AD_____

AWARD NUMBER: W81XWH-14-1-0562

TITLE: Development of a Lubricant Therapy to Prevent Development of Osteoarthritis after Acute Injury of Synovial Joints

PRINCIPAL INVESTIGATOR: Robert L. Sah

RECIPIENT: Dr. Prem Yadav, Ph.D.

REPORT DATE: October 2015

TYPE OF REPORT: Annual

PREPARED FOR: U.S. Army Medical Research and Materiel Command Fort Detrick, Maryland 21702-5012

DISTRIBUTION STATEMENT: Approved for public release; distribution is unlimited.

The views, opinions and/or findings contained in this report are those of the author(s) and should not be construed as an official Department of the Army position, policy or decision unless so designated by other documentation.

					Form Approved		
				wing instructions sear	OMB No. 0704-0188		
data needed, and completing a	and reviewing this collection of in	nformation. Send comments rega	arding this burden estimate or an	y other aspect of this co	blection of information, including suggestions for reducing		
4302. Respondents should be	etense, Washington Headquard aware that notwithstanding any	ers Services, Directorate for Infoi other provision of law, no persoi	mation Operations and Reports	(0704-0188), 1215 Jeff for failing to comply with	erson Davis Highway, Suite 1204, Arlington, VA 22202- n a collection of information if it does not display a currently		
valid OMB control number. PL	EASE DO NOT RETURN YOU		RESS.	3 [
October 2015		Annual		3	0 Sep 2014 - 29 Sep 2015		
4. TITLE AND SUBTIT	LE			5a.			
Development o	f a Lubricant '	Therany to Prev	ent Developmen	t of			
Ogtooprthriti	a after Agute	Indiapy do 1100	ial Tointa	5b.	5b. GRANT NUMBER		
USCEDAL CHILLES	s allel Acule	IIIJULY OL SYNOV	Tal UOINCS	W8	1XWH-14-1-0562		
				5c.	PROGRAM ELEMENT NUMBER		
6. AUTHOR(S)				5d.	PROJECT NUMBER		
Robert L. Sah							
				5e.	TASK NUMBER		
				5f.			
E-Mail: rsah@ucso					EPEOPMING OPOANIZATION PERCET		
Iniversity of	California Ca	and ADDRESS(ES) an Diego		8.6	VERFORMING ORGANIZATION REPORT		
9500 Gilman Dr	cive, Mail Code	e 0412		'			
La Jolla, CA	2093-0412						
9. SPONSORING / MC	NITORING AGENCY N	AME(S) AND ADDRES	S(ES)	10.	SPONSOR/MONITOR'S ACRONYM(S)		
			. ,				
U.S. Army Medica	I Research and Ma	teriel Command					
Fort Detrick, Maryl	and 21702-5012			11.	SPONSOR/MONITOR'S REPORT		
					NUMBER(S)		
12. DISTRIBUTION / A	VAILABILITY STATE	IENT					
Approved for put	lia ralagga diatri	hution is unlimited					
Approved for put	JIC TElease, distri						
13. SUPPLEMENTAR	I NUTES						
14. ABSTRACT							
clinical chall	lenge and may b	be associated w	ith the poor lu	ubricant qu	ality of the synovial		
fluid in the f	injured joint.	Scope. The aim	s of the studie	es here wei	re to test whether the		
pathological a	accumulation of	f a specific su	bstance found :	in joint fl	uid following an injury		
mediated alter	red synovial f	luid lubricatio	n of articular	cartilage	and whether modulation of		
lubricating mo	plecules could	restore lubric	ation function	and prever	nt cartilage wear. <u>Major</u>		
Findings. Thro	ough studies in	n year 1, we ha	ve delineated t	the time-de	ependent changes in		
lubricant mole	ecules and fund	ction of joint	riuid post-inju	iry in a ra	abbit model as well as		
Collection and	analysis of 1	uman synovial	fluid is ongoir	na (Aim 1a)	and we have completed		
most of the analysis of rabbit synovial fluid (Aim 1b). From this data, the time-dependent							
relationship between joint fluid molecules and function could be determined.							
15. SUBJECT TERMS							
Synovial Fluid, Joint							
16. SECURITY CLASSIFICATION OF:			17. LIMITATION	18. NUMBER	19a. NAME OF RESPONSIBLE PERSON		
	L ADOTD CT		UF ADJIKAUI	OF PAGES			
a. REPORT	D. ABSTRACT	C. THIS PAGE	linglocation	8	19D. IELEPHONE NUMBER (include area code)		
Unclassified	Unclassified	Unclassified	Unclassified				
Choladonica	Cholabolilou	Cholabolilou	1	l			

