FDA 510(k) Premarket Notification

K963204 DMS-1000C DERMOABRADER

CLEARED

Dec 9, 1996
116 days to decision

K963204 · Product code: **GFE** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k963204/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Brush, Dermabrasion, Powered (GFE)
Date received	Aug 15, 1996
Decision date	Dec 9, 1996
Days to decision	116 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Aesthetic Lasers, Inc.**
Location	Annapolis, MD, US
Contact	FRED DE JACMA
510(k) history	1 submissions · 1 cleared · 1996-1996