

**K101711 CERULEAU ELECTROSURGICAL PROBE**Jul 1, 2010  
13 days to decisionK101711 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k101711/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 18, 2010
Decision date	Jul 1, 2010
Days to decision	13 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nuortho Surgical, Inc.</b>
Location	Newington, NH, US
Contact	MARTIN J LEIGHTON
510(k) history	1 submissions · 1 cleared · 2010-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k101711/> 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 20, 2026