FDA 510(k) Premarket Notification

K904993 LAPAROSCOPIC ELECTROSURGICAL DEVICE

CLEARED

Jan 25, 1991
79 days to decision

K904993 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k904993/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 7, 1990
Decision date	Jan 25, 1991
Days to decision	79 days
Third-party review	No

APPLICANT

Company	**Everest Medical Corp.**
Location	Brooklyn Center, MN, US
Contact	DAVID J PARINS
510(k) history	21 submissions · 21 cleared · 1987-2000