

**K912781 ELECTROSHIELD**Dec 30, 1991  
189 days to decisionK912781 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k912781/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 24, 1991
Decision date	Dec 30, 1991
Days to decision	189 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Electroscope, Inc.</b>
Location	Boulder, CO, US
Contact	ROGER C.ODELL
510(k) history	6 submissions · 6 cleared · 1991-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k912781/> 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 3, 2026