FDA 510(k) Premarket Notification

K921022 DISPOSABLE CURETTE

CLEARED

Dec 3, 1992
275 days to decision

K921022 · Product code: **MDM** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k921022/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Manual, Surgical, General Use (MDM)
Date received	Mar 3, 1992
Decision date	Dec 3, 1992
Days to decision	275 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Dermtec, Inc.**
Location	Kershaw, SC, US
Contact	BREWER, JR.
510(k) history	1 submissions · 1 cleared · 1992-1992