

K233157 VaporMAX LDDMay 3, 2024
219 days to decisionK233157 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k233157/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 27, 2023
Decision date	May 3, 2024
Days to decision	219 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	MOJo LDD; MegaJOULE LDD

APPLICANT

Company	Innovaquartz, LLC
Location	Phoenix, AZ, US
Contact	Stephen Griffin
510(k) history	4 submissions · 4 cleared · 2014-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k233157/> 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 14, 2026