

**K223107 MOLLI 2**Jan 18, 2023  
110 days to decisionK223107 · Product code: **NEU** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k223107/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Marker, Radiographic, Implantable (NEU)
Date received	Sep 30, 2022
Decision date	Jan 18, 2023
Days to decision	110 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Molli Surgical, Inc.</b>
Location	Toronto, CA
Contact	Joseph De Croos
510(k) history	5 submissions · 5 cleared · 2021-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k223107/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 25, 2026