

K222055 Disposable Sterile LancetNov 15, 2022
126 days to decisionK222055 · Product code: **FMK** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k222055/>**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	Jul 12, 2022
Decision date	Nov 15, 2022
Days to decision	126 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Tianjin Rilifine Medical Device Co., Ltd.
Location	Tianjin, CN
Contact	Qiusheng Jiang
510(k) history	2 submissions · 2 cleared · 2022-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k222055/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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