

AWARD NUMBER: W81XWH-19-1-0862

TITLE: A Comparative Approach to Human Auditory Synaptopathy

PRINCIPAL INVESTIGATOR: Marjorie Leek, Ph.D.

CONTRACTING ORGANIZATION: Loma Linda Veterans Association for Research & Education, Redlands, CA

REPORT DATE: October 2023

TYPE OF REPORT: Annual

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

DISTRIBUTION STATEMENT: Approved for Public Release;  
Distribution 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.

# REPORT DOCUMENTATION PAGE

*Form Approved*  
*OMB No. 0704-0188*

Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing this collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to Department of Defense, Washington Headquarters Services, Directorate for Information Operations and Reports (0704-0188), 1215 Jefferson Davis Highway, Suite 1204, Arlington, VA 22202-4302. Respondents should be aware that notwithstanding any other provision of law, no person shall be subject to any penalty for failing to comply with a collection of information if it does not display a currently valid OMB control number. **PLEASE DO NOT RETURN YOUR FORM TO THE ABOVE ADDRESS.**

<b>1. REPORT DATE</b> October 2023		<b>2. REPORT TYPE</b> Annual		<b>3. DATES COVERED</b> 30Sep2022 -29Sep2023	
<b>4. TITLE AND SUBTITLE</b>  A Comparative Approach to Human Auditory Synaptopathy				<b>5a. CONTRACT NUMBER</b> W81XWH-19-1-0862	
				<b>5b. GRANT NUMBER</b> RH180051	
				<b>5c. PROGRAM ELEMENT NUMBER</b>	
<b>6. AUTHOR(S)</b>  Marjorie R. Leek, Ph.D., Edward J. Walsh, Ph.D.  E-Mail: Marjorie.Leek@va.gov				<b>5d. PROJECT NUMBER</b>	
				<b>5e. TASK NUMBER</b>	
				<b>5f. WORK UNIT NUMBER</b>	
<b>7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES)</b> Loma Linda Veterans Association for Research and Education LLVARE 25884 Business Center Drive, Suite A Redlands, CA 92373				<b>8. PERFORMING ORGANIZATION REPORT NUMBER</b>	
<b>9. SPONSORING / MONITORING AGENCY NAME(S) AND ADDRESS(ES)</b>  U.S. Army Medical Research and Development Command Fort Detrick, Maryland 21702-5012				<b>10. SPONSOR/MONITOR'S ACRONYM(S)</b>	
				<b>11. SPONSOR/MONITOR'S REPORT NUMBER(S)</b>	
<b>12. DISTRIBUTION / AVAILABILITY STATEMENT</b>  Approved for Public Release; Distribution Unlimited					
<b>13. SUPPLEMENTARY NOTES</b>					
<b>14. ABSTRACT</b> Exposure to noise can cause damage to structures in the inner ear, often resulting in a loss of hearing. Recent findings in noise-exposed animals raise a new specter that even moderate noise exposures may result in damage specifically located in the synaptic region between the sensory cells in the cochlea and primary auditory neurons. There is no way currently that scientists and clinicians can diagnose possible auditory synaptic damage in humans, and diagnosis is critical for the development of innovative treatments. The objective of this project is to develop a statistical model that will accurately predict the likelihood of synaptopathy in humans who have had noise exposures in their lives. The development of the statistical model will be supported by collecting non-invasive measurements in both humans and guinea pigs. Findings from the animal testing have identified several metrics that show promise for differentiating noise-exposed from control animals, including newly created analyses of evoked potential and otoacoustic emission testing. These metrics will be tested further with increasing animal data to determine if they are candidates for inclusion in the statistical model of synaptopathy under development. Successful metrics will subsequently be applied to the human data to predict synaptopathy.					
<b>15. SUBJECT TERMS</b> Human hearing loss, guinea pig, Noise exposures, Neural deficits, Statistical model development					
<b>16. SECURITY CLASSIFICATION OF:</b>			<b>17. LIMITATION OF ABSTRACT</b>  Unclassified	<b>18. NUMBER OF PAGES</b>  14	<b>19a. NAME OF RESPONSIBLE PERSON</b> USAMRDC
<b>a. REPORT</b>  Unclassified	<b>b. ABSTRACT</b>  Unclassified	<b>c. THIS PAGE</b>  Unclassified			<b>19b. TELEPHONE NUMBER</b> (include area code)

