

**K910334 ELECTROCAUTERY PROBE**Feb 21, 1991  
27 days to decisionK910334 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k910334/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 25, 1991
Decision date	Feb 21, 1991
Days to decision	27 days
Third-party review	No

**APPLICANT**

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Company	<b>Wiltek Medical, Inc.</b>
Location	Winston-Salem, NC, US
Contact	JON S WILSON
510(k) history	21 submissions · 21 cleared · 1989-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k910334/>; 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 July 1, 2026