

K240434 RELIEEV Suction Curette (Flexible 3.0/ Standard 3.6)Sep 25, 2024
224 days to decisionK240434 · Product code: **HHK** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k240434/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Curette, Suction, Endometrial (and Accessories) (HHK)
Date received	Feb 14, 2024
Decision date	Sep 25, 2024
Days to decision	224 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Li Medical Corporation , Ltd.
Location	Xizhi Dist., TW
Contact	Jago Chen
510(k) history	3 submissions · 3 cleared · 2024-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k240434/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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