Award Number:  W81XWH-12-2-0018

TITLE:   NRC/AMRMC Resident Research Associateship Program

PRINCIPAL INVESTIGATOR:   Howard R. Gamble, Ph.D.

CONTRACTING ORGANIZATION:  National Academy of Sciences
                             Washington, DC 20001

REPORT DATE: April 2013

TYPE OF REPORT: Annual

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

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<td>During this reporting period, the NRC promoted research opportunities at AMRMC institutes through a broad outreach plan. A total of 36 applications were received during the period and of these, 30 were reviewed by NRC panels. A total of 17 award offers were made and all 17 applicants accepted the awards. The productivity of these Associates is listed in the technical report</td>
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19b. TELEPHONE NUMBER (include area code)
During the reporting period, the NRC conducted the following activities in support of the subject contract:

**Outreach and Promotion**

The promotional schedule to advertise the National Research Council (NRC) Research Associateship Programs included the following: 1) attendance at meetings of major scientific and engineering professional societies; 2) advertising in programs and career centers for these and other professional society meetings; 3) direct mailing and emailing of announcements and program materials to presidents, graduate deans, and heads of appropriate science and engineering departments and minority-affairs offices of all academic degree-granting institutions in the United States; 4) posting announcements on internet job sites, electronic newsletters and professional society websites; 5) print advertising in high profile publications (e.g., Science magazine, the Chronicle of Higher Education); and, 6) maintaining a presence on social media sites such as Facebook.

The NRC attended a number of minority focused events in which we maintained exhibit booths, participated in workshops and advertised in meeting literature, newsletters and websites or submitted materials for distribution. In addition, ads were placed in a variety of minority publications (e.g., Affirmative Action, Black Collegian).

In advertising the Research Opportunities available to prospective applicants, the NRC maintained an up-to-date listing of all active Research Advisers, current Adviser contact information and details of each Research Opportunity.

**Processing and Review of Applications**

Applications to the Research Associateship Program were submitted via a web-based application system. Each of the four application cycles opened two months prior to the application deadline. NRC staff provided support to prospective applicants including providing application instructions, technical support and additional information as requested.

A summary of applications for the reporting period is shown in Table 1.

For each applicant, the NRC received and processed an application form, a research proposal, transcripts, a statement of previous and current research, and confidential reference reports. An application file check was made prior to the review and each applicant was notified if required documents were missing.

The NRC convened panels in five broad discipline areas for the competitive review of applications in the Research Associateship Programs. Results of the review were made available to Laboratory Program Representatives immediately following the conclusion of the each review.

A summary of the outcome of the review of applications for the reporting period is shown in Table 1.

**Administration of Awards**

The NRC made awards to applicants based on sponsor authorization. A summary of awards authorized and the acceptance or declination by the applicant during the current reporting period is shown in Table 1.

For Associates beginning or continuing tenure, the NRC provided the administrative functions described in the contract Statement of Work. These functions included stipend payments, management of a major medical benefits insurance program, and reimbursement for relocation and travel to professional meetings.
A summary of NRC Research Associates on tenure during the reporting period is shown in Table 2.

**Outcomes Reporting**

All NRC Associates who completed tenure were required to submit a final report that described the outcome of their Associateship award. Final reports received by the NRC during the current reporting period are attached to this technical report.

The activities of Associates submitting final reports during this reporting period, including publications, presentations and patents, as well as an assessment of their experience in the program, are summarized in Table 3. Specific research accomplishments of Associates completing tenure during the reporting period are summarized in Table 4.

**Table 1.** Applications and Awards

**Table 2.** Associates on Tenure

**Table 3.** Associates Activity

**Table 4.** Summary of Associate Research

**Attachments:** Associate Final Reports
Table 1: Applications and Awards

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Table 3: Associates’ Activities

14 Associates ended tenure during the report period
24 months was the average tenure length
42 months was the longest
9 months was the shortest
13 submitted final reports

In the final reports, Associates indicated the following scholarly activity while on tenure.
- 17 Articles published in refereed journals
- 0 Patent applications
- 6 International presentations
- 34 Domestic presentations
- 3 Awards

After ending their tenure, Associates indicated their future plans as follows:
- 1 Permanent position at the NRC host agency
- 6 Contract or temporary position at the NRC host agency
- 1 Research/administrative position with another U.S. government agency
- 1 Research/administrative position with foreign government agency
- 1 Research/teaching at US college/university
- 1 Research/teaching position at a foreign college or university
- 0 Research/administrative position in private industry in the U.S.
- 0 Research/administrative position in private industry outside of the U.S.
- 0 Research/administrative position with a non-profit
- 0 Self-employed/consulting
- 2 Postdoctoral Research
- 0 Other
- 0 No information provided

In their final reports, Associates were asked to evaluate certain aspects of their experiences on a scale of 1 (low) to 10 (high). The average rating for each item follows:

9.4 Short-term value (lab)-Development of knowledge, skills, and research productivity at lab
9.2 Long-term value (career)-How your Research Associateship affected your career to date
8.8 Laboratory Support-Equipment, funding, orientation, safety & health training, etc.
9.5 Adviser Mentoring-Quality of mentoring from the Research Adviser
9.2 LPR Support-Quality of administrative support from the LPR
9.2 NRC Support-Quality of administrative support from the NRC
### Table 4: Summary of Associate Research

<table>
<thead>
<tr>
<th>Associate</th>
<th>Tenure Dates</th>
<th>Summary</th>
</tr>
</thead>
</table>
| Boutte, Angela | 6/15/2011-1/14/2013       | 1. Determined the time course of brain derived biomarkers elevated in brain tissue and serum of a rat model of penetrating ballistic-like brain injury (PBBI) (on-going)  
2. Determined the time course of brain derived biomarkers elevated in brain tissue and serum of a rat model post-traumatic stress disorder (fear conditioning) and blast injury (on-going)  
3. Determined the effect of therapeutic drugs to prevent increases in brain and serum biomarkers after PBBI  
4. Initiated proteomics analysis of a rat model post-traumatic stress disorder (PTSD) and blast injury (on-going)  
5. Grants - Submitted grant proposal to define the markers and mechanism of severe and mild long term brain injury (Combat Casualty Care Program), co-authored pilot grant applications to determine efficacy of therapeutics in PBBI.  
6. Publications - Under the guidance of Dr. Dave, our lab published our proteomics study in the journal Electrophoresis. In a separate cohort using proteomics and bioinformatics prediction of biomarkers, I aided in defining novel biomarkers of PBBI in brain tissue that I have independently confirmed. A bioinformatics-themed manuscript is complete and pending submission. |
2. Peptide-oligonucleic acid conjugates for antisense  
3. Co-polymer enhanced liposome for antisense ODN delivery |
| Johnston, Sara | 3/25/2009-5/1/2012        | 1. MPXV Active Disease Surveillance Program identifying significant increases in prevalence in DRC  
2. Identification of potential novel therapeutic (IFN-beta) against MPXV  
3. Identification of MPXV variants actively circulating in the DRC  
4. In vitro and in vivo evaluation of novel therapeutics against Henipaviruses  
5. Development and implementation of diagnostic ELISA assays against MPXV |
2. Study of differences in respiratory pattern between healthy animals and animals with the Acute Respiratory Distress Syndrome  
3. Development of a new technique for the placement of bicaval dual-lumen catheters for venovenous extracorporeal gas exchange  
4. Effects of radiation dose reduction on lung quantitative CT scan results in healthy in the Acute Respiratory Distress Syndrome: low-dose chest CT as a valuable tool for quantification and monitoring of pulmonary disease reducing patient exposure |
| Leung, Lai Yee | 2/16/2010-1/31/2013       | 1. Established polytrauma models associated with hypoxemia and hemorrhagic hypotension that will be used for future neuroprotective drug studies.  
2. Acute physiological changes were characterized in the polytrauma models. The patterns of these changes were found to be unique under different injury combinations.  
3. Hemorrhagic shock increased the incidence and duration of cortical spreading depolarization within 2 hours following PBBI whereas hypoxemia only prolonged the depolarization.  
4. Histopathological changes were characterized (3, 7 days post-injury) in the polytrauma models. Hemorrhagic shock increased neuronal degeneration and astrocytic activation following PBBI.  
5. The sequence of PBBI, hemorrhagic hypotension and hypoxemia affected the mortality rate and neurological deficits. PBBI, HS followed by HX resulted in the highest mortality rate (50%) and more neurological deficits among all polytrauma groups. |
| McCoy, Margaret | 7/27/2009-10/4/2012       | 1. Identified the mechanism of action of SAPN-induced Ab that provides sterile immunity in mice  
2. Designed and carried out experiments to examine and characterize the processing and presentation of SAPN within the immune system |
3 Examined the phenotypes of SAPN-specific T-cell populations and proved that CD8+ T-cells from mice immunized with SAPN are able to, by themselves, induce sterile immunity in mice - this is the first malaria vaccine to be able to show this.
4 Examined needleless approaches to vaccine delivery, including pleuronic lecithin organogel creams
5 Identified potential effects on vaccine efficacy resulting from the addition of mosquito saliva
6 The youngest investigator at the WRAIR to successfully write and complete a non-human primate trial for malaria

<table>
<thead>
<tr>
<th>Meledeo, Michael</th>
<th>3/1/2010-7/6/2012</th>
</tr>
</thead>
</table>
| 1 Apartimers can be used to completely inhibit the anti-coagulant effects of activated protein C (aPC); however, aPC does not appear to be the sole sufficient cause of the acute coagulopathy of trauma.
2 Exposing in vitro cultures of endothelial cells (ECs) to laminar flow (as a model of their physiological environment) will induce a number of changes to both EC morphology and gene expression in a variety of inflammatory and morphology pathways.
3 While not all of the changes in gene expression result in an altered proteome, there are a number of significant differences in protein expression between static cultured ECs and those exposed to flow.
4 An analysis of the EC glycocalyx through confocal microscopy and western blotting of membrane proteins and associated glycoforms has led to an enhanced understanding of the structure and function of the endothelial glycocalyx layer.
5 All of these have provided advancement in the formulation of in vitro models of endothelium; in the future it should be possible to use these models as a platform for the testing of both therapeutics and diagnostics for vascular dysfunction.

<table>
<thead>
<tr>
<th>Melendrez, Melanie</th>
<th>3/22/2010-5/21/2012</th>
</tr>
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</table>
| 1 Variants analysis of dengue quasispecies populations showed that variants are host or vector specific despite containing the same consensus sequence. Diversity was found to not be constrained within the vector as suggested in some publications.
2 Selection analysis showed the populations to be expanding, evolving at a faster rate than 'average' for the dataset, and were under selective pressure with a predominance of nonsynonymous mutations when compared with the consensus sequence.
3 Phylogenetic analysis revealed that dengue quasispecies sequences isolated in 2010 were distinct from other circulating consensus sequences from Thailand and full E gene offered higher resolution than partial E gene sequences.
4 Amino acid (aa) analysis suggested several positions where changes would affect replication, antibody binding or VLP assembly according to the literature. Multivariate analysis predicted uncharacterized aa positions that would have impact if altered.
5 This work revealed the importance of full E gene surveillance for assessment of aa changes, illustrated the variability and pathogenic potential of dengue quasispecies variant diversity and established a baseline in which to make future comparisons.