Table of Contents

<u>Page</u>

1.	Introduction	4
2.	Keywords	4
3.	Overall Project Summary	4
4.	Key Research Accomplishments	4
5.	Conclusion	5
6.	Publications, Abstracts, and Presentations	8
7.	Inventions, Patents and Licenses	8
8.	Reportable Outcomes	8
9.	Other Achievements	8
10.	References	8
11.	Appendices	8

- 1. INTRODUCTION: Deficient lubrication of articular cartilage by synovial fluid after synovial joint injury may advance the deterioration of cartilage, especially in the superficial zone, and hasten the development of osteoarthritis (OA). We recently delineated two mechanisms by which acute injury disrupts the normal lubrication of articular cartilage by synovial fluid: (1) diminished concentration of high-MW HA in SF, and (2) detrimental interaction of blood with SF. The scope of these studies was to test whether the impaired lubrication of articular cartilage by synovial fluid after joint injury contributes to the development of post-traumatic OA and is due, in part, to the pathological accumulation of a specific substance which can be countered by an enhanced lubricant therapy.
- 2. KEYWORDS: cartilage wear, lubrication, friction, synovial fluid, joint, osteoarthritis, post-traumatic osteoarthritis, injury, blood.

3.	ACCOMPLISHMENTS:	
3. a.	. What were the major goals o	f

were the major goals of the project?				
Specific Aim 1 (per proposal)	Timeline	Site 1		
1.1. From 60 patients, collect hSF & analyze for lubricant content, quality, & function.	Months	Completion [%]		
(a) Obtain IRB approval (VA Palo Alto).	1-3	100		
(b) Collect hSF, & transport to UCSD.	3-24	50		
(c) Analyze hSF (as fluid is collected) for biochemical properties.	3-30	50		
(d) Analyze hSF for lubricant function.	3-30	50		
Milestone(s) Achieved:				
A. VA Palo Alto IRB Approval	3	100		
B. Characterization of hSF with time after injury and severity/type of injury	15	75		
1.2. After rabbit ACL Transection (ACLT), collect & analyze (1) rSF for lubricant content, quality, & function and (2) rAC for content and structure.	Months	Completion [%]		
(a) Obtain IACUC approval (UCSD)	1-3	100		
(b) Perform rabbit ACLT surgeries, collect fluids & articular cartilage.	3-4.5	100		
(c) Analyze rSF for biochemical properties.	3-6	67		
(d) Analyze rSF for lubricant function.	3-6	33		
(e) Analyze rAC for biochemical and molecular properties.	3-9	75		
Milestone(s) Achieved:				
(A) UCSD IACUC Approval	3	100		
(B) Characterization of rSF and rAC with time after injury (rabbit ACLT)	9	50		

3.b. What was accomplished under these goals?

3.b.i. <u>Aim 1a.</u>

<u>Major Activities</u>. IRB approval was obtained at VA Palo Alto (IRB#33169) and UCSD (IRB#141410X), and the study was approved by the HRPO (A18193). hSF samples were collected at time points early (AIa, 0-14days, n=4), mid (AIb, 15-30days, n=4), and late (AIc, 31-90days, n=12) post-injury. They were analyzed for the concentrations of lubricant molecules, proteoglycan-4 (PRG4) and hyaluronan (HA), and lubricant function. Values obtained for injury samples were compared to historical normal hSF (NL-hSF), SF from knees with a chronic meniscus tear (MEN) and from osteoarthritic knees (OA-hSF).

