

K972358 ORATEC MODEL ENDOTAC PROBEJan 23, 1998
212 days to decisionK972358 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k972358/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 25, 1997
Decision date	Jan 23, 1998
Days to decision	212 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k972358/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026