	321-2300	2251	
	TIC. NAME AND T			• 			
10. DIRECT COSTS REQUESTED FOR BUDGET PERIOD	-4						
\$14,439	128. COUNTY OF	APPLICA	NT ORGANI	ZATION	SHOWN IN ITE	Nº d	
12A. CONGRESSIONAL DISTRICT OF APPLICANT ORGANIZATION SHOWN IN ITEM 4 Tenth	Santa C						
I USE THIS SPACE FOR CORRECTIONS TO ITEMS I THROUGH 6			WHERE AN	SWER(S)	APPLY		
Note correction of spelling in title, I	tem 1. Item	2B - a	add degr	ee.			
Item 2C - Dr. Lederberg's Soc. Sec. No.			5				

. .

14. CERTIFICATION AND ACCEPTANCE. WE THE UNDERSIGNED, CERTIFY THAT THE STATEMENTS HEREIN ARE TRUE AND COMPLETE TO THE BEST OF OUR KNOWLEDGE AND ACCEPT AS TO ANY GRANT AWARDED. THE DELIGATION TO COMPLY WITH PUBLIC HEALTH SERVICE TERMS AND CONDITIONS IN EFFECT AT THE TIME OF THE AWARD.

SIGNATURES	154. PRINCIPAL INVESTIGATOR OR PROGRAM DIRECTOR	DATE .
(Signatures required of original conv only. Us		8/16/71
ink "Per" signature: not acceptable.)		DATE
FH3 2590-1 OPTIONAL REV. 1-70	RETHEN COMPLETED APPLICATION TO DHS REFORES 1	

RETURN COMPLETED APPLICATION TO PHS BEFORE: 1 OCTOBER 1971

			SECTION	11				<b>F</b> ()
SECTION II-BI		12 MONTHS)	FROM	1/71	THROUGH		NT NUMB	
A. ITEMIZE DIRECT COS			12/2	L//L	11/30/7	<u> </u>	<u>1 146</u>	50-06
PERSONNEL				TIME OR EFFORT %/HRS.	SALARY REQUESTED	FRINGE BI (See Instr	NEFITS actions)	TOTAL
NAME (Last, First, Initial) (a)		TITLE OF POSITION (b)		%/HRS. (C)	(d)	(e)		(f)
Lederberg, J.		PRINCIPAL INVESTIGATOR			none			
Hwang, J.		Scientific Prog	gramme		3,600	561		
Wang, L,		Sr. Res. Asst.		_16%	2,000	467		
Leo, M.		Computer Operat	tor	35%	3,000			
Staff benefit	s:							
9 mo. @ 15.2%								
3 mo.,@ 1.6.7%								
		S	ubtotals —	>	\$ 8,600	\$ 1,339	)	
(Indicate cost of eacl	h item listed below)		TOTAL.	(Columns (d)			>	\$ 2.000
CONSULTANT COSTS (See I	Instructions)							<u>9,939</u>
CONSOLIANI COSIS (SEE	instructions)							\$
EQUIPMENT								
				······				
·								
SUPPLIES Cultur	<u>e medium anc</u>	<u>chemicals</u>				300	0	
		<u></u>						
								\$ 300
	DOMESTIC							\$
TRAVEL	FOREIGN							\$
PATIENT COSTS (See instr	uctions)							
ALTERATIONS AND RENOVA	TIONS		·····			······································		\$
		·····						S
OTHER EXPENSES (Itemize	)							
Computer Time		\$4,000						
Communication	ns, Publicat:							
Costs		200						\$ 4,200
TOTAL DIRECT COST (Ente	er on Page 1, Item 10)							\$14,439
INDIRECT		% S&W*		Date of DHEV	Y Agreement:		] Not Rec	· · · · · · · · · · · · · · · · · · ·
COST		"NTDC"	2/2-	4/71				negotiation with.
(See Instructions)	"If this is a spe	cial rate (e.g. off-site), explai	n		·		<u> </u>	

PHS 2590-1 (All previous editions obsolete) Rev. 1-70

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PAGE 2

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#### SECTION II—BUDGET (Continued)

Grant Number

GM-14650-06

#### JDGET (Continued)

B. Supplemental information regarding ITEMS in the proposed budget for the next period which require explanation or justification. (See Instructions)

The major emphasis of the work supported will be in connection with the continuing computer analysis of our serological data on the HL-A system. Mrs. Hwang has been largely responsible for the development of our system of programs and continues, with the help of Mrs. Leo, to supervise the analysis of our data. During the coming year we will be obtaining data on the frequencies of HL-A antigens in a variety of populations, and also different disease states, specifically Hodgkin's disease and systemic lupus erythematosus. The experimental work is being done in collaboration with Drs. McDevitt and Grumet of the Department of Medicine. In addition, our system of programs is used by Dr. Rose Payne of the Department of Medicine, who was closely associated with Dr. Bodmer when he was at Stanford.

A small amount of experimental work connected with the correlation of HL-A typing on fibroblasts and lymphocytes is being carried on by Mrs. Wang. This represents a continuation of the studies started by Dr. Bodmer when he was at Stanford.

