

**K233002 BEED 2,5, BEED 3,8**Mar 20, 2024  
180 days to decisionK233002 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k233002/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 22, 2023
Decision date	Mar 20, 2024
Days to decision	180 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Excelsior Resources, LLC</b>
Location	Ft. Lauderdale, FL, US
Contact	Christopher P Weber
510(k) history	1 submissions · 1 cleared · 2024-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k233002/> 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