<table>
<thead>
<tr>
<th>Pichugin, Alexander</th>
<th>1/12/2009-7/11/2012</th>
</tr>
</thead>
</table>
| 1 10 novel liver stage Pb antigens reduce LS and BS parasite burden in C57Bl/6 mice.
2 3 novel liver stage Pb antigens sustain protection during 6 months after the last immunization.
3 3 novel liver stage Pb antigens enhance protection induced by PbCSP.
4 Established caged MHC-tetramer technology to use for discovery of T cell epitopes in malaria antigens.
5 Identified 3 immunodominant CD8 T cell epitopes from Pb PEVA.

<table>
<thead>
<tr>
<th>Rajendran, Gnana</th>
<th>9/7/2010-8/31/2012</th>
</tr>
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</table>
| 1 Several compounds were synthesized by the SAR of DQ and submitted for in vitro testing against blood stage malaria, specifically P. falciparum D6, W2, C235 and C2B strains, and assessed for metabolic stability in the mouse and human microsomes.
2 In many cases the compounds solubility was improved but the compounds either lost potency against the C2B resistant strain of malaria or microsomal stability.
3 A few interesting trends were discovered. An unprecedented ester replacement, to the ethyl or morpholine amide was discovered that maintained potency against D6,W2,C235 but unfortunately lost activity against the key C2B atovaquone resistant strain.
4 Additionally, many compounds were synthesized that maintained in vitro potency with significantly lower clogP (main focus of the research effort).
5 Efforts are currently toward acquiring a complete set of data to select profile compounds to be scaled up and tested in in vivo models.

|----------------|----------------|
| 1 Intravital Microscopy was successfully employed for investigating EG shedding in hemorrhagic shock/resuscitation for the very first time;
| 2 | Intravital microscopy integrated with systemic hemodynamics evaluations may be essential and more accurate tools to identify changes and study mechanisms of EG shedding and systemic responses to hemorrhage and resuscitation therapy; |
| 3 | Compared to baseline and to the sham group, there was a 50% reduction in endothelial glycocalyx (EG) thickness after hemorrhage and 60% increase in the levels of plasma Syndecan-1; |
| 4 | Although resuscitation with LR and Hextend could stabilize hypotensive rats hemodynamically, these fluids were unable to restore EG thickness or coagulopathy (weak clots and prolonged coagulation time); |
| 5 | Rats who received fresh frozen plasma (FFP) restored venular EG thickness to baseline level in addition to improve the systemic hemodynamics and coagulation response (restored homeostasis). |

**Vecchi, Vittoria**  
**11/14/2011-10/22/2012**

| 1 | Effects of radiation dose reduction on lung quantitative CT scan results in healthy in the Acute Respiratory Distress Syndrome: low-dose chest CT as a valuable tool for quantification and monitoring of pulmonary disease reducing patient exposure |
| 2 | Use of quantitative CT for in vivo lung weight measurement: evaluate and monitor the time course of lung edema in ARDS measuring lung weight by qCT |
| 3 | Pressure-guided positioning of bicaval dual-lumen catheters for veno-venous extracorporeal gas exchange |
| 4 | Low-flow extracorporeal gal exchange for the treatment of ARDS caused by smoke inhalation and cutaneous burn in pigs |
| 5 | Extracorporeal Gas Exchange in awake spontaneously breathing sheep before and after the induction of ARDS |

**Wong, Benjamin**  
**2/7/2011-9/4/2012**

| 1 | Developed and characterized novel system for inhalational exposure of conscious animals to chemical agents |
| 2 | Utilized above system to examine the toxicokinetics (TK) of nerve agents and their analogs in rats |
| 3 | Investigated utility of bronchodilators in a treatment regimen for inhalational chemical agent exposure |
### SUMMARY OF RESEARCH DURING TENURE

Itemize significant findings in concise form, utilizing key concepts/words.

1. Determined the time course of brain derived biomarkers elevated in brain tissue and serum of a rat model of penetrating ballistic-like brain injury (PBBI) (on-going)
2. Determined the time course of brain derived biomarkers elevated in brain tissue and serum of a rat model post-traumatic stress disorder (fear conditioning) and blast injury (on-going)
3. Determined the effect of therapeutic drugs to prevent increases in brain and serum biomarkers after PBBI
4. Initiated proteomics analysis of a rat model post-traumatic stress disorder (PTSD) and blast injury (on going)
5. Grants - Submitted grant proposal to define the markers and mechanism of severe and mild long term brain injury (Combat Casualty Care Program), co-authored pilot grant applications to determine efficacy of therapeutics in PBBI.
6. Publications - Under the guidance of Dr. Dave, our lab published our proteomics study in the journal Electrophoresis. In a separate cohort using proteomics and bioinformatics prediction of biomarkers, I aided in defining novel biomarkers of PBBI in brain tissue that I have independently confirmed. A bioinformatics-themed manuscript is complete and pending submission.

(USMA Davies Fellow: please add summary of teaching, including classes taught.)

N/A

### RESEARCH IN PROGRESS

Describe in no more than 100 words.

Post traumatic stress disorder (PTSD), traumatic brain injuries (TBIs) (e.g., blast overpressure (BOP), concussive, or penetrating) affect thousands military personnel. To successfully identify a panel of protein biomarkers that can (1) predict onset/existence and (2) define mechanisms, I am continuing studies in multiple brain regions and bio-fluids. Initial testing of preparative proteomics protocols was successful; each anatomical region (beginning with cerebral cortex) will be analyzed using proteomics and immunological methods. Biomarker abundance changes in tissues and biofluids after penetrating ballistic-like brain injury (PBBI) are being expanded to include therapeutic and sub-acute effects and we have fully developed studies to define biomarkers of mild, concussive, TBI.

### PUBLICATIONS AND PAPERS RESULTING FROM NRC ASSOCIATESHIP RESEARCH

Provide complete citations: author(s), title, full name of journal, volume number, page number(s), and year of publication.

#### a) Publications in peer-reviewed journals


Angela M Boutté, Changping Yao, Firas Kobaissy, Xi-Chun May Lu, Zhiqun Zhang, Kevin K. Wang, Kara Schmid, Frank C. Tortella and Jitendra R. Dave
b) Books, book chapters, other publications

Protein Biomarkers in Traumatic Brain Injury: An Omics Approach Angela Boutte

c) Manuscripts in preparation, manuscripts submitted

1. SYSTEMS BIOLOGY META-ANALYSES OF GENOMIC DATASETS TO IDENTIFY CONSERVED MECHANISMS AND NOVEL BIOMARKERS OF TRAUMATIC BRAIN INJURY.

2. Time dependant biomarker abundance in brain tissue and serum after acute PBBI

3. Spatial-temporal biomarkers in a model of blast with and without fear conditioning stress in brain tissue and serum after acute and chronic injury

4. The effect of blast injury with and without fear conditioning on the cerebral cortex proteome

10) PATENT OR COPYRIGHT APPLICATIONS RESULTING FROM NRC ASSOCIATESHIP RESEARCH

Provide titles, inventors, and dates of applications.

N/A

11) PRESENTATIONS AT SCIENTIFIC MEETINGS OR CONFERENCES

Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.

International

N/A

Domestic


12) SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES

Include dates, names and locations of seminars.


13) PROFESSIONAL AWARDS RECEIVED DURING TENURE

N/A

14) POST-TENURE POSITION / JOB TITLE

Research Biologist

15) NAME AND ADDRESS OF POST-TENURE POSITION / JOB ORGANIZATION

Walter Reed Army Inst. For Research

16) POST-TENURE POSITION STATUS / CATEGORY Please indicate only one.

☒ Permanent position at the NRC host agency

☐ Contract or temporary position at the NRC host Agency

Abbreviate Host Laboratory/Center

☐ Research/Administrative position with another U.S.-government agency

☐ Research/Administrative position with a foreign-government agency

☐ Research/teaching position at a U.S. college or university

☐ Research/teaching position at a foreign college or university

☒ Research/administration position in private industry in the U.S.

☐ Research/administration position in private industry outside of the U.S.

☐ Research/administration position with a non profit

☐ Self-employed/consulting

☐ Postdoctoral research

☐ Other (Please specify, possible)

☐ No information provided

17) APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM

On a scale of 1 – 10 (poor - excellent), please rate the following:

SHORT TERM VALUE

☒ Development of knowledge, skills, and research productivity
LONG TERM VALUE
10 How the NRC Associateship award affected your career to date

LAB SUPPORT
9 Quality of support from the Laboratory—equipment, funding, orientation, safety and health guidelines, etc.

ADVISER/MENTOR SUPPORT
10 Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)

LPR SUPPORT
10 Quality of administrative support from the Laboratory (e.g., NIST, NRL, IWR, FHWA) NRC Program Representative (LPR)

NRC SUPPORT
10 Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)

18) PLEASE PROVIDE ANY SUGGESTIONS FOR PROGRAM IMPROVEMENT.

Please do NOT scan to PDF. Send the Final Report as MSWord document via e-mail to your NRC Program Coordinator

No handwritten signature required; but you may upload a scanned signature file below:

Maria Crocco: mcrocco@nas.edu
Asha Davis: adavis@nas.edu
Linda Sligh: lsligh@nas.edu
Jason Thornhill: jthornhill@nas.edu
Peggy Wilson: pwilson@nas.edu

Id# Rev. Jan 2013 Proj/Act ID#
**FINAL REPORT**

1) **Associate Last or Family Name**

Chen

2) **First Name**

Guojun

3) **M.I.**

4) **Today's Date**

April 18, 2012

4) **Dates of Tenure**

from 08-08-2011 to August 8, 2012

4) **Host Agency**

USAMC

(e.g., AFRL)

4) **Laboratory or Center**

(e.g., Wright Patterson AFB)

4) **Division / Directorate / Department**

(e.g., High-Speed Propulsion)

5) **Name of Laboratory NRC Adviser (and USMA Mentor, if applicable)**

David. Devore

6) **TITLE OF RESEARCH PROPOSAL**

7) **SUMMARY OF RESEARCH DURING TENURE**

Itemize significant findings in concise form, utilizing key concepts/words.

1) virus-like particle for antisense ODN delivery

2) peptide-oligonucleic acid conjugates for antisense

3) Co-polymer enhanced liposome for antisense ODN delivery

4)

5)

(USMA Davies Fellow: please add summary of teaching, including classes taught.)

8) **RESEARCH IN PROGRESS**

Describe in no more than 100 words.

