

K260307 Diode Laser Therapy Systems (V19)Apr 15, 2026
75 days to decisionK260307 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k260307/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jan 30, 2026
Decision date	Apr 15, 2026
Days to decision	75 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Weifang Mingliang Electronics Co., Ltd.
Location	Weifang, CN
Contact	MingLiang Li
510(k) history	3 submissions · 3 cleared · 2020-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k260307/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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