<u>Specific Objectives.</u> The objectives for Year 1 were to a) obtain IRB approval of the study, b) collect human synovial fluid (hSF) and transport to UCSD, c) analyze hSF for biochemical properties (as it is collected), and d) analyze hSF for lubricant function.

Significant Results, Key Outcomes, Major Findings, Developments, or Conclusions. Lubricant molecule concentrations and quality as well as lubricant function were modulated by injury relative to NL samples. Relative to NL samples, protein concentrations were higher (+115-157%) in injury samples. Similarly, PRG4 concentrations were higher (+57-87%) in injury than NL samples. Relative to NL samples, HA concentrations were lower (-69-89%) in injury samples, with a shift in HA molecular weight (MW) to lower MW forms in injury samples. GAG concentrations were similar between injury groups. Generally, the kinetic ($\mu_{kinetic}$) and start-up ($\mu_{static,Tps=120}$) friction coefficients were higher in AIa than NL samples (+23-25%).

The concentrations and quality of SF lubricant constituents were altered following injuries involving the ACL and/or meniscus, in a manner consistent with our previous studies. SF samples will be analyzed for the suspected pathological substance as well as glycosaminoglycan (GAG) and collagen content to assess the extent of active cartilage degradation. It would be of interest to assess these properties in additional samples to better define the time-dependent changes in human SF at early time points following acute joint injury.

3.b.ii. Aim 1b.

<u>Major Activities</u>. IACUC approval was obtained at UCSD (IACUC Protocol #S10018). Unilateral anterior cruciate ligament transection (ACLT) surgeries in the rabbit were performed to define the time-dependent changes in synovial fluid (rSF) and articular cartilage (rAC, Aim 1b). Analysis of rSF and rAC for iron and iron-binding proteins, biochemical content, and lubricant function are on-going.

<u>Specific Objectives.</u> The objectives for Year 1 were to a) obtain IACUC approval of the study, b) perform rabbit ACLT surgeries, collect fluids & articular cartilage (for Aim 1b), c) analyze rSF for biochemical properties, d) analyze rSF for lubricant function, and e) analyze rAC for biochemical properties.

Significant Results, Key Outcomes, Major Findings, Developments, or Conclusions. Lubricant molecule concentrations and quality were modulated by injury relative to contralateral control (CTRL) samples. Relative to CTRL samples, ACLT rSF protein) concentrations were generally higher at days 1, 4, and 7 (+38-72%) and similar at days 14, 28, and 42. Relative to CTRL samples, rSF HA concentrations tended to be lower in ACLT samples at days 1 (-73%), 4 (-69%), 7 (-52%), 14 (-62%), and 28 (-50%) and similar at day 42, with HA MW being shifted to lower MW forms in ACLT samples.

The concentrations and quality of SF lubricant constituents were altered following injury involving the ACL, in a manner consistent with our previous studies. SF samples will be analyzed for the suspected pathological substance as well as lubrication function.

3.c. What opportunities for training and professional development has the project provided?

Trained undergraduate and graduate students in biochemical techniques and project management skills.

3.d. How were the results disseminated to communities of interest?

Nothing to report.

3.e. What do you plan to do during the next reporting period to accomplish the goals?

In the next reporting period, it is planned to complete on-going studies for both Aim 1a and Aim 1b. For Aim 1a, we plan to continue analyzing hSF as the samples arrive to UCSD. In addition, we plan to analyze the hSF for iron and iron-binding proteins. We are aiming for a publication by June 2016 of hSF biochemical and lubricating properties following injury and the relationship between iron and lubrication function of hSF.

For Aim 1b, we plan to measure iron and iron-binding proteins in the rSF and rAC, we plan to complete the lubricant analyses in rSF, and we plan to assess the rSF for lubrication function. In the first quarter of next year, we plan to begin the second rabbit study (Aim 2b) to assess the ability of a chelator to restore lubrication function in the joint. We are targeting a publication of the time-dependent, post-injury biochemical and lubrication properties of rSF for March 2016.