Virus-like particle project was stopped due to the lack of fresh plant infected tissue. Therefore, two extra backup plans were carried out. One is peptide-oligonucleic acid (PNA) and the other one is liposome delivery system. For PNA project, conjugation are being synthesized and characterized. The preliminary bacteria tests have indicated that PNA can improve the antisense antimicrobial capability. For copolymer enhanced liposome, the copolymers were synthesized. However, the complexing the biological testing are not carried out yet.

9) **PUBLICATIONS AND PAPERS RESULTING FROM NRC ASSOCIATESHIP RESEARCH**

Provide complete citations: author(s), title, full name of journal, volume number, page number(s), and year of publication.

a) Publications in peer-reviewed journals

b) Books, book chapters, other publications

c) Manuscripts in preparation, manuscripts submitted

Peptide-oligonucleic acid for mecA antisense in S.aureus

10) **PATENT OR COPYRIGHT APPLICATIONS RESULTING FROM NRC ASSOCIATESHIP RESEARCH**

Provide titles, inventors, and dates of applications.

11) **PRESENTATIONS AT SCIENTIFIC MEETINGS OR CONFERENCES**

Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.

International
12) SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES  Include dates, names and locations of seminars.

13) PROFESSIONAL AWARDS RECEIVED DURING TENURE

14) POST-TENURE POSITION / JOB TITLE

15) NAME AND ADDRESS OF POST-TENURE POSITION / JOB ORGANIZATION

Emory/Atlanta

16) POST-TENURE POSITION STATUS / CATEGORY  Please indicate only one.

- □ Permanent position at the NRC host agency
- □ Contract or temporary position at the NRC host Agency
- □ Research/Administrative position at a foreign government agency
- □ Research/Administrative position at a foreign college or university
- □ Research/Administration position in private industry in the U.S.
- □ Research/Administration position in private industry outside of the U.S.
- □ Research/administration position with a non-profit
- □ Self-employed/consulting
- □ Postdoctoral research
- □ Other (Please specify, possible) ______
- □ No information provided

17) APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM

On a scale of 1 – 10 (poor - excellent), please rate the following:

SHORT TERM VALUE

- Development of knowledge, skills, and research productivity
- Comments

LONG TERM VALUE

- How the NRC Associateship award affected your career to date
- Comments

LAB SUPPORT

- Quality of support from the Laboratory—equipment, funding, orientation, safety and health guidelines, etc.
- Comments

ADVISER/MENTOR SUPPORT

- Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)
- Comments

LPR SUPPORT

- Quality of administrative support from the Laboratory (e.g., NIST, NRL, IWR, FHWA) NRC Program Representative (LPR)
- Comments

NRC SUPPORT

- Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)
- Comments

18) PLEASE PROVIDE ANY SUGGESTIONS FOR PROGRAM IMPROVEMENT.

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No handwritten signature required; 
Asha Davis: adavis@nas.edu
but you may upload a scanned Linda Sligh: bsligh@nas.edu
signature file below:

<table>
<thead>
<tr>
<th>Id#</th>
<th>Rev. July 2011</th>
<th>Proj/Act ID#</th>
</tr>
</thead>
<tbody>
<tr>
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</tbody>
</table>

Jason Thornhill:  jthornhill@nas.edu
Peggy Wilson:     pwilson@nas.edu
Suzanne White:    swhite@nas.edu
**FINAL REPORT**

1) **Associate Last or Family Name**
   - Johnston

2) **First Name**
   - Sara

3) **M.I.**
   - C

4) **Today's Date**
   - April 26, 2012

5) **Dates of Tenure**
   - from March 25, 2009 to April 30, 2012

6) **Host Agency**
   - USAMRIID

7) **Laboratory or Center**
   - Fort Detrick Army Garrison

8) **Division / Directorate / Department**
   - Virology Division

9) **Name of Laboratory NRC Adviser (and USMA Mentor, if applicable)**
   - Dr. Arthur Goff

10) **TITLE OF RESEARCH PROPOSAL**
    - Identification and Characterization of Viral Immunomodulators that Affect the Host Specificity of Orthopoxviruses

11) **SUMMARY OF RESEARCH DURING TENURE**
    - Itemize significant findings in concise form, utilizing key concepts/words.

   1) MPXV Active Disease Surveillance Program identifying significant increases in prevalence in DRC
   2) Identification of potential novel therapeutic (IFN-beta) against MPXV
   3) Identification of MPXV variants actively circulating in the DRC
   4) In vitro and in vivo evaluation of novel therapeutics against Henipaviruses
   5) Development and implementation of diagnostic ELISA assays against MPXV

   (USMA Davies Fellow: please add summary of teaching, including classes taught.)

12) **RESEARCH IN PROGRESS**
    - Describe in no more than 100 words.

   Continued efforts associated with MPXV active disease surveillance which are ongoing, in vivo characterization of IFN-beta against MPXV, continued therapeutic testing against Henipaviruses.

13) **PUBLICATIONS AND PAPERS RESULTING FROM NRC ASSOCIATESHIP RESEARCH**
    - Provide complete citations: author(s), title, full name of journal, volume number, page number(s), and year of publication.

   a) Publications in peer-reviewed journals


      "Assessment of high-throughput screening (HTS) methods for high-consequence pathogens" Brian M Friedrich, Corinne E Scully, Jennifer M Brannan, Monica M Ogg, Sara C Johnston, Lisa E Hensley, Gene G Olinger, and Darci R Smith: Bioterrorism & Biodefense 2011, S3


      "Major increase in human monkeypox incidence 30 years after smallpox vaccination campaigns cease in the Democratic Republic of Congo" Anne W Rimoin, Prime M Mulembakani, Sara C Johnston, James O Lloyd Smith, Neville K Kisalu, Timothee L Kinkela, Seth Blumberg, Henri A Thomassen, Brian L Pike, Joseph N Fair, Nathan D Wolfe, Robert L
b) Books, book chapters, other publications

c) Manuscripts in preparation, manuscripts submitted

- “Risk Factors for Human Monkeypox in the Democratic Republic of the Congo”
  Manuscript in Review (Emerging Infectious Diseases).
- “Identification of Genomic Destabilization in Monkeypox Clinical Samples”
  Manuscript in Preparation (Science)
- “Pathogen-host Associations and Range Shifts of Human Monkeypox in Response to Climate Change in Central Africa” Manuscript in Preparation (PLOS One).

10) PATENT OR COPYRIGHT APPLICATIONS RESULTING FROM NRC ASSOCIATESHIP RESEARCH

Provide titles, inventors, and dates of applications.

11) PRESENTATIONS AT SCIENTIFIC MEETINGS OR CONFERENCES

Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.

International

- “A Major Increase in the Incidence of Human Monkeypox Thirty Years After Smallpox Vaccination Campaigns Cease in the Democratic Republic of Congo”
  Oral presentation at the 11th Meeting of the WHO Advisory Committee on Variola Virus, Geneva, Switzerland
  November 2009

Domestic

- “Human Monkeypox Emergence Since the Cessation of Global Smallpox Vaccination”
  Oral presentation at the Pennsylvania Vector Control Association Training Conference, State College, PA
  November 2011
- “Evaluation of Rash Illness in DRC Using Pan-Orthopox, Monkeypox Specific, and VZV Specific Assays”
  Poster presentation at the Chemical and Biological Defense Science and Technology Conference, Orlando, FL
  November 2010
- “Major Increase in Human Monkeypox Incidence Thirty Years After Smallpox Vaccination Campaigns Cease in the DRC”
  Poster presentation at the XVIII International Poxvirus, Asfivirus, and Iridovirus Symposium, Sedona, AZ
  June 2010
- “Vaccinia Virus E3L Blocks the Formation of Stress Granule (SG)-Related “Factory Granules” that Inhibit the Replication of a ΔE3L Virus”
  Oral presentation at the XVIII International Poxvirus, Asfivirus, and Iridovirus Symposium, Sedona, AZ
  June 2010
- “The Actin Motor Myosin V Associates with Intracellular Enveloped Virions”
  Oral presentation at the XVIII International Poxvirus, Asfivirus, and Iridovirus Symposium, Sedona, AZ
  June 2010
- “Human Monkeypox Genomic Divergence and Determinants of Pathogenesis”
  Poster presentation at the 58th annual meeting of the American Society of Tropical Medicine
  Washington, DC
  November 2009

12) SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES

Include dates, names and locations of seminars.

"The Development of Antiviral Strategies to Combat Emerging/Re-Emerging Pathogens"
  Oral presentation at Tufts University Cummings School of Veterinary Medicine
  North Grafton, MA
  April 2012

13) PROFESSIONAL AWARDS RECEIVED DURING TENURE
14) POST-TENURE POSITION / JOB TITLE

Research Microbiologist 4 contracted through ClinRM

15) NAME AND ADDRESS OF POST-TENURE POSITION / JOB ORGANIZATION

USAMRIID, 1425 Porter St. Fort Detrick, MD 21702

16) POST-TENURE POSITION / STATUS / CATEGORY  Please indicate only one.

☐ Permanent position at the NRC host agency
☐ Contract or temporary position at the NRC host Agency
☐ Research/Administrative position with another U.S.-government agency
☐ Research/Administrative position with a foreign-government agency
☐ Research/teaching position at a U.S. college or university
☐ Research/teaching position at a foreign college or university
☐ Research/administration position in private industry in the U.S.
☐ Research/administration position in private industry outside of the U.S.
☐ Research/administration position with a non profit
☐ Self-employed/consulting
☐ Postdoctoral research
☐ Other (Please specify, possible) _____
☐ No information provided

17) APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM

On a scale of 1 – 10 (poor - excellent), please rate the following:

SHORT TERM VALUE
☐ Development of knowledge, skills, and research productivity
 Comments

LONG TERM VALUE
☐ How the NRC Associateship award affected your career to date
 Comments

LAB SUPPORT
☐ Quality of support from the Laboratory--equipment, funding, orientation, safety and health guidelines, etc.
 Comments

ADVISER/MENTOR SUPPORT
☐ Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)
 Comments

LPR SUPPORT
☐ Quality of administrative support from the Laboratory (e.g., NIST, NRL, IWR, FHWA) NRC Program Representative (LPR)
 Comments

NRC SUPPORT
☐ Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)
 Comments

18) PLEASE PROVIDE ANY SUGGESTIONS FOR PROGRAM IMPROVEMENT.

