FDA 510(k) Premarket Notification

K990955 MODIFICATION OF:BIODERM FOAM WOUND DRESSING

CLEARED

May 18, 1999
57 days to decision

K990955 · Product code: **KMF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k990955/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bandage, Liquid (KMF)
Date received	Mar 22, 1999
Decision date	May 18, 1999
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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