

**K230274 Medical Diode Laser (M2)**Mar 31, 2023  
58 days to decisionK230274 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k230274/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 1, 2023
Decision date	Mar 31, 2023
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Wuhan Pioon Technology Co., Ltd.</b>
Location	Wuhan, CN
Contact	Feng Zhang
510(k) history	7 submissions · 7 cleared · 2023-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k230274/> 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 26, 2026