FDA 510(k) Premarket Notification

K896531 NOVADERM

CLEARED

Apr 13, 1990
148 days to decision

K896531 · Product code: **KMF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k896531/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bandage, Liquid (KMF)
Date received	Nov 16, 1989
Decision date	Apr 13, 1990
Days to decision	148 days
Third-party review	No

APPLICANT

Company	**Syncare Corp.**
Location	Charleston, SC, US
Contact	TODD DERBIN
510(k) history	1 submissions · 1 cleared · 1990-1990