

**K253219 LeafPure-Echo (LM-LPNdA)**Nov 25, 2025  
57 days to decisionK253219 · Product code: **GEX** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k253219/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Sep 29, 2025
Decision date	Nov 25, 2025
Days to decision	57 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Shenzhen Leaflife Technology Co., Ltd.</b>
Location	Shenzhen, CN
Contact	Qiang Cheng
510(k) history	7 submissions · 7 cleared · 2020-2025

**REGULATORY CONSULTANT**

---

Consulting firm	<b>Aestheticcert Compliance, LLC</b>
Contact	Asher Zeng

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.net](https://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k253219/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 14, 2026