Please do NOT scan to PDF. Send the Final Report as MSWord document via e-mail to your NRC Program Coordinator

No handwritten signature required; but you may upload a scanned signature file below:

Asha Davis: adavis@nas.edu
Linda Sligh: lsligh@nas.edu
Jason Thornhill: jthornhill@nas.edu
Peggy Wilson: pwilson@nas.edu
Suzanne White: swhite@nas.edu

Id# Rev. July 2011 Proj/Act ID#
1) Associate Last or Family Name  |  First Name  |  M.I.
Langer  |  Thomas  |  

3) Today's Date  |  Dates of Tenure

4) Host Agency  |  Laboratory or Center  |  Division / Directorate / Department
AMRMC (e.g., AFRL)  |  USA ISR (e.g., Wright Patterson AFB)  |  (e.g., High-Speed Propulsion)

5) Name of Laboratory NRC Adviser (and USMA Mentor, if applicable)
Andriy Batchinsky and Leopoldo Cancio

6) TITLE OF RESEARCH PROPOSAL
PHYSIOLOGY AND PATHOPHYSIOLOGY OF SPONTANEOUS BREATHING DURING TOTAL EXTRACORPOREAL RESPIRATORY SUPPORT IN HEALTHY SHEEP AND IN SHEEP WITH THE ACUTE RESPIRATORY DISTRESS SYNDROME

7) SUMMARY OF RESEARCH DURING TENURE
Itemize significant findings in concise form, utilizing key concepts/words.
1) Development of a large animal model to study spontaneous breathing during extracorporeal gas exchange
2) Study of differences in respiratory pattern between healthy animals and animals with the Acute Respiratory Distress Syndrome
3) Development of a new technique for the placement of bicaval dual-lumen catheters for venovenous extracorporeal gas exchange
4) Effects of radiation dose reduction on lung quantitative CT scan results in healthy in the Acute Respiratory Distress Syndrome: low-dose chest CT as a valuable tool for quantification and monitoring of pulmonary disease reducing patient exposure
5) (USMA Davies Fellow: please add summary of teaching, including classes taught.)

8) RESEARCH IN PROGRESS
Describe in no more than 100 words.

Mechanical ventilation (MV) is the current standard of care for the treatment of the acute respiratory distress syndrome. MV can however worsen lung injury. Extracorporeal Gas Exchange is a tempting alternative to treat ARDS. We have established a model to start investigating this treatment option in order to be able to provide, in the future, a safe and successful treatment to patients with ARDS.

9) PUBLICATIONS AND PAPERS RESULTING FROM NRC ASSOCIATESHIP RESEARCH
Provide complete citations: author(s), title, full name of journal, volume number, page number(s), and year of publication.

a) Publications in peer-reviewed journals

b) Books, book chapters, other publications
0

c) Manuscripts in preparation, manuscripts submitted
Extracorporeal Gas Exchange in awake spontaneously breathing sheep before and after the induction of ARDS - manuscript in preparation
10) **PATENT OR COPYRIGHT APPLICATIONS RESULTING FROM NRC ASSOCIATESHIP RESEARCH**

Provide titles, inventors, and dates of applications.

N/A

11) **PRESENTATIONS AT SCIENTIFIC MEETINGS OR CONFERENCES**

Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.

**International**


**Domestic**


12) **SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES**

Include dates, names and locations of seminars.

0

13) **PROFESSIONAL AWARDS RECEIVED DURING TENURE**

0

14) **POST-TENURE POSITION / JOB TITLE**

Intensive Care Medicine Resident

15) **NAME AND ADDRESS OF POST-TENURE POSITION / JOB ORGANIZATION**

Istituto di Anestesia e Rianimazione, Università degli Studi di Milano, Milan, Italy

16) **POST-TENURE POSITION STATUS / CATEGORY**

Please indicate only one.

- [ ] Permanent position at the NRC host agency
- [ ] Contract or temporary position at the NRC host Agency (Abbreviate Host Laboratory/Center)
- [ ] Research/Administrative position with another U.S.-government agency
- [ ] Research/Administrative position with a foreign-government agency
- [ ] Research/teaching position at a U.S. college or university
- [X] Research/teaching position at a foreign college or university
- [ ] Research/administration position in private industry in the U.S.
- [ ] Research/administration position in private industry outside of the U.S.
- [ ] Research/administration position with a non profit
- [ ] Self-employed/consulting
- [ ] Postdoctoral research
- [ ] Other (Please specify, possible)
- [ ] No information provided

17) **APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM**

On a scale of 1 – 10 (poor - excellent), please rate the following:

**SHORT TERM VALUE**

- [8] Development of knowledge, skills, and research productivity
  
  Comments: Great opportunity to develop research skills

**LONG TERM VALUE**

- [8] How the NRC Associateship award affected your career to date
  
  Comments

**LAB SUPPORT**

- [7] Quality of support from the Laboratory--equipment, funding, orientation, safety and health guidelines, etc.
  
  Comments

**ADVISER/MENTOR SUPPORT**

- [8] Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)
  
  Comments

**LPR SUPPORT**

- [8] Quality of administrative support from the Laboratory (e.g., NIST, NRL, IWR, FHWA) NRC Program Representative (LPR)
Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)

Really great support. Thank you Jason and Peggy!

Please do NOT scan to PDF. Send the Final Report as MSWord document via e-mail to your NRC Program Coordinator.

No handwritten signature required; but you may upload a scanned signature file below:

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Asha Davis: adavis@nas.edu
Linda Sligh: bsligh@nas.edu
Jason Thornhill: jthornhill@nas.edu
Peggy Wilson: pwilson@nas.edu
Suzanne White: swhite@nas.edu
CHARACTERIZING A CLINICALLY/MILITARILY RELEVANT RAT MODEL OF POLYTRAUMA ASSOCIATED WITH PENETRATING BRAIN INJURY: NEUROPATHOLOGICAL EFFECTS OF TRANSIENT HYPOTENSION AND/OR ACUTE HYPOXEMIA AFTER PENETRATING BALLISTIC-LIKE BRAIN INJURY (PBBI)

1) ESTABLISHED POLYTRAUMA MODELS ASSOCIATED WITH HYPOXEMIA AND HEMORRHAGIC HYPOTENSION THAT WILL BE USED FOR FUTURE NEUROPROTECTIVE DRUG STUDIES.

2) ACUTE PHYSIOLOGICAL CHANGES WERE CHARACTERIZED IN THE POLYTRAUMA MODELS. THE PATTERNS OF THESE CHANGES WERE FOUND TO BE UNIQUE UNDER DIFFERENT INJURY COMBINATIONS.

3) HEMORRHAGIC SHOCK INCREASED THE INCIDENCE AND DURATION OF CORTICAL SPREADING DEPOLARIZATION WITHIN 2 HOURS FOLLOWING PBBI WHEREAS HYPOXEMIA only prolonged the depolarization.

4) HISTOPATHOLOGICAL CHANGES WERE CHARACTERIZED (3, 7 DAYS POST-INJURY) IN THE POLYTRAUMA MODELS. HEMORRHAGIC SHOCK INCREASED NEURONAL DEGENERATION AND ASTROCYTIC ACTIVATION FOLLOWING PBBI.

5) THE SEQUENCE OF PBBI, HEMORRHAGIC HYPOTENSION AND HYPOXEMIA AFFECTED THE MORTALITY RATE AND NEUROLOGICAL DEFICITS. PBBI, HS FOLLOWED BY HX RESULTED IN THE HIGHEST MORTALITY RATE (50%) AND MORE NEUROLOGICAL DEFICITS AMONG ALL POLYTRAUMA GROUPS.

RESEARCH IN PROGRESS

Part of the histopathological characterization is underway. Brain samples collected following the different sequences of polytrauma are being processed for immunostaining. The quantifications of histopathological changes will be performed in these immunotained brain slices. The behavioral and EEG studies in the polytrauma models will soon be started. I have submitted a core funding proposal (FY14-18) as a principal investigator to Combat Casualty Care Research Program of MRMC to continue the research efforts on polytrauma. The proposal aims at investigating the effects of polytrauma on protein biomarkers, the vital organs, etc. as well as polytrauma associated with mild TBI.

PUBLICATIONS AND PAPERS RESULTING FROM NRC ASSOCIATESHIP RESEARCH


Leung LY, Wei G, Shear DA, Tortella FC. The acute effect of hemorrhagic shock on cerebral blood flow, brain tissue oxygen tension and spreading depolarization following penetrating ballistic-like brain injury. Journal of Neurotrauma. (Final revision)


b) Books, book chapters, other publications

c) Manuscripts in preparation, manuscripts submitted

Leung LY, Wei G, Shear DA, Tortella FC. Temporal and Spatial Profile of Histopathological Changes Caused by Hemorrhagic Shock in a Rat Model of Penetrating Ballistic-like Brain Injury (PBBI). (In preparation)

10) PATENT OR COPYRIGHT APPLICATIONS RESULTING FROM NRC ASSOCIATESHIP RESEARCH
Provide titles, inventors, and dates of applications.
N/A

11) PRESENTATIONS AT SCIENTIFIC MEETINGS OR CONFERENCES
Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.

International
N/A

Domestic


12) SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES Include dates, names and locations of seminars.

8 November, 2011. CNS Injury Conference, hosted by Center for Brain Injury and Repair, University of Pennsylvania.

13) PROFESSIONAL AWARDS RECEIVED DURING TENURE
N/A

14) POST-TENURE POSITION / JOB TITLE
Neurobiologist

15) NAME AND ADDRESS OF POST-TENURE POSITION / JOB ORGANIZATION

Walter Reed Army Institute of Research
503 Robert Grant Avenue, 2W12
Silver Spring, Maryland

16) POST-TENURE POSITION STATUS / CATEGORY

Please indicate only one.

- Permanent position at the NRC host agency
- Contract or temporary position at the NRC host Agency
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- Research/administration position in private industry outside of the U.S.
- Research/administration position with a non profit
- Self-employed/consulting
- Postdoctoral research
- Other (Please specify, possible)
- No information provided

17) APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM

On a scale of 1 – 10 (poor - excellent), please rate the following:

SHORT TERM VALUE

10 Development of knowledge, skills, and research productivity
Comments

LONG TERM VALUE

10 How the NRC Associateship award affected your career to date
Comments

LAB SUPPORT

10 Quality of support from the Laboratory--equipment, funding, orientation, safety and health guidelines, etc.
Comments

ADVISER/MENTOR SUPPORT

10 Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)
Comments

LPR SUPPORT

10 Quality of administrative support from the Laboratory (e.g., NIST, NRL, IWR, FHWA) NRC Program Representative (LPR)
Comments

NRC SUPPORT

10 Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)
Comments

18) PLEASE PROVIDE ANY SUGGESTIONS FOR PROGRAM IMPROVEMENT.

Please do NOT scan to PDF. Send the Final Report as MSWord document via e-mail to your NRC Program Coordinator

No handwritten signature required; but you may upload a scanned signature file below:

Asha Davis: adavis@nas.edu
Linda Shlig: bsligh@nas.edu
Jason Thornhill: jthornhill@nas.edu
Peggy Wilson: pwilson@nas.edu
Suzanne White: swhite@nas.edu

Id# Rev. July 2011 Proj/Act ID#
**FINAL REPORT**

1) **Associate Last or Family Name**

McCoy

2) **First Name**

Margaret

3) **M.I.**

E

4) **Today's Date**

September 4, 2012

5) **Dates of Tenure**

from July 29, 2012  

to October 4, 2012

6) **Host Agency**

WRAIR (e.g., AFRL)

**Laboratory or Center**

David E. Lanar (e.g., Wright Patterson AFB)

**Division / Directorate / Department**

Malaria Vaccine Branch (e.g., High-Speed Propulsion)

7) **Name of Laboratory NRC Adviser (and USMA Mentor, if applicable)**

David. E. Lanar  N/A

8) **TITLE OF RESEARCH PROPOSAL**

Examination of immune responses and efficacy in mouse and Rhesus animal models of a circumsporozoite proteine (CSP)-based Self-Assembling Polypeptide Nanoparticle (SAPN) malaria vaccine.

