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# Kane Biotech Completes Enrollment in U.S. Case Series Studies of its revyve® Antimicrobial Wound Gel and Spray

WINNIPEG, Manitoba, Sept. 10, 2025 (GLOBE NEWSWIRE) -- Kane Biotech Inc. (TSX-V:KNE) ("Kane Biotech" or "Kane") today announces that it has enrolled 28 participants in its revyve® Antimicrobial Wound Gel and Spray U.S. Case Series Studies, exceeding its 25-participant target.



*US FDA 510(k) cleared revyve Antimicrobial Wound Gel*

For the clinical series, Kane engaged wound care and burn specialists in the U.S. Data is anticipated to be presented at various medical meetings in 2025 and 2026.

"Interest from clinicians and patients in the U.S. case series is encouraging," said Dr. Robert Huizinga, Interim CEO. "revyve products can meaningfully improve outcomes for those suffering from challenging wounds and burns."

## **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. Its revyve® Antimicrobial Wound Gel and revyve® Antimicrobial Wound Gel Spray are US FDA 510(k) cleared. To learn more about revyve, visit [revyvegel.com](https://revyvegel.com) or [revyvegel.ca](https://revyvegel.ca).

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A photo accompanying this announcement is available at <https://www.globenewswire.com/NewsRoom/AttachmentNg/29d77582-1218-4bbc-9c31-ebd8cec3ea69>



Source: Kane Biotech Inc.

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