

**K222958 mJOULE RF System and Accessories**Apr 23, 2024  
574 days to decisionK222958 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k222958/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 27, 2022
Decision date	Apr 23, 2024
Days to decision	574 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sciton, Inc</b>
Location	Palo Alto, CA, US
Contact	Jay Patel
510(k) history	23 submissions · 23 cleared · 2000-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k222958/> 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 13, 2026