Because the staff benefit rate has increased over that previously anticipated, total direct costs are slightly more than the previously recommended amount.

SECTION III							
SECTION III—FISCAL DATA FOR CURRENT BUDGET PERIOD (USUALLY 12 MONTHS)	FROM 1.2/1/71	тнгоидн 11/30/72	GRANT NUMBER GM-14650-06				

The following pertains to your CURRENT PHS budget. Do not include cost sharing funds. This information in conjunction with that provided on Page 2 will be used in determining the amount of support for the NEXT budget period.

<u>,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,</u>	A. BUDGET CATEGORIES	CURRENT BUDGET (As approved by awarding unit) (1)	ACTUAL EXPENDITURES THRU 7/31/71 (Insert Date) (2)	ESTIMATED ADDITIONAL EXPENDITURES AND OBLIGATIONS FOR REMAINDER OF CURRENT BUDGET PERIOD (3)	TOTAL ESTIMATED EXPENDITURES AND OBLIGATIONS (Col. 2 plus Col. 3) (4)	ESTIMATED UNOBLIGATED BALANCE (Subtract Col. 4 from Col. 1) (5)
Personnel	(Salaries)	13,890	7,338	6,552	13,890	0
Fringe Ber	nefits .	1,952	1,020	955	1,975	-23
Consultant	Costs					
Equipment	t					
Supplies		3,000	2,400	900	3,300	-300
	Domestic					
TRAVEL	Foreign					
Patient Co	sts	1,000	200	0	200	800
Alterations	s and Renovations					
Other		12,100	6,335	6,242	12,577	-477
Total Dire	ct Costs	31,942	17,293	14,649	31,942	0
Indirect C	osts (If included in award)	8,195	4,329	3,866	8,195	0
TOTALS	······································	\$40,137	\$21,622	\$18,515	\$40,137	\$ 0

Use space below to:

B. List all items of equipment purchased or expected to be purchased during this budget period which have a unit cost of \$1000 or more.

C. Explain any significant balance or deficit shown in any category of Column 5.

D. List all other research support for Principal Investigator by source, project title, and annual amount.

The taking of blood and tissue samples has for the most part been completed, so there will be a balance left in the "Patient Cost" allocation. However, this is more than offset by costs related to the analysis of the samples and computerization of the data.

Research Support:

(NIH) AI-5160 Genetics of Bacteria

\$56,000 p.a.

#### SECTION IV

APPLICANT: REPEAT GRANT NUMBER SHOWN ON PAGE 1	GRANT NUMBER	
SECTION IV-SUMMARY PROGRESS REPORT	GM-14650-	06
PRINCIPAL INVESTIGATOR OR PROGRAM DIRECTOR (Last, First, Initial)	PERIOD COVERED BY THIS REP	
Lederberg, Joshua	FROM	THROUGH
NAME OF ORGANIZATION	10/1/71	11/00/70
Stanford University	12/1/71	11/30/72
FITLE (Repeat title shown in Item 1 on first page)		
Constine of Human Tierre Anticone		

Genetics of Human Tissue Antigens

1. List publications: (a) published and not previously reported; (b) in press, Provide five reprints if not previously submitted.

2. List all additions and deletions in professional personnel and any changes in effort.

3. Progress Report. (See Instructions)

## 1. Publications.

- Santachiara, A.S., M. Nabholz, V. Miggiano, A. J. Darlington and W. F. Bodmer, 1970. Genetic analysis with man-mouse hybrids: linkage between human lactate dehydrogenase B and peptidase B. Nature 227: 248-251.
- Bodmer, J.G. and W. F. Bodmer, 1970. Studies on African Pygmies. IV: A comparative study of the HL-A polymorphism in the Babinga Pygmies and other African and Caucasian populations. Am. Journ. Hum. Genet. 22: 396-411.
- Miggiano, V.C., M. Nabholz and W. F. Bodmer, 1970. Detection of HL-A and other antigens on fibroblast micro-monolayers using a fluorochromatic cytotoxicity assay. Histocompatibility Testing, 1970, 623-629.
- Gabb, B.W. and W. F. Bodmer, 1970. A micro complement fixation test for platelet antibodies. Histocompatibility Testing, 1970, 543-547.
- Bodmer, J., A. Coukell, W. F. Bodmer, R. Payne and E. Shanbrom, 1970. A new allele for the LA series of HL-A antigens: the analysis of a complex serum. Histocompatibility Testing, 1970, 175-185.
- Bodmer, W. F., J. G. Bodmer and M. Tripp, 1970. Recombination between the LA and 4 loci of the HL-A system. Histocompatibility Testing 1970, 187-191.
- Mattiuz, P. L., D. Ihde, A. Piazza, R. Ceppellini and W. F. Bodmer, 1970. New approaches to the population genetic and segregation analysis of the HL-A system. Histocompatibility Testing 1970, 193-205.
- Hulett, R., A. Coukell and W. F. Bodmer, 1970. Tissue typing instrumentation using the fluorochromatic cytocoxicity assay. Transplantation 10: 135-137.
- Payne, R., J. Bodmer, W. F. Bodmer and E. Shanbrom, 1970. Production of defined human leukocyte typing sera. Histocompatibility Testing, 1970, 207-220.
- Coukell, A., J. G. Bodmer and W. F. Bodmer, 1971. HL-A types of forty-four Hodgkins patients. Transplantation Proc. (in press)
- McDevitt, H.O. and W. F. Bodmer, 1971. Histocompatibility antigens, immune responsiveness and susceptibility to disease. American Journal of Med. (in press)
- Grumet, F.C., A. Coukell, J. G. Bodmer, W. F. Bodmer and H.O. McDevitt, 1971. Histocompatibility antigens associated with systemic lupus erythematosis: A possible genetic predisposition to disease. New England J. Med. (in press)

