

TECHNICAL REPORT

12/18/2020

Microbiologic Properties of Tobramycin 1.2gram vials mixed with BASSA-GEL™ against selected pathogens was assessed and the results are conveyed here.

Disclosure: This study was funded by Richie’s Specialty Pharmacy (“RSP”) pursuant to its research agreement with the UH College of Pharmacy (“UHCOP”). RSP is owned solely by Richie Ray, R.Ph. and he serves as its Pharmacist-In-Charge. He is a 1996 graduate of the UHCOP and RSP has had a research agreement with UHCOP since May 2012. RSP promotes appropriate pharmacy practices and is an advocate for infection treatments based upon supporting clinical data such as that provided by this report.

Executive Summary: Tobramycin 1.2gram vials (1 vial) mixed with BASSA-GEL™ (“DRUG”) was tested against the identified pathogens and the results of these tests are reported as follows. Should there be only a “blue-line” reported that means the DRUG was so effective against the pathogen that the detection limit was below the assay of the experiment.

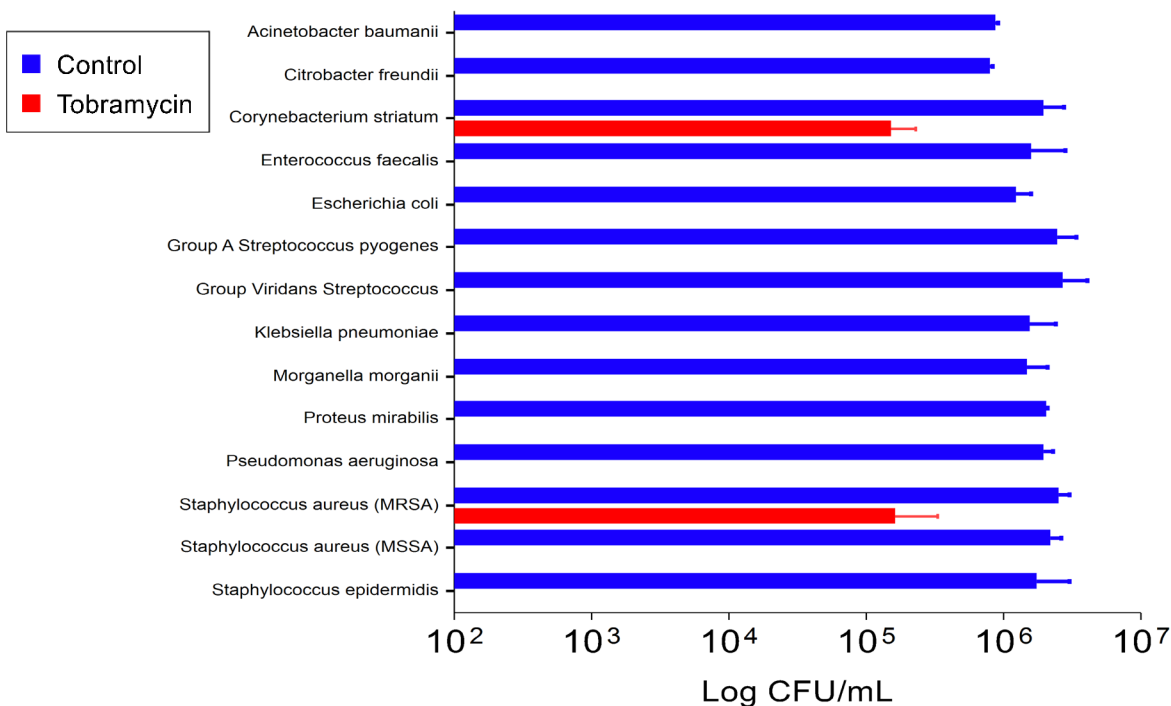
Methods overview: Methods for this laboratory study were adapted from Bearden *et al* and from FDA Docket No. FDA-1975-N-0012.^{1,2} All experiments were performed using the commercially available formulations. Reductions in bacterial counts between agents were determined.

Methods and Results:

Bacterial strains: Pathogens selected are defined in ATCC or CDC AR strains (Table 1, page 2).

Antimicrobial agents: Tobramycin 1.2 g vial (NDC 39822-0412-01) – 1 vial mixed with BASSA-GEL™

Experiment: Pre-sterilized discs were saturated with 1 x10⁷⁻⁸ CFU/mL of bacterial culture, allowed to incubate for 24 hours to mimic *ex vivo* wound infection, exposed to the gel/drug solution or positive control (phosphate buffer saline, PBS), and then incubated aerobically at 37°C for 24 hours. After this time, disks were washed, diluted, and then cultured onto blood agar plates for colony forming unit (CFU/mL) counts using serial dilution spread plate technique. The results are reported below (mean log CFU/mL ± standard error). As stated above in the executive summary, should there be only a “blue-line” reported that means the DRUG was so effective against the pathogen that the detection limit was below the assay of the experiment.



Interpretation: Tobramycin with BASSA-GEL™ was tested in a model mimicking a bandaged wound. The experiment demonstrated significant reductions in all gram-negative and most gram-positive bacterial species tested.

Table 1. Organisms Included in Testing

Organism	ATCC number
<i>Acinetobacter baumannii</i>	BAA747
<i>Citrobacter freundii</i>	8090
<i>Corynebacterium striatum</i>	BAA-1293
<i>Enterococcus faecalis</i>	BAA-29212
<i>Escherichia coli</i>	25922
<i>Klebsiella pneumoniae</i>	BAA-2524
<i>Streptococcus pyogenes</i>	19615
<i>Morganella morganii</i>	25830
<i>Proteus mirabilis</i>	CDC AR-29
<i>Pseudomonas aeruginosa</i>	27853
<i>Staphylococcus aureus (MSSA)</i>	29213
<i>Staphylococcus aureus (MRSA)</i>	BAA-41
<i>Staphylococcus epidermidis</i>	12228

References

1. Bearden DT, Allen GP, Christensen JM. Comparative in vitro activities of topical wound care products against community-associated methicillin-resistant *Staphylococcus aureus*. *J Antimicrob Chemother* 2008;62:769-72.
2. Huang DB, Okhuysen PC, Jiang ZD, DuPont HL. Enteroaggregative *Escherichia coli*: an emerging enteric pathogen. *Am J Gastroenterol* 2004;99:383-9.