9) **SUMMARY OF RESEARCH DURING TENURE**

Itemize significant findings in concise form, utilizing key concepts/words.

1) Identified the mechanism of action of SAPN-induced Ab that provides sterile immunity in mice

2) designed and carried out experiments to examine and characterize the processing and presentation of SAPN within the immune system

3) Examined the phenotypes of SAPN-specific T-cell populations and proved that CD8+ T-cells from mice immunized with SAPN are able to, by themselves, induce sterile immunity in mice- this is the first malaria vaccine to be able to show this.

4) Examined needleless approaches to vaccine delivery, including pleuronic lecithin organogel creams

5) Identified potential effects on vaccine efficacy resulting from the addition of mosquito saliva

6) The youngest investigator at the WRAIR to successfully write and complete a non-human primate trial for malaria

(USMA Davies Fellow: please add summary of teaching, including classes taught.)

N/A

10) **RESEARCH IN PROGRESS**

Describe in no more than 100 words.

I have successfully completed all of my murine and Rhesus experiments with only minor finishing touches to be wrapped up.

11) **PUBLICATIONS AND PAPERS RESULTING FROM NRC ASSOCIATESHIP RESEARCH**

Provide complete citations: author(s), title, full name of journal, volume number, page number(s), and year of publication.

a) Publications in peer-reviewed journals

1) Protective antibody and CD8+ T-cell responses to the Plasmodium falciparum protein induced by a polypeptide nanoparticle vaccine. Stephen Kaba, Margaret E. McCoy, Tais Doll, Peter Burkhard, Qin Guo, Debleena Dasgupta, Yongkun Yang, Christian Mittelholzer, Roberta Spaccapelo, Andrea Crisanti, and David E. Lanar. in press PLoS ONE September 2012

b) Books, book chapters, other publications

N/A

c) Manuscripts in preparation, manuscripts submitted


10) PATENT OR COPYRIGHT APPLICATIONS RESULTING FROM NRC ASSOCIATESHIP RESEARCH
Provide titles, inventors, and dates of applications.
N/A

11) PRESENTATIONS AT SCIENTIFIC MEETINGS OR CONFERENCES
Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.

International

Domestic
1) Cross-presentation of exogenous peptides delivered in self-assembling polypeptide nanoparticles: Implications for malaria vaccine development; Margaret E. McCoy and David E. Lanar. WRAIR, Young Investigator's Meeting. Silver Spring, MD. September 2009.
6) A nanoparticle vaccine targeting plasmodium falciparum circumsporozoite protein confers protective humoral and cellular immunity; Margaret E. McCoy, SA Kaba, Q Guo, D Dasgupta, TAPF Doll, Y Yang, C Mittelholzer, R Spaccapelo, A Crisanti, P Burkhard and D Lanar. 61st Annual Meeting, ASTMH, Philadelphia, PA. December 4-8 2011.

12) SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES
Include dates, names and locations of seminars.
1) Lectured at Blari High School, Silver Spring, MD 2011 and 2012
2) Lectured at Bishop O'Connell High School, Arlington, VA 2011

13) PROFESSIONAL AWARDS RECEIVED DURING TENURE
1) Young Investigator Award, 2nd runner-up, ASTMH December 2011
2) The Maurice R. Hilleman Early Stage Career Investigator Award, Runner-up, the National Foundation for Infectious Diseases, April 27, 2010

14) POST-TENURE POSITION / JOB TITLE
Law student, the University of Texas at Austin

15) NAME AND ADDRESS OF POST-TENURE POSITION / JOB ORGANIZATION
Austin, TX

16) POST-TENURE POSITION STATUS / CATEGORY
Please indicate only one.
☐ Permanent position at the NRC host agency
☐ Contract or temporary position at the NRC host Agency
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☐ Research/administration position at a foreign college or university
☐ Research/administration position in private industry in the U.S.
☐ Research/administration position in private industry outside of the U.S.
☐ Research/administration position with a non profit
☐ Self-employed/consulting
☐ Postdoctoral research
☐ Other (Please specify, possible) Law School
17) APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM

On a scale of 1 – 10 (poor - excellent), please rate the following:

SHORT TERM VALUE

10 Development of knowledge, skills, and research productivity

Comments
This position has truly expanded my knowledge, experience and research focusing abilities.

LONG TERM VALUE

10 How the NRC Associateship award affected your career to date

Comments
This fellowship has set me apart and provided me with an outstanding base from which to launch the next phase of my career.

LAB SUPPORT

10 Quality of support from the Laboratory--equipment, funding, orientation, safety and health guidelines, etc.

Comments
Dr. Lanar financially supported me in more than one endeavour of mine in the lab and has encouraged my independent thoughts on experimental designs and writing.

ADVISER/MENTOR_SUPPORT

10 Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)

Comments
Dr. Lanar financially supported me in more than one endeavour of mine in the lab and has encouraged my independent thoughts on experimental designs and writing.

LPR SUPPORT

10 Quality of administrative support from the Laboratory (e.g., NIST, NRL, IWR, FHWA) NRC Program Representative (LPR)

Comments
This position just recently changed hands, but I have to say that Dr. Sara Rothman is the most amazing mentor and support for all of us. She is an icon and a role model to which any after her can only desire to imitate.

NRC SUPPORT

8 Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)

Comments
I had several major issues with payroll. Omegatravel and Jason and his crew (Ms. Winstead included) have been wonderful and very helpful whenever I needed their assistance!!

18) PLEASE PROVIDE ANY SUGGESTIONS FOR PROGRAM IMPROVEMENT.

N/A

Please do NOT scan to PDF. Send the Final Report as MSWord document via e-mail to your NRC Program Coordinator

No handwritten signature required; but you may upload a scanned signature file below:

Asha Davis: adavis@nas.edu
Linda Sligh: lsligh@nas.edu
Jason Thornhill: jthornhill@nas.edu
Peggy Wilson: pwilson@nas.edu
Suzanne White: swhite@nas.edu

Id# Rev. July 2011 Proj/Act ID#
A strategy for the control of acute coagulopathy of trauma

1) Associate Last or Family Name: Meledeo
2) First Name: Michael
3) M.I.: A
4) Today's Date: July 6, 2012
5) Dates of Tenure: from March 1, 2010 to July 6, 2012
6) Host Agency: AMRMC (e.g., AFRL)
7) Laboratory or Center: ISR (e.g., Wright Patterson AFB)
8) Division / Directorate / Department: Damage Control Resuscitation (e.g., High-Speed Propulsion)
9) Name of Laboratory NRC Adviser (and USMA Mentor, if applicable): Bowman Philip

7) SUMMARY OF RESEARCH DURING TENURE

1) Aptamers can be used to completely inhibit the anti-coagulant effects of activated protein C (aPC); however, aPC does not appear to be the sole sufficient cause of the acute coagulopathy of trauma.

2) Exposing in vitro cultures of endothelial cells (ECs) to laminar flow (as a model of their physiological environment) will induce a number of changes to both EC morphology and gene expression in a variety of inflammatory and morphology pathways.

3) While not all of the changes in gene expression result in an altered proteome, there are a number of significant differences in protein expression between static cultured ECs and those exposed to flow.

4) An analysis of the EC glycocalyx through confocal microscopy and western blotting of membrane proteins and associated glycoforms has led to an enhanced understanding of the structure and function of the endothelial glycocalyx layer.

5) All of these have provided advancement in the formulation of in vitro models of endothelium; in the future it should be possible to use these models as a platform for the testing of both therapeutics and diagnostics for vascular dysfunction.

8) RESEARCH IN PROGRESS

The research in progress will continue over the next few months since I will be working just down the hall from my NRC position. This includes a more thorough analysis of the proteomic effects of laminar flow on the endothelial cells through a small grant with the UTSA Proteomics Core. There is a little more work to be done with the confocal analysis of ECs particularly in a time course study of the formation of the EGL. Finally, only recently has the method for isolating and concentrating the membrane proteins for the purposes of identifying enhancement or inhibition of EGL members; more work can quickly be completed on identifying the affected members through western blotting.

9) PUBLICATIONS AND PAPERS RESULTING FROM NRC ASSOCIATESHIP RESEARCH

a) Publications in peer-reviewed journals


b) Books, book chapters, other publications

n/a

c) Manuscripts in preparation, manuscripts submitted


10) PATENT OR COPYRIGHT APPLICATIONS RESULTING FROM NRC ASSOCIATESHIP RESEARCH
Provide titles, inventors, and dates of applications.


11) PRESENTATIONS AT SCIENTIFIC MEETINGS OR CONFERENCES
Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.

International
n/a

Domestic


12) SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES
Include dates, names and locations of seminars.


13) PROFESSIONAL AWARDS RECEIVED DURING TENURE

14) POST-TENURE POSITION / JOB TITLE
Staff Scientist, Contractor with Cherokee Nation Tech.

15) NAME AND ADDRESS OF POST-TENURE POSITION / ORGANIZATION

USAISR Blood Research Group
3650 Chambers Pass, Bldg 3610
Ft. Sam Houston, TX 78234

16) POST-TENURE POSITION STATUS / CATEGORY

Please indicate only one.

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- Research/administration position in private industry in the U.S.
- Research/administration position in private industry outside of the U.S.
- Research/administration position with a non profit
- Self-employed/consulting
- Postdoctoral research
- Other (Please specify, possible)
- No information provided

17) APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM
On a scale of 1 – 10 (poor - excellent), please rate the following:

- Development of knowledge, skills, and research productivity

Comments
I used many things I already knew how to do in this associateship, but I did have the opportunity to learn more about WHY we do the things we do; additionally, I had the opportunity to learn several new things. Productivity was not consistently stressed.
LONG TERM VALUE

How the NRC Associateship award affected your career to date

Comments

Although from a scientific performance metric (that being the number of publications produced), this position was not as good for my career as it could have been (a large part of that was my own fault), it did afford me the opportunity to show my skills and efforts to others in the host institute. This directly led to my new position, and hopefully to further career advancement in the future.

LAB SUPPORT

Quality of support from the Laboratory—equipment, funding, orientation, safety and health guidelines, etc.

Comments

Equipment and funding are unsurpassed in my experience. We had everything we needed. Safety and irrelevant training were overstressed, which is not surprising considering that those programs have to be tailored to the least common denominator.

ADVISER/MENTOR SUPPORT

Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)

Comments

Dr. Bowman is one of the most knowledgable people I’ve ever known, and he was an invaluable resource for any questions I had. He taught me a great deal about the research subjects on which I was working. My only complaints are that his expectations for my work were rarely known to me.

