

K963204 DMS-1000C DERMOABRADERDec 9, 1996
116 days to decisionK963204 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k963204/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 4, 2026