

**K012679 REPROCESSED RF ARTHROSCOPY PROBES**Nov 8, 2001  
86 days to decisionK012679 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k012679/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 14, 2001
Decision date	Nov 8, 2001
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sterilmed, Inc.</b>
Location	Plymouth, MN, US
Contact	PATRICK FLEISCHHACKER
510(k) history	64 submissions · 64 cleared · 2001-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k012679/> 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 May 19, 2026