LPR SUPPORT

Quality of administrative support from the Laboratory (e.g., NIST, NRL, IWR, FHWA) NRC Program Representative (LPR)

Comments

Dr. Dubick was not frequently needed or seen. He did travel a lot which made it difficult to find him, although that was a rare necessity.

NRC SUPPORT

Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)

Comments

Administrative support was excellent. I had several problems with the travel group, but I understand that there was a lot of turmoil in that department with various people covering roles while some were out on leave. Everything worked out in the end.

18) PLEASE PROVIDE ANY SUGGESTIONS FOR PROGRAM IMPROVEMENT.

My primary suggestion would be to identify to both the mentors and the associates what the NRC's expectations are for roles and performance markers. I had certain expectations (my own problem) which were not met, and by the time I realized that they were not going to be met, I felt like it was too late to say anything to anyone about it.

Please do NOT scan to PDF. Send the Final Report as MSWord document via e-mail to your NRC Program Coordinator

No handwritten signature required; but you may upload a scanned signature file below:

Asha Davis: adavis@nas.edu
Linda Sligh: hsligh@nas.edu
Jason Thornhill: jthornhill@nas.edu
Peggy Wilson: pwilson@nas.edu
Suzanne White: swhite@nas.edu

Id# Rev. July 2011 Proj/Act ID#
Dengue Virus Quasispecies' role in Viral Fitness and Adaptation to Changing Environments

1) Variants analysis of dengue quasispecies populations showed that variants are host or vector specific despite containing the same consensus sequence. Diversity was found to not be constrained within the vector as suggested in some publications.

2) Selection analysis showed the populations to be expanding, evolving at a faster rate than 'average' for the dataset, and were under selective pressure with a predominance of nonsynonymous mutations when compared with the consensus sequence.

3) Phylogenetic analysis revealed that dengue quasispecies sequences isolated in 2010 were distinct from other circulating consensus sequences from Thailand and full E gene offered higher resolution than partial E gene sequences.

4) Amino acid (aa) analysis suggested several positions where changes would affect replication, antibody binding or VLP assembly according to the literature. Multivariate analysis predicted uncharacterized aa positions that would have impact if altered.

5) This work revealed the importance of full E gene surveillance for assessment of aa changes, illustrated the variability and pathogenic potential of dengue quasispecies variant diversity and established a baseline in which to make future comparisons.

Future work will focus on analyzing quasispecies population dynamics over the course of an in vitro transmission cycle and characterizing changes within the population during the course of fever prior to defervescence. The merits of using cloning versus next generation sequencing technologies to assist in both of these projects will also be assessed.


10) PATENT OR COPYRIGHT APPLICATIONS RESULTING FROM NRC ASSOCIATESHIP RESEARCH
Provide titles, inventors, and dates of applications.

11) PRESENTATIONS AT SCIENTIFIC MEETINGS OR CONFERENCES
Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.
International

12) SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES
Include dates, names and locations of seminars.

13) PROFESSIONAL AWARDS RECEIVED DURING TENURE

14) POST-TENURE POSITION / JOB TITLE
Bioinformatics Supervisor

15) NAME AND ADDRESS OF POST-TENURE POSITION / ORGANIZATION
Walter Reed Army Institute of Research-Viral Diseases Branch. Robert Grant Ave. Silver Spring, MD

16) POST-TENURE POSITION STATUS / CATEGORY
Please indicate only one.

☐ Permanent position at the NRC host agency
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☐ Postdoctoral research
☐ Other (Please specify, possible) ______
☐ No information provided

17) APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM
On a scale of 1 – 10 (poor - excellent), please rate the following:

SHORT TERM VALUE
☐ 10 Development of knowledge, skills, and research productivity
Comments
AFRIMS, Bangkok provided many opportunities for learning about several infectious diseases important in SE Asia. I was able to observe and participate in methodologies associated with serology (viral culturing), molecular biology (sequencing and various PCR methodologies) and field collection of mosquitoes, training provided by the Entomology Dept. at AFRIMS. The organization encouraged collaboration and laboratory space was readily available my project along with any supplies I required.

LONG TERM VALUE
☐ 10 How the NRC Associateship award affected your career to date
Comments
Having come from an environmental microbiology background with minimal training in infectious disease, tenure with NRC has encouraged me to continue in infectious disease ecology and bioinformatic analysis. It has opened opportunities for me to continue in these fields and continue working and developing under my mentors.

**LAB SUPPORT**

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<td>10</td>
<td>Quality of support from the Laboratory--equipment, funding, orientation, safety and health guidelines, etc.</td>
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**Comments**

All protocols were clearly communicated and training was provided in English whenever required. Department personnel assisted me in locating and obtaining any supplies or equipment I needed for my work and clearing lab space.

**ADVISER/MENTOR SUPPORT**

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<td>Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)</td>
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**Comments**

Correspondence was always prompt and helpful.

**LPR SUPPORT**

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**Comments**

Correspondence was always prompt and helpful.

**NRC SUPPORT**

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<td>Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)</td>
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</tbody>
</table>

**Comments**

Omega travel was very helpful with all travel arrangements. See comments below about issues pertaining to being an internationally placed postdoc.

18) **PLEASE PROVIDE ANY SUGGESTIONS FOR PROGRAM IMPROVEMENT.**

There is not much support or guidance for internationally placed NRC postdocs except the coverage of my plane ticket. Aetna insurance also does not provide very much international support. Upon arriving to Bangkok there was no point of contact to assist in making our transition smooth. I was fortunate to have the support of the Virology secretary who served that role when I asked her to. There also was not much guidance for moving abroad specifically, I am now serving as a resource for some new NRC postdocs applying to the program as to how logistically to make moving to Bangkok Thailand feasible with respect to visas, housing, APO address, medical care, communications with NRC etc. If NRC is unable to provide this kind of abroad logistical support and the supporting agency where the fellow will work does not have this understanding to provide a point of contact for the fellow when they arrive, then the transition is challenging. Please feel free to post me as a resource to postdoctoral fellows with NRC that will be relocating abroad. I would be willing to give them information on the logistics of relocating to Thailand and advice on being an 'abroad' postdoc in general.

Please do **NOT** scan to PDF. Send the Final Report as MSWord document via e-mail to your NRC Program Coordinator

No handwritten signature required; but you may upload a scanned signature file below:

Asha Davis: adavis@nas.edu
Linda Sligh: lsligh@nas.edu
Jason Thornhill: jthornhill@nas.edu
Peggy Wilson: pwilson@nas.edu
Suzanne White: swhite@nas.edu

Id# Rev. July 2011 Proj/Act ID#
1) Associate Last or Family Name: Pichugin
First Name: Alexander
M.I.: V

3) Today's Date: July 6, 2012
Dates of Tenure: from January 12, 2009 to July 11, 2012

4) Host Agency: AMRMC (e.g., AFRL)
Laboratory or Center: WRAIR (e.g., Wright Patterson AFB)
Division / Directorate / Department: Malaria Vaccine Branch (e.g., High-Speed Propulsion)

5) Name of Laboratory NRC Adviser (and USMA Mentor, if applicable): Dr. Urszula Krzych

6) TITLE OF RESEARCH PROPOSAL

Prioritization and selection of pre-erythrocytic liver stage Plasmodia antigens as vaccine candidates.

7) SUMMARY OF RESEARCH DURING TENURE

1) 10 novel liver stage Pb antigens reduce LS and BS parasite burden in C57Bl/6 mice.
2) 3 novel liver stage Pb antigens sustain protection during 6 months after the last immunization.
3) 3 novel liver stage Pb antigens enhance protection induced by PbCSP.
4) Established caged MHC-tetramer technology to use for discovery of T cell epitopes in malaria antigens.
5) Identified 3 immunodominant CD8 T cell epitopes from Pb PEVA.

(USMA Davies Fellow: please add summary of teaching, including classes taught.)

8) RESEARCH IN PROGRESS

We showed that ten of 23 initial Pb DNA constructs induce the protection in C57Bl/6 mice after IM and Gene Gun delivery and subsequent challenge with P. berghei. Six orthologues of ten protective in P. berghei model antigens were also protective in P. yoelii-BALB/c model.

We have established novel state-of-art high throughput method of caged MHC-tetramers to evaluate frequency of peptide-specific CD8 T cells in mice protected by GAP and RAS. We have screened ~400 of Kb- and Db-restricted CD8 peptides from 28 initial Pb antigens and determined three immunodominant epitopes on two protective vaccine candidates.

9) PUBLICATIONS AND PAPERS RESULTING FROM NRC ASSOCIATESHIP RESEARCH

Provide complete citations: author(s), title, full name of journal, volume number, page number(s), and year of publication.

a) Publications in peer-reviewed journals

Anjali Yadava, Saule Nurmukhambetova, Alexander V Pichugin, Joanne M Lumsden
"Cross-Species Immunity Following Immunization with a Circumsporozoite Protein-Based Vaccine for Malaria" J Infect Dis. 2012, 205(9):1456-63.

b) Books, book chapters, other publications

c) Manuscripts in preparation, manuscripts submitted

"Early Liver Stage Transcriptome of Plasmodium falciparum Reveals Novel Malaria Vaccine Candidates", in preparation

10) PATENT OR COPYRIGHT APPLICATIONS RESULTING FROM NRC ASSOCIATESHIP RESEARCH

Provide titles, inventors, and dates of applications.
11) PRESENTATIONS AT SCIENTIFIC MEETINGS OR CONFERENCES

Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.

International

Alexander V. Pichugin, Lindsey Ehrler, Sharvan Sehrawat, Cate Speake, Patrick Duffy, Hidde Ploegh, Urszula Krzych
"Application of novel caged MHC-tetramer technology for the discovery of immunodominant CD8 T cell epitopes on Plasmodium liver stage antigens"

Alexander V. Pichugin, Cate Speake, Lindsey Ehrler, Lindsay Holladay, Valentino Garcia, Bob Morrison, Patricia DeLaVega, Isaac Chalom, D. Gray Heppner, Patrick Duffy and Urszula Krzych
"Novel Plasmodium berghei pre-erythrocytic liver stage antigens as potential vaccine candidates"

Domestic

Alexander V. Pichugin, Stasya zarling, Lindsey Ehrler, Sharvan Sehrawat, Cate Speake, Patrick Duffy, Hidde Ploegh, Urszula Krzych
"Identification of Plasmodiun berghei novel liver stage CD8 T cell epitopes by caged MHC class I tetramer technology"

12) SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES

Include dates, names and locations of seminars.

13) PROFESSIONAL AWARDS RECEIVED DURING TENURE

14) POST-TENURE POSITION / JOB TITLE

Senior Immunologist

15) NAME AND ADDRESS OF POST-TENURE POSITION / JOB ORGANIZATION

Clinical Research Management
411 Aviation Way Ste. 220
Frederick, MD 21701

16) POST-TENURE POSITION STATUS / CATEGORY

Please indicate only one.

