MediWound

Next-Generation Enzymatic Therapeutics For Non-Surgical Tissue Repair



NexoBrid®

Disruptive Therapy for Burn Care FDA and EMA approved

- Approved in 40+ countries
- 18,000+ patients treated to date
- Poised to replace standard of care for eschar removal in severe burns
- Minimizes need for surgery and significantly reduces blood loss
- Topical application at bedside
- Preserves viable tissue and improves patient outcomes (scar quality and function)
- c-GMP sterile manufacturing facility to support global demand



EscharEx®

Bioactive Enzymatic Debridement Drug Candidate for Chronic Wounds

- Rapid, effective debridement and facilitation of wound closure
- Easy to use topical application for all patient settings
- Debrides chronic ulcers within 4-8 applications
- Promotes granulation tissue and reduces bacteria & biofilm
- Demonstrated superiority over SANTYL®
- Targets a $4B market
- De-risked program; based on 3 successful Phase II trials
  - Ongoing Phase III trial for venous leg ulcers (VLU)
  - DFU/PU – Trials preparation underway
- Research collaborations with Mölnlycke, Solventum, Medline, Essity, B. Braun, Coloplast, Convatec and MIMEDX

Global Collaborations

BARDA · European Commission · Mölnlycke · essity · convatec · MEDLINE · B|BRAUN SHARING EXPERTISE · solventum · KAKEN · VERICEL · Coloplast · MIMEDX

Validated Enzymatic Technology Platform

Solid Balance Sheet

$28.4M cash (as of September 30, 2026)

$17.3M financing in November 2026

$140M U.S. government support

No debt