FDA 510(k) Premarket Notification

K240434 RELIEEV Suction Curette (Flexible 3.0/ Standard 3.6)

CLEARED

Sep 25, 2024
224 days to decision

K240434 · Product code: **HHK** · Obstetrics & Gynecology
Source: 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