☐ Permanent position at the NRC host agency
☒ Contract or temporary position at the NRC host Agency Abbreviate Host Laboratory/Center WRAIR
☐ Research/Administrative position with another U.S.-government agency
☐ Research/Administrative position with a foreign-government agency
☐ Research/teaching position at a U.S. college or university
☐ Research/teaching position at a foreign college or university
☐ Research/administration position in private industry in the U.S.
☐ Research/administration position in private industry outside of the U.S.
☐ Research/administration position with a non profit
☐ Self-employed/consulting
☐ Postdoctoral research
☐ Other (Please specify, possible) _____
☐ No information provided

17) APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM

On a scale of 1 – 10 (poor - excellent), please rate the following:

SHORT TERM VALUE

☐ Development of knowledge, skills, and research productivity
Comments

LONG TERM VALUE

☐ How the NRC Associateship award affected your career to date
Comments

LAB SUPPORT

☐ Quality of support from the Laboratory--equipment, funding, orientation, safety and health guidelines, etc.
Comments
ADVISER/MENTOR SUPPORT
10 Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)
Comments

LPR SUPPORT
10 Quality of administrative support from the Laboratory (e.g., NIST, NRL, IWR, FHWA) NRC Program Representative (LPR)
Comments

NRC SUPPORT
10 Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)
Comments

18) PLEASE PROVIDE ANY SUGGESTIONS FOR PROGRAM IMPROVEMENT.

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No handwritten signature required; but you may upload a scanned signature file below:

Alexander Pichugin

Id# Rev. July 2011 Proj/Act ID#

Asha Davis: adavis@nas.edu
Linda Sligh: lsligh@nas.edu
Jason Thornhill: jthornhill@nas.edu
Peggy Wilson: pwilson@nas.edu
Suzanne White: swwhite@nas.edu
**Final Report**

1) **Associate Last or Family Name**
   - Rajendran

2) **First Name**
   - Gnana

3) **M.I.**
   - R

4) **Today's Date**
   - September 4, 2012

5) **Dates of Tenure**
   - From September 7, 2010 to August 31, 2012

6) **Host Agency**
   - AMRMC (e.g., AFRL)

7) **Laboratory or Center**
   - WRAIR (e.g., Wright Patterson AFB)

8) **Division / Directorate / Department**
   - ET (e.g., High-Speed Propulsion)

9) **Name of Laboratory NRC Adviser (and USMA Mentor, if applicable)**
   - Dr. Michael Kozar

10) **Title of Research Proposal**
    - Design and synthesis of decaquinate derivatives toward new antimalarial agents

11) **Summary of Research During Tenure**
    - Itemize significant findings in concise form, utilizing key concepts/words.
    - 1) Several compounds were synthesized by the SAR of DQ and submitted for in vitro testing against blood stage malaria, specifically P. falciparum D6, W2, C235 and C2B strains, and assessed for metabolic stability in the mouse and human microsomes.
    - 2) In many cases the compounds solubility was improved but the compounds either lost potency against the C2B resistant strain of malaria or microsomal stability.
    - 3) A few interesting trends were discovered. An unprecedented ester replacement, to the ethyl or morpholine amide was discovered that maintained potency against D6, W2, C235 but unfortunately lost activity against the key C2B atovaquone resistant strain.
    - 4) Additionally, many compounds were synthesized that maintained in vitro potency with significantly lower clogP (main focus of the research effort).
    - 5) Efforts are currently toward acquiring a complete set of data to select profile compounds to be scaled up and tested in in vivo models.
    (USMA Davies Fellow: please add summary of teaching, including classes taught.)

12) **Research in Progress**
    - Describe in no more than 100 words.
    - Please see the attached document.

13) **Publications and Papers Resulting from NRC Associateship Research**
    - Provide complete citations: author(s), title, full name of journal, volume number, page number(s), and year of publication.
    - a) Publications in peer-reviewed journals
    - b) Books, book chapters, other publications
    - c) Manuscripts in preparation, manuscripts submitted

14) **Patent or Copyright Applications Resulting from NRC Associateship Research**
    - Provide titles, inventors, and dates of applications.

15) **Presentations at Scientific Meetings or Conferences**
    - Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.
International

Domestic

12) SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES  Include dates, names and locations of seminars.

13) PROFESSIONAL AWARDS RECEIVED DURING TENURE

14) POST-TENURE POSITION / JOB TITLE

Scientist

15) NAME AND ADDRESS OF POST-TENURE POSITION / JOB ORGANIZATION

WRAIR, 503 Robert Grant Ave, Silver Spring, MD

16) POST-TENURE POSITION STATUS / CATEGORY  Please indicate only one.

☐ Permanent position at the NRC host agency
☒ Contract or temporary position at the NRC host Agency
Abbreviate Host Laboratory/Center WRAIR
☐ Research/Administrative position with another U.S.-government agency
☐ Research/Administrative position with a foreign-government agency
☐ Research/teaching position at a U.S. college or university
☐ Research/teaching position at a foreign college or university
☐ Research/administration position in private industry in the U.S.
☐ Research/administration position in private industry outside of the U.S.
☐ Research/administration position with a non profit
☐ Self-employed/consulting
☐ Postdoctoral research
☐ Other (Please specify, possible) ______
☐ No information provided

17) APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM
On a scale of 1 – 10 (poor - excellent), please rate the following:

SHORT TERM VALUE

10  Development of knowledge, skills, and research productivity
Comments

LONG TERM VALUE

10  How the NRC Associateship award affected your career to date
Comments

LAB SUPPORT

10  Quality of support from the Laboratory—equipment, funding, orientation, safety and health guidelines, etc.
Comments

ADVISER/MENTOR SUPPORT

10  Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)
Comments

LPR SUPPORT

10  Quality of administrative support from the Laboratory (e.g., NIST, NRL, IWR, FHWA) NRC Program Representative (LPR)
Comments

NRC SUPPORT

10  Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)
Comments

18) PLEASE PROVIDE ANY SUGGESTIONS FOR PROGRAM IMPROVEMENT.
Please do NOT scan to PDF. Send the Final Report as MSWord document via e-mail to your NRC Program Coordinator

No handwritten signature required; but you may upload a scanned signature file below:

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<th>Id#</th>
<th>Rev. July 2011</th>
<th>Proj/Act ID#</th>
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</table>
7) SUMMARY OF RESEARCH DURING TENURE  Itemize significant findings in concise form, utilizing key concepts/words.

1) Intravital Microscopy was successfully employed for investigating EG shedding in hemorrhagic shock/resuscitation for the very first time;

2) Intravital microscopy integrated with systemic hemodynamics evaluations may be essential and more accurate tools to identify changes and study mechanisms of EG shedding and systemic responses to hemorrhage and resuscitation therapy;

3) Compared to baseline and to the sham group, there was a 50% reduction in endothelial glycocalyx (EG) thickness after hemorrhage and 60% increase in the levels of plasma Syndecan-1;

4) Although resuscitation with LR and Hextend could stabilize hypotensive rats hemodynamically, these fluids were unable to restore EG thickness or coagulopathy (weak clots and prolonged coagulation time);

5) Rats who received fresh frozen plasma (FFP) restored venular EG thickness to baseline level in addition to improve the systemic hemodynamics and coagulation response (restored homeostasis).

(USMA Davies Fellow: please add summary of teaching, including classes taught.)

8) RESEARCH IN PROGRESS  Describe in no more than 100 words.

We added a few more resuscitation groups (i.e. hypertonic saline, normal saline) to our protocol. So the next step is to test other resuscitation fluids and their effects on the endothelial glycocalyx (EG) degradation, coagulation and vascular permeability. Also we will continue to investigate the link between the EG and coagulopathy, in our rat hemorrhage model. Other measurements, such as vascular permeability (through fluorescence leakage) will also be started soon. A new Confocal microscope is being setup in our lab and we will be performing EG measurements of specific components (proteoglycans and glycosaminoglycans) in vivo and in situ in the cremaster muscle by next year.

9) PUBLICATIONS AND PAPERS RESULTING FROM NRC ASSOCIATESHIP RESEARCH

Provide complete citations: author(s), title, full name of journal, volume number, page number(s), and year of publication.

a) Publications in peer-reviewed journals

b) Books, book chapters, other publications

None

c) Manuscripts in preparation, manuscripts submitted


10) PATENT OR COPYRIGHT APPLICATIONS RESULTING FROM NRC ASSOCIATESHIP RESEARCH
Provide titles, inventors, and dates of applications.
None

11) PRESENTATIONS AT SCIENTIFIC MEETINGS OR CONFERENCES
Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.
International
None

Domestic

Torres, L.N., Sondeen, J.L., Ji, L., Dubick, M.A. and Torres Filho I.P. – In vivo comparison of resuscitation fluids on the preservation of the endothelial glycocalyx after hemorrhagic shock in rats. 2012 Military Health System Research Symposium (former ATACCC), Fort Lauderdale, Florida, 2012 (posters were not published in any journal supplement).

12) SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES Include dates, names and locations of seminars.
None

13) PROFESSIONAL AWARDS RECEIVED DURING TENURE
None

14) POST-TENURE POSITION / JOB TITLE
Research Investigator

15) NAME AND ADDRESS OF POST-TENURE POSITION / JOB ORGANIZATION
US Army Institute of Surgical Research (USAISR)

16) POST-TENURE POSITION STATUS / CATEGORY Please indicate only one.
☐ Permanent position at the NRC host agency
☒ Contract or temporary position at the NRC host Agency
Abbreviate Host Laboratory/Center USAISR
☐ Research/Administrative position with another U.S.- government agency
☐ Research/Administrative position with a foreign- government agency
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☐ Research/teaching position at a foreign college or university
☐ Research/administration position in private industry in the U.S.
☐ Research/administration position in private industry outside of the U.S.
☐ Research/administration position with a non profit
☐ Self-employed/consulting
☐ Postdoctoral research
☐ Other (Please specify, possible) No information provided

17) APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM
On a scale of 1 – 10 (poor - excellent), please rate the following:

SHORT TERM VALUE
10 Development of knowledge, skills, and research productivity
Comments

LONG TERM VALUE
10 How the NRC Associateship award affected your career to date
Comments

LAB SUPPORT
☐ Quality of support from the Laboratory--equipment, funding, orientation, safety and health guidelines, etc.
Comments
ADVISER/MENTOR SUPPORT
10 Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)

Comments

LPR SUPPORT
8 Quality of administrative support from the Laboratory (e.g., NIST, NRL, IWR, FHWA) NRC Program Representative (LPR)

Comments

NRC SUPPORT
7 Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)

Comments

I could be luckier to have Jason as my coordinator: he was absolutely great, helpful and very supportive. The moving company showed efficiency, accountability, and professionalism! I am 100% satisfied. The driver was very experienced and made my move as stress-free as possible. The only complain I have is regarding the travel reimbursements: it took a long time (more than a month or 2) to receive my reimbursements, although the staff handling the travel were always very nice and knowledgeable.

