

**K885200 600 & 900 SERIES ELECTRODES & 755 & 756 HANDLE**Jan 13, 1989  
28 days to decisionK885200 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k885200/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 16, 1988
Decision date	Jan 13, 1989
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Olsen Electrosurgical, Inc.</b>
Location	Concord, CA, US
Contact	EUGENE OLSEN
510(k) history	11 submissions · 11 cleared · 1988-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k885200/>; 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