

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

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

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Device record: <https://www.510kdatabase.net/k992408/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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