FDA 510(k) Premarket Notification

K965256 DMS-1000C DERMOABRADER

CLEARED

Dec 9, 1996
115 days to decision

K965256 · Product code: **GFD** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k965256/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dermatome (GFD)
Date received	Aug 16, 1996
Decision date	Dec 9, 1996
Days to decision	115 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Mattioli Engineering, Srl**
Location	Florence, IT
Contact	GIAN FRANCO BERNABEI
510(k) history	2 submissions · 2 cleared · 1994-1996