# TABLE OF CONTENTS

	<u>Page(s)</u>
1. Introduction .....	4
2. Keywords .....	4
3. Accomplishments .....	4-7
4. Impact .....	7-8
5. Changes/Problems .....	8-9
6. Products .....	9-10
7. Participants & Other Collaborating Organizations .....	11-13
8. Special Reporting Requirements .....	13
9. Appendices .....	14-33

## 1.Introduction

Exposure to noise has long been known to cause damage to structures in the inner ear, often resulting in a loss of hearing. This is a significant health concern for millions of people and it will intensify as the population ages and loud noises become a more prevalent feature of the soundscape. Recent findings in noise-exposed animals raise a new specter that even moderate noise exposures may result in damage specifically located in the synaptic region between the inner hair cells in the cochlea and primary auditory neurons. Although this synaptopathy has been demonstrated in several animal models, there is no way currently that scientists and clinicians can diagnose possible synaptic damage in humans, and diagnosis is critical for the development of new, innovative treatment plans. The objective of this project is to develop and populate a statistical model that is designed to accurately predict the presence and degree of synaptic damage resulting from noise exposure in guinea pigs, and subsequently to use that model to predict the likelihood of synaptopathy in humans who have had significant noise exposures in their lives. Auditory function will be assessed using a variety of non-invasive tests following noise exposure in non-human animals, and cochleae will subsequently be analyzed for evidence of synapse loss or damage. Performance on tests of auditory function in non-human animals will be compared with human performance to develop a predictive model that can be refined and extended by adding results from behavioral tests of temporal processing and speech perception in noise by human listeners. Finally, a comprehensive statistical predictive model will be developed to evaluate the likelihood and severity of synaptic damage in individual humans.

## 2.Keywords

Human hearing loss, guinea pig, Noise exposures, Neural deficits, Statistical model development

## 3.Accomplishments

**What were the major goals of the project?**

- a. Establish and set up final electrophysiological and acoustic diagnostic tests. Milestone 1 to be completed by end of 4<sup>th</sup> month (i.e., January 31, 2020). This goal was fully realized in the Walsh (animal) lab in November 2020. Completion of this goal in the Leek (human) lab was achieved during this annual reporting period.
- b. Acquire test data on 12 guinea pigs per quarter. This was partially completed during the current reporting period.
- c. Perform anatomical assessment of 12 guinea pigs per quarter. This aspect of the project was partially completed during the current reporting period.
- d. Acquire physiological and behavioral data on 10 human subjects per quarter. This has been partially completed during this year of the award.

## What was accomplished under these goals?

- a. This goal has been fully accomplished in both the Walsh (animal) lab and the Leek (human) lab. All testing protocols are in place and equipment set-up and calibration have been completed.

On-line meetings of the entire research team have been held during this year to address issues involved in creating the most similar testing procedures between the two sides of the project. These discussions have been informed largely by the successful implementation of data collection in guinea pigs and from human subjects, and by new findings in the literature. We are continuing to seek efficiencies to accelerate the rate of data collection in the two labs without penalty to the ultimate goals of the work.

- b. Electrophysiology/acoustic data sets have been acquired from 84 animals and another 4 are currently acclimatizing and are scheduled for study. Findings based on data obtained from 52 noise-exposed animals and 32 sham animals support our *a priori* view that a subset of auditory response elements reliably differentiate noise-exposed from control animals (see Appendix 1). Findings from early efforts to lay out and consider a subset of essential elements of the proposed multivariate statistical model with significant predictive/diagnostic power during the current reporting period are highly encouraging.
- c. Progress was made on this aspect of the project during the past funding year as well. One hundred and sixty-six inner ear specimens (both right and left cochleae) have been perfused with fixative, 86 have been processed for histological assessment of hair cell and synapse pathology and cochlear images have been acquired on 85 ears. Hair cell and synapse counts are complete in the case of 81 cochleae acquired from 47 noise-exposed and 32 sham animals. Quantitative analysis is currently in progress on immunofluorescent images acquired from 4 additional cochleae. The data acquired are consistent with the expectation that ribbon synapses are lost at predictable levels in noise-exposed animals.
- d. Data collection continues in the human lab. So far, we have completed all measurements for 16 subjects. We currently have additional participants enrolled and moving through the testing. We continue to recruit additional subjects using flyers in the audiology clinic, describing our work to the audiology staff and asking for referrals, and working actively with our public affairs office to place our recruitment advertisement in the rotation of the informational closed-captioning televisions posted around the VA hospital.

