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Kane Biotech Receives FDA Approval to Increase Dosage Allowance of its revyve™ Antimicrobial Wound Gel

Paves the Way for Kane's revyve™ Antimicrobial Wound Gel Spray

WINNIPEG, Manitoba, July 23, 2024 (GLOBE NEWSWIRE) -- Kane Biotech Inc. (TSX-V:KNE; OTCQB:KNBIF) ("Kane Biotech", "Kane" or the "Company") announces that the US Food and Drug Administration (FDA) has eliminated its usage limitation on the Company's 510(k) cleared revyve™ Antimicrobial Wound Gel ("revyve™"). Prior to the removal of this restriction, there was a 90 grams/month limit to the amount of revyve™ product that could be administered to patients.

This now clears the way for the introduction and extended use of Kane's revyve™ Antimicrobial Wound Gel Spray which is expected to be filled in spray cans in a higher quantity making it ideal for application on large wounds.

"This is yet another important development for Kane as it allows for increased use of the revyve™ product line throughout the standard of care in both prescription and over-the-counter (OTC) channels," said Marc Edwards, President & CEO.

About Kane Biotech

Kane Biotech Inc. is a biotechnology company engaged in the research, development and commercialization of technologies and products that prevent and remove microbial biofilms. Kane has a portfolio of biotechnologies, intellectual property (67 patents and patents pending, trade secrets and trademarks) and products developed by Kane's own biofilm research expertise and acquired from leading research institutions. DispersinB®, coactiv+™, coactiv+®, DermaKB™, DermaKB Biofilm™, and revyve™ are trademarks of Kane Biotech Inc. Kane is listed on the TSX Venture Exchange under the symbol "KNE" and on the OTCQB Venture Market under the symbol "KNBIF".

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Source: Kane Biotech Inc.