FDA 510(k) Premarket Notification

K955035 VIDAMED SERIES 7205 ELECTROSURGICAL GENERATOR

CLEARED

Nov 29, 1995
27 days to decision

K955035 · Product code: **GEI** · General & Plastic Surgery
Source: 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