**Please see Appendices for further details of progress in the animal and human labs and preliminary work on model development.**

### **What opportunities for training and professional development has the project provided?**

Although this project was not intended to provide formal training and professional development opportunities, as a matter of course, the more senior members in each lab (Walsh and Leek labs) do encourage professional and scientific growth among the less senior team members. For example, Dr. Venezia, who is an accomplished statistician and modeler, has led our team in understanding the type of modeling we plan to successfully complete this study. He also has led the team in understanding his career development grant, which includes both behavioral testing of human subjects, as well as functional MRI techniques. Research Assistant Kelli Sugai works closely with other team members to learn how to work with human subjects on data collection, including enrollment procedures such as acquiring informed consent from subjects. She also has received training on how to carry out the various auditory tests required for this study. Two postdoctoral fellows in the lab, while not formally part of this study, are included in discussions of this work, and are being trained to recognize signs of hearing loss and other characteristics of audiological evaluation. Each of them brings new research projects to the lab in which all lab members are involved. Members of the Leek lab have learned more about working with animal subjects, carrying out the regulatory procedures for animal work, and learning how animal electrophysiology will be performed to mimic (as much as possible) the human testing. It is our practice for the entire Auditory Research Group at the VA hospital (some 13 scientists and technicians working with humans or with small animals) to meet every two weeks (currently virtually on Zoom) to hear about the ongoing research and data from each lab or to review a current article that is of interest to the labs. Because our group has a wide range of interests and expertise, this is an enjoyable and valuable learning experience, and encourages the social bonding of the group, as well as keeping us all up to speed on new developments in auditory research.

Specific career training and professional development has been implemented during the year in the Walsh lab. This training has enhanced the skills and capabilities of both Research Technicians, Dr. Xiaohui Lin and Ms. Ashley Vazquez, in this area of auditory research. Although Dr. Lin had previous experience in the auditory neurosciences, Drs. McGee and Walsh extended the training of both assistants in the realm of auditory evoked potentials during the past year. The training was designed to enhance data acquisition and analyses capabilities and both technicians successfully acquired these skills. Software adjustments were also made to advance quantitative analyses of immunofluorescent cochlear images and both technicians are now proficient in their implementation. In addition, Dr. Lin was able to join Drs. Walsh and McGee at a meeting of the "Southern California Hearing Research Conference and Inaugural Neil Segil Symposium in Hearing and Communications Neuroscience" hosted by the University of Southern California. Dr. Lin presented the poster, "Envelope following responses following partial cochlear deafferentation in guinea pigs" that was presented previously at the Acoustical Society of America meeting in Nashville. Presentations by well-recognized keynote speakers were outstanding, as were presentations from participants in general. The meeting was a generally enriching experience.

**How were the results disseminated to communities of interest?**

During the last year, a presentation regarding results of this project was made at the Fall (183<sup>rd</sup>) Meeting of the Acoustical Society of America held in Nashville, TN in December 2022, as well as at the “Southern California Hearing Research Conference and Inaugural Neil Segil Symposium in Hearing and Communications Neuroscience” held at USC in 2023.

McGee, J., Lin, X., Vazquez, A., Li, H., Venezia, J., Leek, M.R., and Walsh, E.J. (2022). *Envelope following responses following partial cochlear deafferentation in guinea pigs*. (Abstract #3pPP12 published in J. Acoust. Soc. Am.. 152 (no.4, Pt.2): A197.

In addition, an abstract was submitted to the Association for Research in Otolaryngology for the 47<sup>th</sup> annual midwinter meeting that will be held in February of 2024 in Anaheim, CA.

McGee, J., Lin, X., Vazquez, A., Li, H., Venezia, J., Leek, M.R., and Walsh, E.J. (2024). *Analysis of Receiver Operating Characteristics to Evaluate the Performance of Predictor Variables for the Diagnosis of Cochlear Synaptopathy*. Assoc. Res. Otolaryngol. 47.

**What do you plan to do during the next reporting period to accomplish the goals?**

We have received approval for another no-cost extension year to support continued data collection in both the animal and human labs through September 2024.

