

**K202818 Voyant Fine Fusion Device (EB230)**Dec 16, 2021  
448 days to decisionK202818 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k202818/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 24, 2020
Decision date	Dec 16, 2021
Days to decision	448 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Applied Medical</b>
Location	Fort Worth, TX, US
Contact	Blake Stacy
510(k) history	14 submissions · 14 cleared · 1985-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k202818/> 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 26, 2026