

**K170610 PlasmaBlade T**Apr 13, 2017  
43 days to decisionK170610 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k170610/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 1, 2017
Decision date	Apr 13, 2017
Days to decision	43 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Advanced Energy</b>
Location	Portsmouth, NH, US
Contact	Lydia Sakakeeny
510(k) history	8 submissions · 8 cleared · 2014-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k170610/> 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 28, 2026