4. IMPACT:

4.a. What was the impact on the development of the principal discipline(s) of the project?

Nothing to report. However, a publication is pending.

4.b. What was the impact on other disciplines?

Nothing to report. However, a publication is pending.

4.c. What was the impact on technology transfer?

Nothing to report.

4.d. What was the impact on society beyond science and technology?

Nothing to report. However, a publication is pending.

5. CHANGES/PROBLEMS:

5.a.Changes in approach and reasons for change.

Nothing to report.

5.b. Actual or anticipated problems or delays and actions or plans to resolve them.

Analyses of rabbit synovial fluids required some scaling of the various assays. That work has been completed and analyses are proceeding.

5.c. Changes that had a significant impact on expenditures.

Nothing to report.

5.d. Significant changes in use or care of human subjects, vertebrate animals, biohazards, and/or select agents.

Nothing to report.

- 5.d.i. Significant changes in use or care of human subjects
- 5.d.ii. Significant changes in use or care of vertebrate animals.
- 5.d.iii. Significant changes in use of biohazards and/or select agents

6. PUBLICATIONS, ABSTRACTS, AND PRESENTATIONS:

- 6.a. Lay Press:
- 6.b. Peer-Reviewed Scientific Journals:
- 6.c. Invited Articles:
- 6.d. Abstracts:
- 6.e. Presentations:

7. INVENTIONS, PATENTS AND LICENSES:

Nothing to report.

8. REPORTABLE OUTCOMES:

Nothing to report.

9. OTHER ACHIEVEMENTS:

Nothing to report.

10. REFERENCES:

11. APPENDICES:

Development of a Lubricant Therapy to Prevent Osteoarthritis after Acute Injury Log Number: W81XWH-14-1-0562

Award Number: OR130385

PI: Robert L. Sah

Org: University of California, San Diego

Award Amount: \$1,128,083

Study/Aim(s)

• Determine time course & extent of increased iron in SF and articular cartilage after knee joint injury

• Optimize the dosage & time course of chelation therapy to deplete pathological iron from cartilage

• Assess the ability of chelation-enhanced lubricant therapy to restore lubrication function and prevent wear

Approach

- SF will be obtained from the injured (AI) or normal (NL) joint of patients or rabbits at multiple times after injury
- SF will be analyzed for iron, iron-binding proteins, and lubricant content and function.
- Iron-laden cartilage following chelator treatment will be analyzed for iron content & lubrication properties.
- AI-SF ±chelator ±lubricants will be assessed for wear protective properties relative to NL-SF *in vitro* and *in vivo*.

Timeline and Cost

Activities CY	14	15	16	17
Obtain human & animal protocol approvals				
Determine iron profile after injury				
Optimization of chelation therapy				
Determine SF±therapy function				
Estimated Budget (\$K)	\$95	\$378	\$375	\$282

Updated: 01/06/2015



Accomplishment: Studies are in progress to analyze the SF lubricant properties after injury *in vitro* and *in vivo*. The overall goal is to develop an enhanced lubricant therapy to restore SF lubricant function after acute joint injury and prevent post-traumatic osteoarthritis.

Goals/Milestones

CY14 Goals – Obtain study approvals

- ☑ Obtain UCSD IRB & USAMRMC HRPO approval
- ☑ Obtain IACUC & USAMRMC ACURO approval
- CY15 Goal SF characterization & Optimization of chelation therapy
- Collect & analyze human SF biochemical properties & function
- \Box Collect & analyze rabbit SF biochemical properties & function

 $\Box Determine chelator dose-dependent iron removal from human cartilage$

 $\Box Determine \ post-chelator \ hSF$ lubricant function

 \Box Determine chelator dose-dependent iron removal from rabbit cartilage

Determine post-chelator rSF lubricant function

CY16 Goal – Develop enhanced lubricant therapy

□ Determine enhanced AI-hSF & AI-rSF lubricant function with chelator and lubricant supplements

Comments/Challenges/Issues/Concerns

None

Budget Expenditure to Date

Projected Expenditure:	\$379,938
Actual Expenditure:	\$320,502