

**K233374 Secret Pro**Dec 14, 2023  
73 days to decisionK233374 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k233374/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 2, 2023
Decision date	Dec 14, 2023
Days to decision	73 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Ilooda Co.,, Ltd.</b>
Location	Gwonseon-Gu, Suwon-Si, KR
Contact	Kathy Maynor
510(k) history	16 submissions · 16 cleared · 2015-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k233374/> 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