

**K033880 ELECTRO-LUBE**Mar 10, 2004  
86 days to decisionK033880 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k033880/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Dec 15, 2003
Decision date	Mar 10, 2004
Days to decision	86 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Mectra Labs, Inc.</b>
Location	Bloomfield, IN, US
Contact	CHARLES E ALLGOOD JR.
510(k) history	9 submissions · 9 cleared · 1992-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k033880/> 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 14, 2026