

Southern California ASM 2022

RESEARCH POSTER SESSION

Date: October 15, 2022

Location: Online (Zoom)

Deadline for Abstract Submission:

5:00 PM (PST) on September 14, 2022

WHO SHOULD SUBMIT AN ABSTRACT?

The discipline of microbiology is highly diverse, and encompasses many specialized areas, which include but are not limited to, **clinical, ecological, environmental, industrial infectious disease immunology, marine, pharmaceutical, and public health microbiology.**

Undergraduate, graduate or post-baccalaureate students, and trainees who are participating in research projects pertaining to any of these specialized areas of microbiology are encouraged to submit an abstract.

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Student Participation Benefits

- Attend thought-provoking talks pertaining to microbiology, career development and networking with other microbiologists at the virtual synchronous meeting on October 15, 2022.
- All ePoster presenters will receive an official certificate of participation.
- All ePoster presenters have a chance to be selected for synchronous virtual oral presentation and win one of three awards on October 15, 2022.
- All ePoster presenters have a chance to be selected for a live oral 1-minute flash talk at the SCASM Annual Meeting in La Jolla, CA, October 22, 2022.

Student Awards

- Total of **3** awards, 1 travel grant (1st prize) and 2 free ASM student memberships (2nd and 3rd prize). The winner of the 1st prize will have the option to give a brief testimony in person or pre-record a video for streaming at the Annual SCASM Fall Meeting on October 22, 2022.
- The travel grant will be awarded to attend and present the poster, if accepted, at the national Annual General Meeting of the American Society for Microbiology (ASM) Microbe conference to be held in Houston, Texas, June 15 -19, 2023. The total value of the award is up to **\$1500**, toward conference registration and expenses accrued for travel, lodging, and food. In the case that the meeting will be virtual, the travel grant will cover the registration fees only. The trip will be **reimbursed** after the travel or virtual meeting has occurred with the appropriate itemized receipts provided to the SCASM Treasurer.

Judging Criteria

- Criteria for poster judging are listed on the poster evaluation form on page **(7)**.
- The top 3 posters will be selected for a live presentation at the SCASM Virtual Student Meeting. Based on the scores obtained from the live presentation the best presenter will win the travel grant award. The presenters with the 2nd and 3rd highest cumulative scores will win the ASM student memberships.
- There will be judges representing various areas of microbiology. Judges will include microbiology educators/researchers and post-doctoral fellows from academic institutions in Southern California.
- In the event of a tie, the winner(s) will be determined by the votes of 3 alternate judges.

Abstract Guidelines

- Abstracts must be typed in a 12-point font size, with a word limit of 350.
- Abstract must be submitted using the abstraction submission form on page **(6)**.
- Abstract must include the following: title, author including the principal investigator(s), affiliated institution(s), introduction, objectives, methods, materials, results and conclusion(s).
 - The reason for the study or how the study came about (e.g., hypothesis, discovery or central question) should be clearly stated.
 - Data must be included and must support the stated conclusion(s).

EXAMPLES OF ABSTRACTS THAT MAY BE REJECTED

- Abstracts that are general descriptions of a new product.
- Abstracts that read like advertisements.
- Abstracts that describe future studies.
- Abstracts that do not include data.

Please see page (8) for an example of an acceptable abstract

ePoster Guidelines

- ePoster submission includes a PDF file of the poster and a link to a 1-minute recorded poster presentation.
- Students will receive notification of abstract deposition and instructions on how to prepare and submit the poster and 1-minute flash talk on Wednesday, September 21, 2022.
- Poster must include the following sections: title, author including the principal investigator(s), affiliated institution(s), introduction, research question/hypothesis, objectives, methods, materials, results, conclusion(s), acknowledgment(s). Reference(s) section is optional.

Please see page (7) for the poster judging rubric and page (9) for an example of an acceptable poster

Presentation Guidelines for the 1-minute ePoster Presentation

- The “1-minute flash talk” is an opportunity to quickly and compellingly share the highlights of your poster. This presentation is a scorable item and the selection of the top 3 posters for the virtual live presentations will be influenced by your 1-minute flash talk.

Use the following links below for tips on preparing a quick elevator speech.

<https://www.software.ac.uk/home/cw11/giving-good-lightning-talk>

Abstract Submission Form

- **Deadline** for abstract submission is at **5:00PM (PST)** on **Wednesday, September 14, 2022**.
- **No more than 3 abstracts per PI/lab can be submitted.**
- **No co-presenters are allowed.**
- Students who submit an abstract must be a student member of SCASM (\$15/ one year; \$25/ two years; <https://scasm.org/membership-plans>).
- Abstract submission forms received after the deadline will be automatically rejected.
- **Notification of abstract acceptance** and invitation to submit an eposter and recorded 1-minute poster presentation video will be sent by email on **Wednesday, September 21, 2022**.
- **ePoster and 1-minute poster presentation video** must be uploaded by **5PM (PST)** on **Monday, October 3, 2022**.
- Selection of the top 10 in person presenters that will be invited to present their posters/1-minute flash talk at the SCASM Annual Meeting in La Jolla, CA, October 22,2022 and the top 3 virtual meeting presenters that will be announced by email by **Monday, October 10, 2022**. Presentation instructions and the evaluation rubric will be sent at that time.

