

**K984185 MODIFICATION TO TAC-S MONOPOLAR CAUTERY  
PROBE FAMILY**

Dec 16, 1998  
23 days to decision

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

**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k984185/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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