## GEORGIA INSTITUTE OF TECHNOLOGY OFFICE OF CONTRACT ADMINISTRATION SPONSORED PROJECT INITIATION

Date: September 9, 1977

Project Title: Equipment for Instrumentation and Experimentation in Heart-Muscle Mechanics

Project No: E-23-631

Project Director: Dr. Hyland Y.-L. Chen

Sponsor: National Science Foundation

Dr ch

Agreement Period: From 9/1/77 Until 2/28/79 (12-month budget period plus 6-month flexibility period)

Type Agreement: Grant No. ENG77-17233

Amount: \$19,300 NSF 19,248 GIT (E-23-330) \$38,548

Reports Required: Final Technical Report; Summary of Completed Project

Sponsor Contact Person (s):

Technical Matters

Contractual Matters (thru OCA)

Division of Engineering National Science Foundation Washington, D. C. 20550

### Ms. Mary Frances O'Connell Grants Specialist, Area 4 Division of Grants and Contracts National Science Foundation Washington, D. C. 20550 (202) 632-2858

#### Defense Priority Rating: N/A

# Assigned to: <u>Engineering Science and Mechanics</u> (School/Laboratory) COPIES TO:

Project Director Division Chief (EES) School/Laboratory Director Dean/Director—EES Accounting Office Procurement Office Security Coordinator (OCA) Reports Coordinator (OCA) Library, Technical Reports Section Office of Computing Services Director, Physical Plant EES Information Office Project File (OCA) Project Code (GTRI) Other

CA-3 (3/76)

## **GEORGIA INSTITUTE OF TECHNOLOGY** OFFICE OF CONTRACT ADMINISTRATION

#### SPONSORED PROJECT TERMINATION

Date: 1/29/81

Project Title: Equipment for Instrumentation & Experimentation in Heart-Muscle Mechanics

Project No: E-23-631

Project Director: Dr. Hyland Y.-L. Chen

Sponsor: NSF

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Effective Termination Date: \_\_\_\_\_2/28/79

Clearance of Accounting Charges: \_\_\_\_\_

Grant/Contract Closeout Actions Remaining:

Final Invoice and Closing Documents

\_x Final Fiscal Repair FCTR (Accounting)

X Final Report of Inventions (if positive)

Govt. Property Inventory & Related Certificate

\_ Classified Material Certificate

\_\_\_ Other \_\_\_\_\_

# Assigned to: \_\_\_\_\_ Engineering Science & Mechanics

(School/Kattory)

#### COPIES TO:

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NATIONAL SCIENCE FOUNDATION			<u> </u>		; , , , , , , , , , , , , , , , , ,
Washington, D.C. 20550	L PROJECT F	A			
PLEASE READ INSTRUCT					
PART I-PROJEC	2. NSF Program				
1. Institution and Address	Engineer			SF Award Nun	iber
Georgia Institute of Technology Atlanta, Georgia 30332	4. Award Period			<u>377-17233</u> umulative Awa	, A mount
	From 9/1/			\$19,300	
6. Project Title	<u> </u>				
"Equipment for Instrumentation and E	xperimentat	ion in He	art-Muscle	e Mechanio	cs".
PART II-SUMMARY OF	COMPLETED PR	OJECT (FOR P	UBLIC USE)		<u>_</u>
An experimental system for the s of heart muscle is designed and const The system consists of four majo optical and control systems. The env PO <sub>2</sub> , temperature and programmed elect the rapidly and continuously changing looped, electro-mechanical system is microseconds. (Comparable to the hux The optic system consists of a 1 without artifact introduced by damage activity. The control system consist used to monitor and control the mover and stimulation; it also monitors pO, digital processing units. As a by-product, an optic probe	for subsyster vironmental cric stimul g active st designed w ley's of 40 laser and a es ends, an ts of a min ment of the displace for the vi	ms, i.e., system pr ation of t ate of hea ith a stea 0 microsec ccessories d birefrin icomputer mechanica ment and f ability of	environme ovides co the specim art muscle ady state conds). which me agence due and acces al system, force data	ntal, mec ntrol of en. To c , a fast rise time asures mu to muscu sory I/O pH, temp via spec um is inv	hanical, the pH, ope with close- e of 100 ascle length dar devices perature cial purpose
PART III-TECHNICAL INFOR	MATION (FOR P	ROGRAM MAN	AGEMENT US	ES)	
1. ITEM (Check appropriate blocks)	NONE	ATTACHED	PREVIOUSLY	SEPARATE	FURNISHED LY TO PROGRAM
		Check (v/)	Approx. Date		
a. Abstracts of Theses	X		·	_	· · · · · · · · · · · · · · · · · · ·
b. Publication Citations	X		ļ		
c. Data on Scientific Collaborators	X				
d. Information on Inventions		<u> </u>			
e. Technical Description of Project and Results f. Other (specify)					Jan. 1981
2. Principal Investigator/Project Director Name (Typed)	3. Principal Invo	estigator/Project	Director Signatu	re	4. Date
Hyland Y. L. Chen	1				12 12 81
-	1				1 12 12 X/

