

K860925 DERMASPANDEREMar 28, 1986
17 days to decisionK860925 · Product code: **GFD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k860925/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dermatome (GFD)
Date received	Mar 11, 1986
Decision date	Mar 28, 1986
Days to decision	17 days
Third-party review	No

APPLICANT

Company	Mri Ventures, Inc.
Location	Kansas City, MO, US
Contact	A. E VANDEGRIFT
510(k) history	1 submissions · 1 cleared · 1986-1986

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k860925/> 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 8, 2026