Revolutionizing Treatment of Chronic Diseases

Brett Baker

Founder and Chief Innovation Officer



SAFETY OF TOPICAL PRAVIBISMANE IN DIABETIC FOOT ULCER INFECTIONS (PHASE 2A)

- The primary objective of the randomized, open label, controlled, multi-center Phase 2 study is to assess safety and tolerability, and secondarily efficacy of topical pravibismane to treat DFI
- Microbion completed enrollment of subjects; data analysis ongoing



ADVANTAGES OF TOPICAL PRAVIBISMANE FOR WOUND CARE IN WARFIGHTERS

- Topical pravibismane may address the military health concern in the treatment of antibiotic-resistant, **biofilm-forming infections in open injuries** contaminated with environmental materials
- Pravibismane has the potential to:
 - improve the rates of wound closure and resolution of infection
 - prevent and treat bacterial biofilm formation \checkmark
 - reduce the rate of infection-related amputations
 - \checkmark prevent extended courses of systemic antibiotic treatment
- No refrigeration required, means topical pravibismane can be a part of the medical field kits and could be continued in a hospital setting

SAFETY OF TOPICAL PRAVIBISMANE IN DIABETIC FOOT ULCER INFECTIONS (PHASE 1B)







- Pravibismane was **safe and well-tolerated** at all doses tested
- Pravibismane treated subjects exhibited ullet~3-fold reduction in wound size compared to placebo (85% vs 30%) and ~6-fold lower incidence of ulcer-related **limb amputations** (2.6% vs 15.4%)*



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* study was not powered to demonstrate statistical significance

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