

K200118 Diode Laser Therapy DeviceApr 17, 2020
87 days to decisionK200118 · Product code: **GEX** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k200118/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jan 21, 2020
Decision date	Apr 17, 2020
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Shanghai Apolo Medical Technology Co., Ltd.
Location	Shanghai, CN
Contact	Felix Li
510(k) history	13 submissions · 13 cleared · 2019-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k200118/> Data sourced from FDA 510(k) public records (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. - Generated May 26, 2026