

**K240070 Profound Matrix**Jul 1, 2024  
174 days to decisionK240070 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k240070/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 9, 2024
Decision date	Jul 1, 2024
Days to decision	174 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Candela Corporation</b>
Location	Wayland, MA, US
Contact	Rina Ordonez
510(k) history	8 submissions · 8 cleared · 2015-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k240070/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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