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Kane Biotech to Present revyve® Clinical Data at Symposium on Advanced Wound Care (SAWC) Fall Conference

Clinical data to be presented demonstrates wound healing in complex wounds that were over two years old

WINNIPEG, Manitoba, Sept. 04, 2025 (GLOBE NEWSWIRE) -- Kane Biotech Inc. (TSX-V:KNE) ("Kane Biotech", "Kane" or the "Company") today announces that Interim Chief Executive Officer, Dr. Robert Huizinga, will be presenting at the Symposium on Advanced Wound Care ("SAWC") Fall conference taking place from September 3-6, 2025 in Las Vegas, Nevada.



US FDA 510(k) cleared revyve Antimicrobial Wound Gel and revyve Antimicrobial Wound Gel Spray

SAWC is a multidisciplinary community of wound care professionals — including physicians, nurses, podiatrists, physical therapists, researchers, administrators and wound care companies — that share the latest evidence-based practices, clinical innovations, and research aimed at improving outcomes in wound management.

Two presentations are being given at this meeting:

1. *Clinical Use of a Novel Thermo-reversible Antimicrobial Wound Gel to Fill Deep Cavity Wounds and Reduce Pain and Exudate Levels*

In this case study, revyve Antimicrobial Wound Gel was used on a two-year old non-healing Stage 4 sacral ulcer and a pain score of 10/10 in an elderly patient. Previous treatment included daily dressing changes using silver-based antimicrobial agents, gelling fibre wound fillers. This was a deep cavity wound requiring a thermoreversible gel to cover all of the wound surface. Use of revyve and daily dressing changes resulted in pain scores of 0, and wound size was decreased over three weeks. A reduction in *Klebsiella pneumonia* cultures was noted over the course of treatment. The patient remained on dressing changes with revyve over the two-month course of the study.

2. *Vibrational Debridement with a Novel Thermo-reversible Antimicrobial Wound Gel Turns a Non-healing Venous Leg Ulcer (VLU) to a Healing VLU*

In this case study, revyve Antimicrobial Wound Gel along with a vibrational debridement tool was used on an elderly female with a two-year old non-healing venous leg ulcer. This was a full thickness wound with slough, eschar and necrotic tissue present, alongside granulation tissue. Pain score was 5/10. Pain levels decreased with dressing changes, with the patient reporting a score of 0 from Week three onward. The wound showed progressive healing with a wound size reduction of 60% over eight weeks, along with decreased discharge.

Kane's US Food and Drug Administration (FDA) 510(k) cleared revyve Antimicrobial Wound Gel and Wound Gel Spray products will be exhibited at the conference. Dr. Huizinga will also be meeting with potential US distributors.

About Kane Biotech Inc. (TSX-V: KNE)

Kane Biotech is developing novel wound care treatments that disrupt biofilms and transform healing outcomes. Biofilms are one of the main contributors to antibiotic resistance in wounds which results in serious clinical outcomes and significant cost. revyve[®] addresses both biofilms and wound bacteria. revyve[®] Antimicrobial Wound Gel and revyve[®] Antimicrobial Wound Gel Spray are US FDA 510(k) cleared. revyve[®] Antimicrobial Wound Gel is Health Canada approved. To learn more about revyve, visit revyvegel.com or revyvegel.ca.

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A photo accompanying this announcement is available at
<https://www.globenewswire.com/NewsRoom/AttachmentNg/bd2cf932-40cc-4466-abbe-4bdc61611b29>



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Source: Kane Biotech Inc.