Electrophysiology and acoustic testing in the Walsh lab are proceeding actively and findings thus far are highly promising with regard to the identification of auditory response features differentiating noise-exposed from control animals (see Appendix 1). During the next reporting period, the plan is to carry on with the essential aspects of the data acquisition protocol currently in use. In addition, time and effort on model development aspects of the project will be continued.

Two major goals will be the focus of the next year of this grant (under a second no-cost extension). First is to continue testing animal subjects and accelerate the recruitment and testing of human subjects. The second is to further develop the predictive model based on guinea pig data and extend it to test its value with human data. This, of course, is the major purpose of this work.

**4.Impact**

**What was the impact on the development of the principal discipline(s) of the project**

Nothing to Report.

**What was the impact on other disciplines?**

Nothing to Report.

**What was the impact on technology transfer?**

Nothing to Report.

## What was the impact on society beyond science and technology?

Nothing to Report.

## 5.Changes/Problems

### Changes in approach and reasons for change

Nothing to Report.

### Actual or anticipated problems or delays and actions or plans to resolve them

In our previous progress reports we have described some significant barriers to implementing and progressing on this project as rapidly as we had envisioned. These delays have included major disruptions to both the animal and human work due to shutdowns and limited access to our labs and to human research subjects as a response to the COVID-19 pandemic, and extended delays in the availability of appropriate and functional lab space and equipment.

In the animal lab: Work on acquiring electrophysiological and acoustical data on guinea pigs and anatomically assessing hair cell survival and ribbon synapses in the cochlea will continue in the upcoming period.

In the human lab: One matter has served to slow down our ability to recruit large numbers of human subjects and that is the medical center administration's decision to discontinue posting advertisements for subjects who are interested in participating in research projects at the hospital. With the support of the facilities Research Administration Office, we have met several times with the leaders of the Public Affairs Office to register our dismay at this decision, but so far have not been able to make any progress toward reinstating those ads. We are hopeful that additional discussions in the near future will assist in the resolution of this condition.

## Changes that had a significant impact on expenditures

Nothing to Report.

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

### Significant changes in use or care of human subjects

Nothing to Report

### Significant changes in use or care of vertebrate animals

Nothing to Report.

### Significant changes in use of biohazards and/or select agents

Nothing to Report.

## 6.Products

### Publications, conference papers, and presentations

#### Journal publications

Nothing to report

#### Books or other non-periodical, one-time publications

Nothing to Report.

#### Other publications, conference paper, and presentations

Conference papers:

A presentation regarding results of this project was made at the Fall (183<sup>rd</sup>) Meeting of the Acoustical Society of America held in Nashville, TN in December 2022, as well as at the "Southern California Hearing Research Conference and Inaugural Neil Segil Symposium in Hearing and Communications Neuroscience" held at USC in 2023.

McGee, J., Lin, X., Vazquez, A., Li, H., Venezia, J., Leek, M.R., and Walsh, E.J. (2022). *Envelope following responses following partial cochlear deafferentation in guinea pigs*. (Abstract #3pPP12 published in J. Acoust. Soc. Am. 152 (no.4, Pt.2): A197)

In addition, an abstract for a conference was submitted to the Association for Research in Otolaryngology for the 47<sup>th</sup> annual midwinter meeting that will be held in February of 2024 in Anaheim, CA.

McGee, J., Lin, X., Vazquez, A., Li, H., Venezia, J., Leek, M.R., and Walsh, E.J. (2024). *Analysis of Receiver Operating Characteristics to Evaluate the Performance of Predictor Variables for the Diagnosis of Cochlear Synaptopathy*. Assoc. Res. Otolaryngol. 47.

**Website(s) or other internet site(s)**

Nothing to Report.

**Technologies or techniques**

Nothing to Report.

**Inventions, patent applications, and/or licenses**

Nothing to Report.

**Other Products**

Nothing to Report.