2. Per budget.

# 3. Progress Report (GM-14650-05)

During the current grant year, the major part of Dr. Bodmer's activities was transferred to the University of Oxford where he has taken up a position as Professor of Genetics. Experimental work was continued at Stanford by Mrs. Anne Coukell, and the Stanford Medical School's ACME computer facility continued to be used for our data analysis, while Professor Lederberg took over as principal investigator on the project from January 1, 1971.

A major emphasis of our work during the year has been on the association between HL-A and diseases, specifically lupus erythematosis and Hodgkins disease. In collaboration with Drs. McDevitt and Grumet, we have confirmed the very significant increase in the frequency of the antigen W15 in patients with lupus erythematosis. There were some puzzling anomalies in the typing of these patients which will be followed up by family studies to confirm their antigen phenotype and further serological studies on the nature of the autolymphocytotoxic antibody present in the sera of many of these patients.

Typing of forty-four Hodgkins patients from Dr. Henry Kaplan's clinic at Stanford did not indicate the previously reported increase in the antigens W5 or HL-A5 in these patients. However, it appears likely that this may be because the distribution of types of Hodgkins is different in the patients that we typed. Specifically, these seemed to include a much higher frequency of the nodular sclerosing type of the disease than is normally found in other series.

We are again participating in the next International Histocompatibility Testing Workshop, whose aim is to obtain as comprehensive information as possible on the distribution of the HL-A antigens in different populations. We took part in the testing of sera to be used by the participants in this Workshop and prepared a specially absorbed serum for the detection of a component of one of the newer antigens of the LA series.

We have greatly simplified our procedures for collecting, storing and shipping lymphocytes for typing so that they now require a minimum of processing in the field. This has enabled us to collaborate much more easily with workers in out-of-the-way places in order to obtain blood samples for HL-A typing. Prepared for the Science Information Exchange.

Not for publication or publication reference.

U. S. Department of HEALTH, EDUCATION, AND WELFARE PUBLIC HEALTH SERVICE PROJECT NO. (DO NOT USE THIS SPACE)

# NOTICE OF RESEARCH PROJECT

## TITLE OF PROJECT

Genetics of Human Tissue Antigens

#### GIVE NAMES, DEPARTMENTS, AND OFFICIAL TITLES OF PRINCIPAL INVESTIGATORS OR PROJECT DIRECTORS AND ALL OTHER PROFESSIONAL PERSONNEL ENGAGED ON THE PROJECT.

NAME AND ADDRESS OF APPLICANT INSTITUTION

Stanford University, Stanford, California 94305

SUMMARY OF PROPOSED WORK – (200 words or less – Omit Confidential data.) In the Science Information Exchange summaries of work in progress are exchanged with government and private agencies supporting research in the bio-sciences and are forwarded to investigators who request such information. Your summary is to be used for these purposes.

The main aim of this research program is to further the understanding of the inheritance of antigenic differences of human leukocytes and other human tissues and to use these antigens for studies in somatic cell genetics. Cytotoxicity assays are being used together with intensive absorption studies of sera reacting with human leukocytes for the investigation of the genetics of the major human leukocyte antigen polymorphisms. Studies on the distribution of these antigens in various racial groups will also be undertaken. The specificities of antigens carried by permanent and primary cell culture lines are being investigated. The use of these antigens for studies in somatic cell genetics are being explored.

PROFESSIONAL SCHOOL (medical, dental, etc. WHICH THIS PROJECT SHOULD BE IDENTIFI	.) WITH SIGNATURE OF PRINCIPAL INVES	TIGATOR	DATE			
School of Medicine	former so show	Minine to shark				
DO N	OT WRITE BELOW THIS LINE - FOR OFFICE USE C	INLY /				
SUPPORTING AGENCY		1				
METHOD OF SUPPORT (Check one) Agency Staff (Intramural) Negotiated Contract	Special Research Project Grant Grant	Other (Specify)				
	BER OF FUTURE YEARS TENTATIVELY ASSURED DND CURRENT FISCAL YEAR	BEGINNING DATE	ESTIMATED COMPLETION DATE			