FDA 510(k) Premarket Notification

K992408 ELECTROSURGICAL PROBES FAMILY: LIGAMENT CHISELS, MICRO LIGAMENT CHISELS, ABLATOR PROBES

CLEARED

Jul 30, 1999
10 days to decision

K992408 · Product code: **GEI** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k992408/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 20, 1999
Decision date	Jul 30, 1999
Days to decision	10 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