

**K252367 InbellaMulti System**Oct 15, 2025  
77 days to decisionK252367 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k252367/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jul 30, 2025
Decision date	Oct 15, 2025
Days to decision	77 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Inbella Medical, Inc.</b>
Location	Richmond Hill, CA
Contact	Michael Kreindel
510(k) history	4 submissions · 4 cleared · 2025-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Hogan Lovells US LLP</b>
Contact	Janice Hogan

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k252367/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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