

## DEPARTMENT OF THE AIR FORCE 59TH MEDICAL WING (AETC) JOINT BASE SAN ANTONIO - LACKLAND TEXAS



27 APR 2017

MEMORANDUM FOR SGOIC

ATTN: LT COL STEVEN J. ACEVEDO

FROM: 59 MDW/SGVU

SUBJECT: Professional Presentation Approval

- 1. Your paper, entitled Regulation of EGF and Prostaglandin Expression During Neonatal Gastrointestinal Injury in a Non-Human Primate Explant Model presented at/published to Pediatric Academic Societies Meeting, San Francisco CA, 5-9 May 2017 in accordance with MDWI 41-108, has been approved and assigned local file #17199.
- 2. Pertinent biographic information (name of author(s) title, etc.) has been entered into our computer file. Please advise us (by phone or mail) that your presentation was given. At that time, we will need the date (month, day and year) along with the location of your presentation. It is important to update this information so that we can provide quality support for you, your department, and the Medical Center commander. This information is used to document the scholarly activities of our professional staff and students, which is an essential component of Wilford Hall Ambulatory Surgical Center (WHASC) internship and residency programs.
- 3. Please know that if you are a Graduate Health Sciences Education student and your department has told you they cannot fund your publication, the 59th Clinical Research Division may pay for your basic journal publishing charges (to include costs for tables and black and white photos). We cannot pay for reprints. If you are a 59 MDW staff member, we can forward your request for funds to the designated Wing POC at the Chief Scientist's Office, Ms. Alice Houy, office phone: 210-292-8029; email address: alice.houy.civ@mail.mil.
- 4. Congratulations, and thank you for your efforts and time. Your contributions are vital to the medical mission. We look forward to assisting you in your future publication/presentation efforts.

LINDA STEEL-GOODWIN, Col, USAF, BSC Director, Clinical Investigations & Research Support

Linda Steel-Goodwin

## PROCESSING OF PROFESSIONAL MEDICAL RESEARCH/TECHNICAL PUBLICATIONS/PRESENTATIONS

## **INSTRUCTIONS**

## USE ONLY THE MOST CURRENT 69 MDW FORM 3039 LOCATED ON AF E-PUBLISHING

- 1. The author must complete page two of this form:
  - a. In Section 2, add the funding source for your study [ e.g., 59 MDW CRD Graduate Health Sciences Education (GH8E) (8G5 O&M); 8G5 R&D;
    Trt-Service Nursing Research Program (TSNRP); Defense Medical Research & Development Program (DMRDP); NIH; Congressionally Directed
    Medical Research Program (CDMRP); Grants; etc.)
  - in Section 2, there may be funding available for journal costs, if your department is not paying for figures, tables or photographs for your publication.
     Please state "YES" or "NO" in Section 2 of the form, if you need publication funding support.
- 2. Print your name, rank/grade, sign and date the form in the author's signature block or use an electronic signature.
- Attach a copy of the 59 MDW IRB or IACUC approval letter for the research related study. If this is a technical publication/presentation, state the type (e.g. case report, QA/QI study, program evaluation study, informational report/briefing, etc.) in the "Protocol Title" box.
- Attach a copy of your abstract, paper, poster and other supporting documentation.
- Save and forward, via email, the processing form and all supporting documentation to your unit commander, program director or immediate supervisor for review/approval.
- 6. On page 2, have either your unit commander, program director or immediate supervisor:
  - a. Print their name, rank/grade, title; sign and date the form in the approving authority's signature block or use an electronic signature.
- 7. Submit your completed form and all supporting documentation to the CRD for processing (59crdpubspres@us.af.mil). This should be accomplished no later than 30 days before final clearance is required to publish/present your materials. If you have any questions or concerns, please contact the 59 CRD/Publications and Presentations Section at 292-7141 for assistance.
- The 59 CRD/Publications and Presentations Section will route the request form to clinical investigations, S02 ISG/JAC (Ethics Review) and Public Affairs
  (59 MDW/PA) for review and then forward you a final letter of approval or disapproval.
- Once your manuscript, poster or presentation has been approved for a one-time public release, you may proceed with your publication or presentation submission activities, as stated on this form. Note: For each new release of medical research or technical information as a publication/presentation, a new 59 MDW Form 3039 must be submitted for review and approval.
- If your manuscript is accepted for scientific publication, please contact the 59 CRD/Publications and Presentations Section at 292-7141. This information is reported to the 59 MDW/CC. All medical research or technical information publications/presentations must be reported to the Defense Technical information Center (DITC). See 59 MDWI 41-108, Presentation and Publication of Medical and Technical Papers, for additional information.
- 11. The Joint Ethics Regulation (JER) DoD 5500.07-R, Standards of Conduct, provides standards of ethical conduct for all DoD personnel and their interactions with other non-DoD entities, organizations, societies, conferences, etc. Part of the Form 3039 review and approval process includes a legal ethics review to address any potential conflicts related to DoD personnel participating in non-DoD sponsored conferences, professional meetings, publication/presentation disclosures to domestic and foreign audiences, DoD personnel accepting non-DoD contributions, awards, honoraria, gifts, etc. The specific circumstances for your presentation will determine whether a legal review is necessary. If you (as the author) or your supervisor oheok "NO" in blook 17 of the Form 3039, your research or feohnloal documents will not be forwarded to the 602 ISG/JAC legal office for an ethics review. To assist you in making this decision about whether to request a legal review, the following examples are provided as a guideline:

