FDA 510(k) Premarket Notification

K861285 SKIN MESH DEVICE

CLEARED

Apr 17, 1986
10 days to decision

K861285 · Product code: **GFD** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k861285/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dermatome (GFD)
Date received	Apr 7, 1986
Decision date	Apr 17, 1986
Days to decision	10 days
Third-party review	No

APPLICANT

Company	**Precision Modified Devices**
Location	Salt Lake City, UT, US
Contact	KATHY M SMITH
510(k) history	1 submissions · 1 cleared · 1986-1986