Please use the following link to submit an abstract.

➤ <https://forms.gle/YzmDKRjZQUy6pb2K7>

For questions regarding abstract submissions please email:

Edith Porter, MD

Associate Chair and Professor of Microbiology & Immunology

Department of Biological Sciences

California State University, Los Angeles

Email: eporter@calstatela.edu

ePoster and Poster Presentation Evaluation Form

Judge's Name: _____

Poster Category (check box): Undergraduate Graduate/Post-baccalaureate

Poster First Author: _____

Poster Title: _____

1= Poor, 2= Fair, 3= Good, 4= Excellent, 5= Outstanding

REVIEW CRITERIA	Please circle the score for each category				
Abstract – includes summary of pertinent details according to poster abstract guidelines.	1	2	3	4	5
Introduction and Statement of the Research Question/Hypothesis – clearly states the research question/hypothesis with appropriate background to the bigger picture.	1	2	3	4	5
Objectives – are clearly stated and addressed by the experimentation.	1	2	3	4	5
Methodology – methods/techniques are appropriate and properly applied.	1	2	3	4	5
Results – logical, clearly presented, and appropriately summarized.	1	2	3	4	5
Conclusions, Future Research – based on given results, emphasizes significance and possible implications of study.	1	2	3	4	5
Overall Organization of Poster – graphics, photographs, other visual aids, and text are well prepared, clean, free of errors and appropriate for the presentation, acknowledgements included.	1	2	3	4	5
1-minute flash talk – engaged presentation that reflects the poster and is free of jargon.	1	2	3	4	5
Total Score					

Example of an Acceptable Abstract

Effect of the antimicrobial peptide hBD-2 on flagellin gene expression in *Pseudomonas aeruginosa*

Brent Beadell¹, Kevin Parducho¹, Mabel Bush¹, and Edith Porter¹

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Introduction: The ubiquitous Gram-negative rod-shaped motile bacterium *Pseudomonas aeruginosa* (PA) is an opportunistic pathogen that causes infection in the airways primarily in immunocompromised individuals. PA is known for its multitude of resistance mechanisms against antibiotics. One of the key mechanisms of resistance stems from its ability to form biofilms on host epithelial surfaces limiting the diffusion of antibiotics. Biofilm production is regulated by quorum sensing and involves early cessation of flagella expression followed by an upregulation of exopolysaccharide production. In immunocompetent individuals, likely due to their epithelial cells mounting effective immune responses, PA typically does not form biofilms. Preliminary data from our laboratory indicate that the epithelial antimicrobial peptide, human β -Defensin-2 (hBD-2), reduces biofilm production possibly through quorum sensing interference. We hypothesize that if PA biofilm inhibition occurs in the presence of hBD-2 via quorum sensing, then flagellin gene expression should be upregulated paralleled with a down regulation genes involved in biofilm production. **Objective:** This study aimed to quantify the relative expression of *pslA*, a gene involved in exopolysaccharide synthesis, and *flgF*, a gene coding for flagellin, in PA after exposure to varying concentrations of hBD-2. **Methods:** Mid-logarithmic growth phase PA was incubated at $\sim 2 \times 10^8$ CFU/mL in 10% MH / 140 mM NaCl in the presence and absence of 0.25 and 0.5 microM hBD-2 for 2 h. Thereafter, bacteria were dislodged by addition of 1 mm glass beads and 10 min vortexing. This was followed by RNA extraction, cDNA synthesis, and real time PCR with SYBR Green technology probing for the target genes *pslA*, and *flgF* and the housekeeping gene *gapA*. **Results:** *flgF* gene expression was over 50-fold times increased by both, 0.25 μ M and 0.50 μ M hBD-2 treatments and *pslA* gene expression was increased 2 -3 times compared to the solvent control. **Conclusion:** If confirmed, the substantial increase in flagellin gene expression in the presence of hBD-2 would be consistent with biofilm inhibition of PA through a quorum sensing pathway. Understanding the antimicrobial peptide mediated interference with biofilm could lead to novel clinical treatments against biofilm forming microbes, in particular PA.