For presentations before professional societies and like organizations, the 59 MDW Public Affairs Office (PAO) will provide the needed review to ensure proper discipliners are included and the subject matter of the presentation does not create any cause for DoD concern.

if the sponsor of a conference or meeting is a DoD entity, an ethics review of your presentation is not required, since the DoD entity is responsible to obtain all approvals for the event.

If the sponsor of a conference or meeting is a non-DoD commercial entity or an entity seeking to do business with the government, then your presentation should have an ethics review.

If your travel is being paid for (in whole or in part) by a non-Federal entity (someone other than the government), a legal ethics review is needed. These requests for legal review should come through the 59 MDW Gifts and Grants Office to 502 ISG/JAC.

If you are receiving an honorarium or payment for speaking, a legal ethics review is required.

If you (as the author) or your supervisor check "YES" in block 17 of the Form 3039, your research or technical documents will be forwarded simultaneously to the 502 ISG/JAC legal office and PAO for review to help reduce turn-around time. If you have any questions regarding legal reviews, please contact the legal office at (210) 671-5795/3365, DSN 473.

- NOTE: All abstracts, papers, posters, etc., should contain the following disclaimer statement:
  - "The views expressed are those of the [author(s)] [presenter(s)] and do not reflect the official views or policy of the Department of Defense or its Components"
- NOTE: All abstracts, papers, posters, etc., should contain the following disclaimer statement for research involving humans:
  - "The voluntary, fully informed consent of the subjects used in this research was obtained as required by 32 CFR 219 and DODI 3216.02\_AFI
- NOTE: All abstracts, papers, posters, etc., should contain the following disclaimer statement for research involving animals, as required by AFMAN 40-401\_IP:
  - "The experiments reported herein were conducted according to the principles set forth in the National Institute of Health Publication No. 80-23, Guide for the Care and Use of Laboratory Animals and the Animal Welfare Act of 1966, as amended."

PROCESSING OF PROFESS	SIONAL MEDICAL	RESEARCH/TECHNICAL	PUBLICATIONS	/PRES	ENTATIONS
To: CLINICAL RESEARCH     Acevedo, Ste	othor's Name, Rank, Gradeven, J., Lt Col., O5	de, Office Symbol)	3. GME/GHSE STUD	100000	PROTOCOL NUMBER:
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6. TITLE OF MATERIAL TO BE PUBLISHED ( Regulation of EGF and Prostaglandin Exp		atal Gastrointestinal Injury i	n a Non-Human Pri	mate Er	splant Model
7. FUNDING RECEIVED FOR THIS STUDY?	YES NO FUN	NOING SOURCE: Air Force Re	search Grant		
8. DO YOU NEED FUNDING SUPPORT FOR	PUBLICATION PURPO	SES: YES NO			
9. IS THIS MATERIAL CLASSIFIED? YE	S ⊠ NO				
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11c. POSTER (To be demonstrated at r Pediatric Academic Societies Meetin 11d. PLATFORM PRESENTATION (At	ng, San Francisco, CA	, May 5-9			
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11e. OTHER (Describe: name of meeting	g, city, state, and date of	meeting.)			
12. HAVE YOUR ATTACHED RESEARCH/TE	CHNICAL MATERIALS	BEEN PREVIOUSLY APPROVE	D TO BE PUBLISHED	WPRE88	ENTED?
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April 17, 2017					
14. 59 MDW PRIMARY POINT OF CONTACT (Last Name, First Name, M.I., email) Acevedo, Steven J				15. DUTY PHONE/PAGER NUMBER 210-916-7078	
16. AUTHORSHIP AND CO-AUTHOR(8) List					
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e. Mustafa, Shamim	Civ			UT Health San Antonio	
17. IS A 502 IBG/JAC ETHICS REVIEW REQ		07-R)? ☐ YES ☒ NO			
I CERTIFY ANY HUMAN OR ANIMAL RESEA 219, AFMAN 40-401_IP, AND 59 MDWI 41-10 ACCURATE MANUSCRIPT FOR PUBLICATION	RCH RELATED STUDIES	S WERE APPROVED AND PER INAL VERSION OF THE ATTAC			
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TO: Clinical Research Division 59 MDW/CRD Contact 292-7141 for email Instructions.	24. DATE RECEIVED 4/12/2017	25. ASSIGNED PROCESSING REQUEST FILE 17199	NUMBER			
26. DATE REVIEWED		27. DATE FORWARDED TO 502 ISGUAC				
April 25, 2017						
28. AUTHOR CONTACTED FOR RECOMM	ENDED OR NECESSARY	CHANGES: NO TYES If yes, give date.	□ N/A			
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**TITLE:** Regulation of EGF and Prostaglandin Expression During Neonatal Gastrointestinal Injury in a Non-Human Primate Explant Model

