FDA 510(k) Premarket Notification

K992045 ELECTROSURGICAL ELECTRODE FAMILY

CLEARED

Jan 27, 2000
224 days to decision

K992045 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k992045/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 17, 1999
Decision date	Jan 27, 2000
Days to decision	224 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Genesis Medical, Inc.**
Location	Sunnyvale, CA, US
Contact	THOMAS WEHMAN
510(k) history	1 submissions · 1 cleared · 2000-2000