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NSF Form 98A (5-78) Supersedes All Previous Editions

12-12-21 Form Approved OMB No. 9980013

# RECORD OF INVENTION - Part I

This is an important legal document. Read instructions carefully before filling in data.

PROJECT NO.				
CONTRACT NO CLASSIFICATION	ITY	REC. OF	65105	
1. NAME OF INVENTOR	<u> </u>	POSITION		
Hyland Y. L. Chen 2. DEPARTMENT OR DIVISION	Ass	Assistant Professor		
ESM Department, Georgia Institute of Tec	chnology			
3. DATES OF EMPLOYMENT 1973-date				
4. PRESEN: ADDRESS (No. Street, City, County, State) 20-G Plumtree Drive, Smyrna	TELEPHON 952-6		T OR UNTIL	
5. PERMANENT AODRESS (No. Street, City, County, State) ESM, Ga Tech, Atlanta, Georgia 30332	<u></u>	TELEPHONE 894-27	90	
6. NAMES (5) AND ADDRESS (ES) OF CO-INVENTORS ( If any )				
N/A				
7. DESCRIPTIVE TITLE OF INVENTION	o o m d i u m			
An Optic Probe for the viability of Myo		······································		
8. LIST DRAWINGS. SKETCHES. PHOTOS, REPORTS, DESCRIPTIONS, NOTEBOOK ENTRIES	- FTC. WHICH SHOW OP DES	CBIRE INVENTION		
<u>A schematic is attached for this report</u>				
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		· · ·	····· ······	
			·	
9. EARLIEST DATA AND PLACE INVENTION WAS CONCEIVED (Brief outline of circumsta Ga. Tech, Sept., 1975. During the desig		al system to avoid	d the "damaged	
end" artifact in heart muscle research				
sense of illumination by Hefner and of	Fourier waves	by Pollack. Con	tinuing this	
line of thought, I saw that by using la	ser light as	an electromagneti	c wave, a device	
10. DATE AND PLACE OF FIRST SKETCH. DHAWING DR PHOTO I/a to test the	viability of	myocardium in sit	u could be develope	
11. DATE AND PLACE OF FIRST WRITTEN DESCRIPTION				
12. DISCLOSURE OF INVE		DATE AND PLACE	WAS SIGNATURE	
NAME, TITLE AND ADDRESS	FORM OF DISCLOSURE	OF DISCLOSURE	OBTAINED (VES OR NO)	
Dr. Y. C. Fung, Professor, AMES-Bioengir	neering	around Dec., 1975	no	
UGSD LaJolla, Calif. 92093	discussion			
Dr. Allen Brady, Professor, Dept. of				
Physiology, UCLA. LA, Calif.	discussion (List to	around Dec., 197 ບ pe continued)	no	
12.A IMPORTANT - HAVE ANY PUBLICATIONS OF REPORTS BEEN MADE ON THIS INVENTI	ION7			
13. DATE AND PLACE OF COMPLETION OF FIRST OPERATING MODEL OR FULL SIZE DEVIC	<u>no.</u>			
	N/A			
14. PRESENT LOCATION OF MODEL	N/A			
15. DATE, PLACE, DESCRIPTION AND RESULTS OF FIRST TEST OR OPERATION	F			
NONI	Ľ			
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			, 	
			e 	

