

**K222284 Voyant Maryland Fusion device with Single-Step Activation (EB212, EB213, EB214)**Oct 14, 2022  
74 days to decisionK222284 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k222284/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Aug 1, 2022
Decision date	Oct 14, 2022
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Applied Medical Resources Corporation</b>
Location	Rancho Santa, CA, US
Contact	Blake Stacy
Website	<a href="https://www.appliedmedical.com/">https://www.appliedmedical.com/</a>
510(k) history	12 submissions · 12 cleared · 2017-2026

Applied Medical Resources Corporation is a medical device manufacturer specializing in minimally invasive surgical instruments and access systems, with a manufacturing facility in Rancho Santa, California. The company develops a broad portfolio of surgical tools for general, plastic, and gynecological procedures. Applied Medical has received FDA 510(k) clearances from total submissions since 2017. The company's regulatory focus centers on General & Plastic Surgery devices, which represent the majority of its cleared portfolio. The most recent clearance was in 2026, demons...

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k222284/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 14, 2026