FDA 510(k) Premarket Notification

K914066 ELECTROSURGICAL CUTTING/COAGULATION DEVICE/ACCESS

CLEARED

Nov 7, 1991
57 days to decision

K914066 · 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