FDA 510(k) Premarket Notification

K241627 Safety Lancet (XXXV)

CLEARED

Jun 20, 2024
14 days to decision

K241627 · Product code: **FMK** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k241627/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Single Use Only Blood Lancet With An Integral Sharps Injury Prevention Feature (FMK)
Date received	Jun 6, 2024
Decision date	Jun 20, 2024
Days to decision	14 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Tianjin Huahong Technology Co., Ltd.**
Location	Tianjin, CN
Contact	Ningning Wang
510(k) history	11 submissions · 11 cleared · 2021-2026

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k241627/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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