

K963180 VIDAMED RF GENERATOR DATA RECORDEROct 10, 1996
57 days to decisionK963180 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k963180/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 14, 1996
Decision date	Oct 10, 1996
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Vidamed, Inc.
Location	Menlo Park, CA, US
Contact	NOEL MESSENGER
510(k) history	11 submissions · 11 cleared · 1995-2002

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