Please continue to next page

## 7.Participants & Other Collaborating Organizations

### What individuals have worked on the project?

Name	<i>Marjorie R. Leek, Ph.D.</i>
Project Role	<i>Principal Investigator</i>
Researcher Identifier	<i>None</i>
Nearest person month worked	<i>2</i>
Contributions To Project	<i>Prepared experimental programming for testing protocols; generated stimuli for physiological and behavioral testing of humans, completed regulatory submissions; supervised research audiologist/research assistant, attended and contributed to team meetings and discussions</i>
Funding Support	<i>This project</i>

Name	<i>Edward J. Walsh, Ph.D.</i>
Project Role	<i>Co-Principal Investigator</i>
Researcher Identifier	<i>None</i>
Nearest person month worked	<i>4.8</i>
Contributions To Project	<i>Oversaw lab activities; provided education, ongoing training, and supervision to lab technicians; conducted relevant data analyses; searched and reviewed literature relevant to studies proposed in this grant; participated in administrative meetings addressing a range of relevant program items, responsible for maintaining updated regulatory protocols. Participated in dissemination of project results.</i>
Funding Support	<i>This project</i>

Name	<i>JoAnn McGee, Ph.D.</i>
Project Role	<i>Co-Investigator</i>
Researcher Identifier	<i>None</i>
Nearest person month worked	<i>9</i>
Contributions To Project	<i>Participated in team meetings that address a range of essential program items, generated stimuli and protocols for data acquisition activities, wrote and refined programs for data and statistical analyses; performed analyses including statistical testing, prepared figures for presentation, conducted relevant literature surveys, participated in training lab technicians; contributed to regulatory protocols, participated in dissemination activities for this project.</i>
Funding Support	<i>This project</i>

Name	<i>Jonathan H. Venezia, Ph.D.</i>
Project Role	<i>Co-investigator</i>
Researcher Identifier	<i>None</i>
Nearest person month worked	<i>1</i>
Contributions To Project	<i>Wrote experimental programming for testing protocols; developed initial modeling efforts based on guinea pig and human data; generated stimuli and experimental programming for electrophysiological tests in human subjects; attended and contributed to team meetings and discussions</i>
Funding Support	<i>Institutional funds</i>

Name	<i>Hongzhe Li, Ph.D.</i>
Project Role	<i>Co-Investigator</i>
Researcher Identifier	<i>None</i>
Nearest person month worked	<i>1</i>
Contributions to Project	<i>Performed animal terminal procedures including cardiac perfusion, followed by subsequent cochlear tissue processing; microdissection of cochlear sample for confocal identification of ribbon synapses; confocal image processing and synapse quantification.</i>
Funding Support	<i>This project</i>

Name	<i>Xiaohui Lin, Ph.D.</i>
Project Role	<i>Research Technician</i>
Researcher Identifier	<i>None</i>
Nearest person month worked	<i>12</i>
Contributions To Project	<i>Responsible for animal management and purchase of consumable supplies needed for animal studies; responsible for electrophysiological and acoustic data acquisition, and basic physiological analyses of data acquired from guinea pigs. Participated in preparation of animals for histological processing, served as a secondary resource for anatomical studies. Took part in regulatory protocol submissions. Participated in group meetings associated with this grant.</i>
Funding Support	<i>This project</i>

Name	<i>Kelli Sugai</i>
Project Role	<i>Research Assistant</i>
Researcher Identifier	<i>None</i>
Nearest person month worked	<i>6</i>
Contributions to Project	<i>Continued recruiting potential human subjects for the study; enrolled subjects and collected behavioral and electrophysiological data record keeping/organizational documents; assisted with regulatory submissions; attended and contributed to team meetings.</i>
Funding Support	<i>This project</i>

Name	<i>Ashley Vazquez, B.S.</i>
Project Role	<i>Research Technician</i>
Researcher Identifier	<i>None</i>
Nearest person month worked	<i>12</i>
Contributions To Project	<i>Participated in animal management and took responsibility for anatomical preparation and imaging of cochlear specimens and for quantitative analyses of hair cell numbers and ribbon synapses. Assisted in electrophysiological and acoustic data collection and a portion of physiological analyses of data acquired in guinea pigs. Took part in regulatory protocol submissions. Participated in group meetings associated with this grant.</i>
Funding Support	<i>This project</i>

**Has there been a change in the active other support of the PD/PI(s) or senior/key personnel since the last reporting period?**

Nothing to Report.

**What other organizations were involved as partners?**

Nothing to Report.

## **8.Special Reporting Requirements**

### **Collaborative Awards**

### **Quad Charts**

Quad chart is not attached because we are not expending any funds from the DoD.

## **9. Appendices**

- Appendix 1 Report of Guinea Pig Testing
- Appendix 2 Report of Human Subjects Testing
- Appendix 3 Report of Preliminary Model Development
- Appendix 4 Abstract submitted