AUTHORS: Steven J. Acevedo, MD<sup>1</sup>, Nicholas B. Alana<sup>2</sup>, Nicholas R. Carr, DO<sup>1</sup>, Cynthia L. Blanco MD<sup>3</sup>, Shamim B. Mustafa PhD<sup>3</sup>, Jonathan M. King PhD<sup>2</sup>

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BACKGROUND: Alterations in the Epidermal Growth Factor (EGF) and/or function of the EGF receptor (EGFR), have been related to GI injury. Bioactive compounds like Prostaglandin E2 (PGE2) also appear to have a role in barrier maintenance. A model evaluating the interaction of such compounds in vivo does not exist.

OBJECTIVE: Develop an ex vivo model to evaluate alterations/interactions of several bioactive compounds within the same tissue exposed to anti-inflammatory (indomethacin) and inflammatory (TNF) modulators.

DESIGN/METHODS: Preterm baboons delivered at 125-day gestation (term=184 days) via C-section were sacrificed at birth (D0=125 day, n=5) or chronically ventilated for 14 days (D14=125day + 14days, n=5). Explants were harvested from proximal ileum, cecum, and colon. Tissue was cultured for 24 hours while exposed to an inflammatory agent (TNF 10 ng/ml), or anti-inflammatory agent (indomethacin 50 mM). Secreted PGE2 in culture media was quantified by ELISA. mRNA expression of prostanoid (EP) receptor subtypes, EGFR, AKT, and inducible nitric oxide synthase (iNOS) were evaluated by qRT-PCR. Protein isolates were measured by ELISA for EGFR, p-EGFR, and AKT.

RESULTS: PGE2 secretion was similar between D0 and D14 controls. PGE2 secretion varied across intestinal regions significantly in control groups (p<0.05), pronounced in the terminal ileum. A reduction in PGE2 secretion was noted in D0 and D14 tissues exposed to indomethacin alone or with TNF in all three intestinal regions (p<0.05). There was a 2 to 4-fold downregulation in mRNA expression of EP2/EP4/EGFR/AKT and inducible NO in D0 tissue, exposed to either TNF or indomethacin in the ileum (p<0.05). These changes were not significant in D14 ileum with exception of iNOS elevation in indomethacin exposed tissues (p<0.05). Protein expression was modestly affected by age of tissue or changes in gene expression. TNF did not change PGE2 secretion, but downregulated gene expression and protein content in the EGFR pathway.

CONCLUSIONS: An explant model efficiently replicates in vivo tissue behavior allowing for testing of confounding variables within the same tissue and translation to an in vivo model. Indomethacin suppressed PGE2 secretion in preterm tissue and reduced gene expression in the EGFR pathway as a potential pathway for irreparable GI injury.

The views expressed are those of the author(s)/presenter(s) and do not reflect the official views or policy of the Department of Defense or its Components.

The experiments reported herein were conducted according to the principles set forth in the National Institute of Health Publication No. 80-23, Guide for the Care and Use of Laboratory Animals and the Animal Welfare Act of 1966, as amended.