

K965256 DMS-1000C DERMOABRADERDec 9, 1996
115 days to decisionK965256 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k965256/> 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