

**K222143 APIS (R), VERIS (TM)**Oct 13, 2023  
450 days to decisionK222143 · Product code: **FRO** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k222143/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dressing, Wound, Drug (FRO)
Date received	Jul 20, 2022
Decision date	Oct 13, 2023
Days to decision	450 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sweetbio, Inc.</b>
Location	Memphis, TN, US
Contact	Kayla Graff
510(k) history	2 submissions · 2 cleared · 2019-2023

**REGULATORY CONSULTANT**

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Consulting firm	<b>Kapstone Medical</b>
Contact	Carolyn Guthrie

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k222143/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](https://accessdata.fda.gov). - Generated May 26, 2026