12. DISCLOSURE OF	INVENTION TO OTHERS		
NAME. TITLE AND ADDRESS	FORM OF DISCLOSURE	DATE AND PLACE OF DISCLOSURE	WAS SIGNATURE OBTAINED (VES OR NO)
or. Robert Schlant, Head, Dept. of Card Cmory University, Atlanta, Georgia Dr. L. L. Hefner, Professor, Dept. of (	diology discussion	early 1977	no
or. L. L. Hetner, Protessor, Dept. of ( 1 <u>AB, Birmingham, Alabama</u> Dr. Steve Yandlé, Naval Medical Researc		mid 1977	no
lethesda MD	discussion	Feb. 197 <b>9</b>	no
or. Newman Stevens, Professor, Univ. of	t Winnepeg, Canada discussion	around 1978	noʻ
12. DISCLOSURE OF	INVENTION TO OTHERS		
NAME, TITLE AND ADDRESS	FORM OF DISCLOSURE	DATE AND PLACE OF DISCLOSURE	WAS SIGNATURI OBTAINED (YES OR NO)
Dr. John Lundberg, Professor, Textile	Engineering	around	
Georgia Tech	discussion	early 1976	no
Dr. B. Zimm, Professor of Chemistry UCSD La Jolla, California 92093	discussion	January, 1977	no
Dr. T. Donalds, Cardiology UAB Birmingham, Alabama	discussion	around 1977	no
Dr. C. Astill, NSF	discussion	around 1977	no

Form EES 304 Page 1

16.	NAMES AND ADDRESSES OF	WITNESSES OF	FIRST TEST
	NONE		

-

17. DATE, PLACE, DESCRIPTION AND RESULTS OF LATER TESTS (name witnesses)

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18. IDENTIFY RECORDS OF TESTS AND GIVE PRESENT LOCATION OF RECORDS NONE

19. PRIOR REPORTS OR RECORDS OF INVENTION TO WHICH INVENTION IS RELATED NONE

OTHER KNOWN CLOSELY RELATED PATENTS, PATENT APPLICATIONS AND PUBLICATIONS				
DA TE	TITLE OF INVENTION OR PUBLISHED ARTICLE	NAME OF PUBLICATION		
		•		
	······································	<del></del>		
	1			

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21. EXTENT OF USE: PAST, PRESENT AND CONTEMPLATED (Give dates, places and other pertinent details)

NONE

SIGNATURE OF INVENTOR

4.5

22.	DETAILS OF INVENTION HAVE BEEN RELEASED TO THE FOLLOWING COMPANIES OR ACTIVITIES					
	NAME AND ADDRESS	INDIVIDUAL OF REPRESENTATIVE	CONTRACT NO.	DATE		
NONE						
	· · · · · · · · · · · · · · · · · · ·					

DATE

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RECORD OF INVENTION - Part II 11/12/52

TITLE OF INVENTION

(Attach to Record of Invention Part 1)

REC. OF GGICS

This Disclosure of Invention should be written up in the inventor's own words and generally should follow the outline given below. Sketches, prints, photos and other illustrations as well as reports of any nature in which the invention is referred to, if available, should form a part of this disclosure and reference can be made thereto in the description of construction and operation. See Pages 12 and 13 of the Fundamentals of Patent Technology SEES

See Fages 12 and 13 of the Fundamentals of Falent lechnology SLES. Circular No. 20.

1. INVENTORS NAME (S) HYLAND Y. L. CHEN

#### AN OPTIC PROBE OF MYOCARDIAL VIABILITY

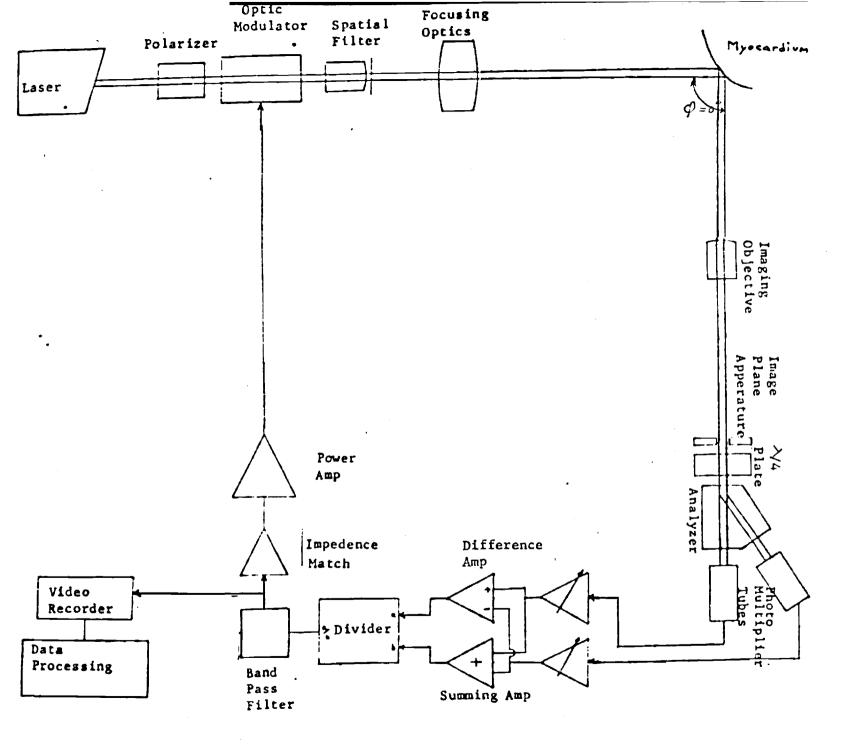
For answers to following questions use remainder of sheet, other side, and attach extrs sheets if necessary.