18) PLEASE PROVIDE ANY SUGGESTIONS FOR PROGRAM IMPROVEMENT.

Please do NOT scan to PDF. Send the Final Report as MSWord document via e-mail to your NRC Program Coordinator.

No handwritten signature required; but you may upload a scanned signature file below:

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Id#                      Rev. July 2011                      Proj/Act ID#
1) Associate Last or Family Name | First Name | M.I.
Vecchi | Vittoria |

3) Today's Date | Dates of Tenure
N/A | from November 14, 2012 to October 22, 2012 |

4) Host Agency | Laboratory or Center | Division / Directorate / Department
AMRMC (e.g., AFRL) | USAISR (e.g., Wright Patterson AFB) | (e.g., High-Speed Propulsion) |

5) Name of Laboratory NRC Adviser (and USMA Mentor, if applicable)
Dr. Andriy Batchinsky |

6) TITLE OF RESEARCH PROPOSAL
Quantitative lung CT-scan as monitoring tool for the Acute Respiratory Distress Syndrome |

7) SUMMARY OF RESEARCH DURING TENURE
Itemize significant findings in concise form, utilizing key concepts/words.

1) Effects of radiation dose reduction on lung quantitative CT scan results in healthy in the Acute Respiratory Distress Syndrome: low-dose chest CT as a valuable tool for quantification and monitoring of pulmonary disease reducing patient exposure
2) Use of quantitative CT for in vivo lung weight measurement: evaluate and monitor the time course of lung edema in ARDS measuring lung weight by qCT
3) Pressure-guided positioning of bicaval dual-lumen catheters for veno-venous extracorporeal gas exchange
4) Low-flow extracorporeal gas exchange for the treatment of ARDS caused by smoke inhalation and cutaneous burn in pigs
5) Extracorporeal Gas Exchange in awake spontaneously breathing sheep before and after the induction of ARDS

(USMA Davies Fellow: please add summary of teaching, including classes taught.)
N/A |

8) RESEARCH IN PROGRESS
Describe in no more than 100 words.

The use of low-dose chest CT, if producing satisfactory image quality for the purpose of the quantitative analysis, could favor the use of qCT as a monitoring tool of severity and time-course of pulmonary disease reducing patient radiation exposure. The aim of the this study is therefore to investigate if and how a reduction in tube current-time product (mAs) during CT image acquisition affects quantitative results in healthy lungs and ARDS. |

9) PUBLICATIONS AND PAPERS RESULTING FROM NRC ASSOCIATESHIP RESEARCH
Provide complete citations: author(s), title, full name of journal, volume number, page number(s), and year of publication.

a) Publications in peer-reviewed journals

b) Books, book chapters, other publications

c) Manuscripts in preparation, manuscripts submitted
- Extracorporeal Gas Exchange in awake spontaneously breathing sheep before and after the induction of ARDS - manuscript in preparation
10) **PATENT OR COPYRIGHT APPLICATIONS RESULTING FROM NRC ASSOCIATESHIP RESEARCH**

Provide titles, inventors, and dates of applications.

N/A

11) **PRESENTATIONS AT SCIENTIFIC MEETINGS OR CONFERENCES**

Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.

**International**


**Domestic**


12) **SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES**

Include dates, names and locations of seminars.

N/A

13) **PROFESSIONAL AWARDS RECEIVED DURING TENURE**

N/A

14) **POST-TENURE POSITION / JOB TITLE**

Radiology Resident

15) **NAME AND ADDRESS OF POST-TENURE POSITION / JOB ORGANIZATION**

Università degli Studi di Milano, Milan, Italy

16) **POST-TENURE POSITION / STATUS / CATEGORY**

- Permanent position at the NRC host agency
- Contract or temporary position at the NRC host Agency
- Research/Administrative position with another U.S.-government agency
- Research/Administrative position with a foreign-government agency
- Research/teaching position at a U.S. college or university
- Research/teaching position at a foreign college or university

Please indicate only one.

17) **APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM**

On a scale of 1 – 10 (poor - excellent), please rate the following:

**SHORT TERM VALUE**

- Development of knowledge, skills, and research productivity
  Comments

**LONG TERM VALUE**

- How the NRC Associateship award affected your career to date
  Comments

**LAB SUPPORT**

- Quality of support from the Laboratory--equipment, funding, orientation, safety and health guidelines, etc.
  Comments

**ADVISER/MENTOR SUPPORT**

- Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)
  Comments
LPR SUPPORT
10 Quality of administrative support from the Laboratory (e.g., NIST, NRL, IWR, FHWA) NRC Program Representative (LPR)
Comments

NRC SUPPORT
10 Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)
Comments

18) PLEASE PROVIDE ANY SUGGESTIONS FOR PROGRAM IMPROVEMENT.

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</table>

Id# Rev. July 2011 Proj/Act ID#
Treating Chemical Agent Exposure Through Multiple Drug Delivery Routes

1) Developed and characterized novel system for inhalational exposure of conscious animals to chemical agents
2) Utilized above system to examine the toxicokinetics (TK) of nerve agents and their analogs in rats
3) Investigated utility of bronchodilators in a treatment regimen for inhalational chemical agent exposure
4) 
5) 

Current research focuses on the inhalation of various chemical agents and the associated effects on different organ systems in order to characterize the role of exposure route on progression and presence of symptoms. The body of literature that describes the inhalation of chemical agents is incomplete, despite being the most likely route by which agent will enter the body. The route of exposure is central to the progression and presence of symptoms, which in turn form the basis for the development of treatments and treatment regimens. Recent development of a novel inhalation model has progressed research toward development and characterization of improved methods of drug delivery in chemical agent exposure scenarios.

1. Development of Model for Nerve Agent Inhalation in Conscious Rats
2. Respiration Toxicity in Non-Anesthetized Rats Following Inhalation Exposure to Soman Vapor
2 others, as yet untitled, in preparation

Provide complete citations: author(s), title, full name of journal, volume number, page number(s), and year of publication.

a) Publications in peer-reviewed journals
N/A

b) Books, book chapters, other publications
N/A

c) Manuscripts in preparation, manuscripts submitted

1. Development of Model for Nerve Agent Inhalation in Conscious Rats
2. Respiration Toxicity in Non-Anesthetized Rats Following Inhalation Exposure to Soman Vapor
2 others, as yet untitled, in preparation

Provide titles, inventors, and dates of applications.
N/A
11) PRESENTATIONS AT SCIENTIFIC MEETINGS OR CONFERENCES
Provide complete references: author(s), title, abstract/proceeding citation, meeting name and location.

International
N/A

Domestic
1. B. Wong, A. Sciuto, G. Murphy.
Development of a Head-Out Vapor Inhalation Model for the Evaluation of Toxicity Following Exposure to Chemical Agents in Non-Anesthetized Rats.
US Army Medical Defense Bioscience Review, Hunt Valley, MD.

2. B. Wong, A. Sciuto, G. Murphy
A Head-Out Vapor Inhalation Model for the Evaluation of Soman Toxicity in Non-Anesthetized Rats
Shoresh Chemical and Biological Defense Conference, Fort Detrick, MD.

12) SEMINARS OR LECTURES DELIVERED AT UNIVERSITIES AND/OR INSTITUTES
Include dates, names and locations of seminars.
N/A

13) PROFESSIONAL AWARDS RECEIVED DURING TENURE
N/A

14) POST-TENURE POSITION / JOB TITLE
Post-Doctoral Research Associate at MRICD

15) NAME AND ADDRESS OF POST-TENURE POSITION / JOB ORGANIZATION
USAMRICD
3100 Rickett's Point Road
APG, MD 21010

16) POST-TENURE POSITION STATUS / CATEGORY
Please indicate only one.
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Abbreviate Host Laboratory/Center APG-MRICD
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☐ Research/teaching position at a foreign college or university

17) APPRAISAL OF RESEARCH ASSOCIATESHIP PROGRAM
On a scale of 1 – 10 (poor - excellent), please rate the following:

SHORT TERM VALUE
☐ Development of knowledge, skills, and research productivity
Comments
Work at MRICD was intended to continue the research from my graduate work; however, the research took on its own directions, resulting in development of knowledge and skills in new areas which complemented my existing knowledge base well. Despite many technical hurdles, challenges proved to help maximize productivity. Overall, work here has provided a different set of challenges, which have resulted in net positive gains in all categories.

LONG TERM VALUE
☐ How the NRC Associateship award affected your career to date
Comments
Cannot currently ascertain effects, as a permanent position at my institute was not available, rendering it difficult to assess the competitive benefits of an NRC fellowship. However, the NRC program has helped me make connections which are likely to have a beneficial effect on my career in the future.

LAB SUPPORT
☐ Quality of support from the Laboratory—equipment, funding, orientation, safety and health guidelines, etc.
Comments
Funding has struggled for the institute recently, and orientation was somewhat confusing. Guidelines for safety and health are excellently implemented in the laboratory setting but less observed in the office setting. Host laboratory/advisor group has done the best possible given the overall situation.

ADVISER/MENTOR SUPPORT

10 Quality of mentoring from the Laboratory NRC Adviser (USMA Mentor, if applicable)

Comments
Dr Sciuto has provided me with excellent support in all aspects of laboratory work and research and career opportunities. I have been able to integrate my skill and mindset into an adaptive laboratory group and have been encouraged to contribute in any possible. He is the type of mentor that is able to help all personality types excel, and I have noted that in both myself and the people I work with in his group.

LPR SUPPORT

8 Quality of administrative support from the Laboratory (e.g., NIST, NRL, IWR, FHWA) NRC Program Representative (LPR)

Comments
Dr Kan has been available for all the necessary assistance I have needed in terms of communication between the institute and NRC.

NRC SUPPORT

9 Quality of administrative support. Please assess respective NRC aspects (e.g., moving company, insurance, Omega, payroll, coordinator, travel, etc.)

Comments
I was on a very tight schedule to move out when I received my appointment, and within 10 hours of having been confirmed, everything concerning my move was sorted out. The moving company was excellent and reliable and the billing was easily sorted out with the NRC. The personnel in charge of communicating with me for health insurance were knowledgeable and of great assistance. Travel and payroll were also excellent in their communication and ability to rapidly process requests.

18) PLEASE PROVIDE ANY SUGGESTIONS FOR PROGRAM IMPROVEMENT.

Communication on whether it was possible to obtain a security clearance through NRC was somewhat lacking, or uninforming. Clear delineations as to the policy would be helpful, especially to the institute.

Please do NOT scan to PDF. Send the Final Report as MSWord document via e-mail to your NRC Program Coordinator

No handwritten signature required; but you may upload a scanned signature file below:

Asha Davis: adavis@nas.edu
Linda Sligh: lsligh@nas.edu
Jason Thornhill: jthornhill@nas.edu
Peggy Wilson: pwilson@nas.edu
Suzanne White: swhite@nas.edu

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