

**K240132 EmoLED (v. 2-USA)**Mar 12, 2024  
55 days to decisionK240132 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k240132/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jan 17, 2024
Decision date	Mar 12, 2024
Days to decision	55 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Emoled, Srl</b>
Location	Sesto Fiorentino, IT
Contact	Lorenzo Targetti
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/k240132/> 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 13, 2026