FDA 510(k) Premarket Notification

K981188 ELECTROSURGICAL CAUTERY PROBES

CLEARED

May 20, 1998
48 days to decision

K981188 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k981188/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 2, 1998
Decision date	May 20, 1998
Days to decision	48 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Microline Pentax, Inc.**
Location	Devers, MA, US
Contact	JACQUELINE E MASSE
510(k) history	12 submissions · 12 cleared · 1988-2002