

**K911274 UNIPOLAR COAGULATOR PROBE SYSTEM**Jun 20, 1991  
90 days to decisionK911274 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k911274/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 22, 1991
Decision date	Jun 20, 1991
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>American Surgical Instruments Corp.</b>
Location	Washington, DC, US
Contact	DORSEY III
510(k) history	6 submissions · 5 cleared · 1990-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k911274/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 30, 2026