FDA 510(k) Premarket Notification

K984185 MODIFICATION TO TAC-S MONOPOLAR CAUTERY PROBE FAMILY

CLEARED

Dec 16, 1998
23 days to decision

K984185 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k984185/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 23, 1998
Decision date	Dec 16, 1998
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Oratec Interventions, Inc.**
Location	Mountain View, CA, US
Contact	SHEILA RAMERMAN
510(k) history	24 submissions · 24 cleared · 1995-2002