

K233036 Ultrasonic Surgical SystemJun 18, 2024
267 days to decisionK233036 · Product code: **LFL** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k233036/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Ultrasonic Surgical (LFL)
Date received	Sep 25, 2023
Decision date	Jun 18, 2024
Days to decision	267 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Hunan Handlike Minimally Invasive Surgery Co., Ltd.
Location	Changsha, CN
Contact	Yang Bihong
510(k) history	2 submissions · 2 cleared · 2018-2024

REGULATORY CONSULTANT

Consulting firm	Share Info (Guangzhou) Medical Consultant , Ltd.
Contact	Lee Cassie

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/k233036/> 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 13, 2026