Example of an Acceptable Poster

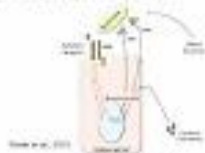


Effect of the Antimicrobial Peptide hBD-2 on Flagellin Gene Expression in *Pseudomonas aeruginosa*

Brent Beadell, Kevin R. Paruchko, Mabel Bush, and Edith Portes
Department of Biological Sciences, California State University Los Angeles, Los Angeles, CA 90032

INTRODUCTION

- The ubiquitous, Gram-negative, rod-shaped model bacterium *Pseudomonas aeruginosa* (PA) is an opportunistic pathogen that causes infection in the airways primarily in immunocompromised individuals (CDC, 2018).
- PA is known for its multitude of resistance mechanisms against antibiotics. One of the key mechanisms of resistance stems from its ability to form biofilms on host epithelial surfaces limiting the diffusion of antibiotics.
- Biofilm production is regulated by quorum sensing and involves early secretion of flagellar components followed by an upregulation of extracellular matrix production. In immunocompetent individuals, study due to their epithelial cells mounting effective immune responses, PA typically does not form biofilms (Dworkin et al., 2015).
- One of the first lines of defense against PA is found in the innate immune response involving the release of antimicrobial peptides (AMPs) by epithelial cells and phagocytes.
- Preliminary data from our laboratory indicate that the cathelicidin antimicrobial peptide, human δ -Cathelicidin (hBD-2), reduces biofilm production possible through quorum sensing interference.
- Our laboratory is interested in exploring three AMPs and hBD-2 for use in novel vaccines and drug creation against PA.



HYPOTHESIS

- If PA biofilm inhibition occurs in the presence of hBD-2 via quorum sensing then flagellar gene expression should be upregulated paralleled with a decrease in biofilm production.



OBJECTIVE

- This study aimed to quantify the relative expression of *polA*, a gene involved in exopolysaccharide synthesis, and *flgC*, a gene coding for flagellin in PA after exposure to varying concentrations of hBD-2.

EXPERIMENTAL APPROACH



Figure 1. Representative way in which PA from PA culture was brought to testing phase. Isolated by subculture isolation with varying concentrations of hBD-2 in 10% dimethyl sulfoxide (DMSO) for 24 hours. PAH extract was performed to evaluate gene expression analysis.



Figure 2. Representative hBD-2 response concentration and purity after purification. After purifying sample using HPLC, purity levels in storage buffer were verified using HPLC. Purity levels were confirmed using HPLC. A representative graph of hBD-2 concentration was generated using HPLC. (A) shows representative concentration and purity samples. Data generated for all samples from same experiment.

RESULTS

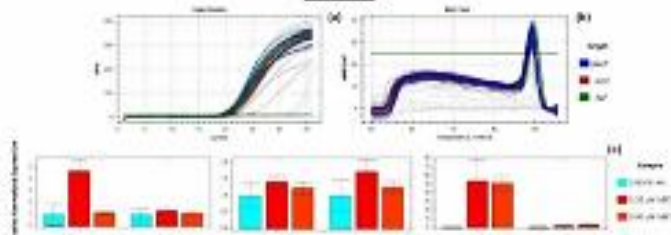


Figure 3. PCR with target primer amplification and relative normalized gene expression. (A) Representative image of PCR amplification results with target primer from our experiment. (B) Shows the corresponding gene ratios associated with the amplification curve. (C) Relative normalized gene expression of flagellar motility gene *flgC* and biofilm exopolysaccharide associated gene *polA*. Values are per gene 2. Results represent mean \pm standard deviation. Significance of *flgC* expression was not seen across all experiments.

CONCLUSION

- This data suggests an increase in the gene expression of flagellar related genes in the presence of hBD-2 consistent with biofilm inhibitor of PA through a quorum sensing pathway.
- Additional low time points are needed to better assess the effects of hBD-2 on the exopolysaccharide production which normally follows the cessation of flagellar expression.
- It might be worthwhile to probe other gene targets associated with the production versus biofilm state in PA cell differentiation.
- Further tests targeting the quorum signaling cascade could help further elucidate an interference of hBD-2 with biofilm formation in PA.

SIGNIFICANCE

- This research may help understand how to utilize hBD-2 and by spreading novel vaccines through induction of its expression and therapeutic drug targeting its mode of action against PA.

REFERENCES

1. *Pseudomonas aeruginosa*. In: *Microbiology: Principles and Concepts*. 9th ed. 2018. <https://doi.org/10.1016/j.mic.2018.05.001>

2. Berman, Kaitlin, et al. "Time to Manage *Pseudomonas Aeruginosa* Infections." *Drugs in Context*, vol. 7, Mar. 2019. doi:10.1016/j.drugcon.2019.01.001.

3. Portes, Edith. "Antimicrobial Peptides: Emerging Effective Molecules of Combat Against Infections." *World Journal of Microbiology*, vol. 3, 2011, pp. 45-51. doi:10.4236/wjmicro.2011.31005.

4. Mawardi, Ehab, and Kevin Paruchko. "Multi-Targeted Approach to the Bacterial Cell of Quorum Sensing." *DUP Academic*. EMUVE University Press, 12 June 2011. http://www.dup.academia.edu/4541514/10.1007/978-1-4419-1111-1_10

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