

**K955035 VIDAMED SERIES 7205 ELECTROSURGICAL
GENERATOR**Nov 29, 1995
27 days to decisionK955035 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k955035/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 2, 1995
Decision date	Nov 29, 1995
Days to decision	27 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k955035/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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