FDA 510(k) Premarket Notification

K870352 RUBYCELL(TM) SYNTHETIC ISLAND DRESSING

CLEARED

Feb 26, 1987
28 days to decision

K870352 · Product code: **FRO** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k870352/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Dressing, Wound, Drug (FRO)
Date received	Jan 29, 1987
Decision date	Feb 26, 1987
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Bioderm, Inc.**
Location	Wheaton, IL, US
Contact	OLIVIER A DELANNOY
510(k) history	14 submissions · 10 cleared · 1987-1999

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k870352/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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