

**K901302 BIPOLAR POLYPECTOMY SYSTEM (TM)**Jun 6, 1990  
78 days to decisionK901302 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k901302/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 20, 1990
Decision date	Jun 6, 1990
Days to decision	78 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Devices, Inc.</b>
Location	Mchenry, IL, US
Contact	SAMUEL DICKSTEIN
510(k) history	49 submissions · 47 cleared · 1977-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901302/>; 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 May 21, 2026