- 3. GENERAL PURPOSE OF INVENTION. STATE IN GENERAL TERMS THE OBJECTS OF THE INVENTION.
- 4. DESCRIBE OLD METHOD (S) IF ANY. OF PERFORMING THE FUNCTION OF THE INVENTION.
- 5. INDICATE THE DISADVANTAGES OF THE OLD MEANS OR DEVICE (S).
- 6. DESCRIBE THE CONSTRUCTION OF YOUR INVENTION, SHOW-ING THE CHANGES, ADDITIONS AND IMPROVEMENTS OVER THE OLD MEANS OR DEVICES
- 7. GIVE DETAILS OF THE OPERATION IF NOT ALREADY DESCRIBED UNDER 6.
- 8. STATE THE ADVANTAGES OF YOUR INVENTION OVER WHAT HAS BEEN DONE BEFORE.
- 9. INDICATE ANY ALTERNATE METHODS OF CONSTRUCTION.
- 10. IF A JOINT INVENTION. INDICATE WHAT CONTRIBUTION WAS MADE BY FACH INVENTOR.
- 11. FEATURES WHICH ARE BELIEVED TO BE NEW.
- 12. AFTER THE DISCLOSURE IS PREPARED. IT SHOULD BE SIGNED BY THE INVENTOR(S). AND THEN READ AND SIGNED AT THE BOTTOM OF EACH PAGE BY TWO WITNESSES USING THE FOLLOWING STATEMENT: "DISCLOSED TO AND UNDERSTOOD BY ME THIS\_\_\_\_\_OAY OF\_\_\_\_\_19\_\_\_

The optic providence of the second field of second and some office of the second s

- 3. The purpose of the invention is to spatially pinpoint rigor, resting and active regions of myocardium in pathological conditions, such as cardiac infarction, to facili-tate diagnosis and surgery by using an optic probe which picks up optical signatures in birefringence of these different regions. The electric activity of myocardium varies significantly with its states, and manifests itself in its birefringent propertie. accordingly.
- 4. & 5. In the case of open heart surgery for cardial infarction the infarcted regions are determined by x-ray angiography. The condition of the myocardium within the infarcted region, either rigor (dead), resting (due to lack of oxygen) or active is determined by inference or the guess of the surgeon. Finite element methods are under development to determine the infarcted region by spotting quantitatively the anomaly of the contraction, i.e. determining the condition of certain regions by mechanical observation, which is global.
- 6. & 7. As a laser beam passes through or scatters from muscle proper as an electromagnetic wave, the electrical activities such as membrane polarization and cross bridge orientation will differentially retard the field propagation. Thus a polarized laser beam will be depolarized. The collected beam will pass through an analyzer and register on light detectors to give a resultant time-phase shift curve. The birefringence of muscle during the course of contraction can then be probed by the time course of the depolarization, producing a distinct optic signature depending on the state of the muscl A brief block diagram is attached as an example of a possible engineering design of the probe.
- This method allows pinpoint (with spatial resolution in the order of 50 μm.) determination of muscle viability, thus eliminating the uncertainty inherent in other methods.

9. The schematic represents only one possible utilization of the basic principle.
10. N/A



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REC. OF <u>661 CS</u>

11. The optic probe tests muscle viability by a method which is categorically different from all classical methods.

DISCLOSED TO AND UNDERSTOOD BY ME ON THIS 6 DAY OF 5 cpt 19 79

INVENTOR

ON THIS \_\_\_\_\_ DAY OB COLL 19