

**K241085 Valleylab <sup>TM</sup>FT10 Energy Platform Software Version 5.0  
(VLFT10GEN VLFT10FXGEN VLFT10LSGEN)**Jul 12, 2024  
84 days to decisionK241085 · Product code: **GEI** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k241085/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 19, 2024
Decision date	Jul 12, 2024
Days to decision	84 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Covidien, LLC</b>
Location	Mansfield, MA, US
Contact	Aliva Mahunta
510(k) history	88 submissions · 85 cleared · 2010-2026

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k241085/> 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 14, 2026