FDA 510(k) Premarket Notification

K001151 ELECTROSURGICAL ELECTRODE FAMILY

CLEARED

Jun 27, 2000
78 days to decision

K001151 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k001151/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 10, 2000
Decision date	Jun 27, 2000
Days to decision	78 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Novasys Medical, Inc.**
Location	Sunnyvale, CA, US
Contact	THOMAS WEHMAN
510(k) history	5 submissions · 5 cleared · 2000-2004