

**K914066 ELECTROSURGICAL CUTTING/COAGULATION
DEVICE/ACCESS**Nov 7, 1991
57 days to decisionK914066 · Product code: **GEI** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k914066/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 11, 1991
Decision date	Nov 7, 1991
Days to decision	57 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Laparomed Corp.
Location	Irvine, CA, US
Contact	PAUL LUBOCK
510(k) history	10 submissions · 10 cleared · 1991-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k914066/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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