FDA 510(k) Premarket Notification

K912781 ELECTROSHIELD

CLEARED

Dec 30, 1991
189 days to decision

K912781 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k912781/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 24, 1991
Decision date	Dec 30, 1991
Days to decision	189 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Electroscope, Inc.**
Location	Boulder, CO, US
Contact	ROGER C.ODELL
510(k) history	6 submissions · 6 cleared · 1991-2000