

K241771 LiftendoSep 16, 2024
88 days to decisionK241771 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k241771/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jun 20, 2024
Decision date	Sep 16, 2024
Days to decision	88 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	Liftendo 2.0

APPLICANT

Company	Medical San Ind?stria DE Equipamentos M?dicos Ltda.
Location	Lajeado, BR
Contact	Francis Laion de Padua
510(k) history	2 submissions · 2 cleared · 2023-2024

REGULATORY CONSULTANT

Consulting firm	United Regulatory & Logistics
Contact	Rodrigo Abreu

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)**510k Database** - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k241771/> 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