FDA 510(k) Premarket Notification

K885200 600 & 900 SERIES ELECTRODES & 755 & 756 HANDLE

CLEARED

Jan 13, 1989
28 days to decision

K885200 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k885200/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 16, 1988
Decision date	Jan 13, 1989
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Olsen Electrosurgical, Inc.**
Location	Concord, CA, US
Contact	EUGENE OLSEN
510(k) history	11 submissions · 11 cleared · 1988-1994