

**K904993 LAPAROSCOPIC ELECTROSURGICAL DEVICE**Jan 25, 1991  
79 days to decisionK904993 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k904993/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Nov 7, 1990
Decision date	Jan 25, 1991
Days to decision	79 days
Third-party review	No

**APPLICANT**

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Company	<b>Everest Medical Corp.</b>
Location	Brooklyn Center, MN, US
Contact	DAVID J PARINS
510(k) history	21 submissions · 21 cleared · 1987-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904993/>; 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 29, 2026