

K223313 Disposable Safety LancetsJan 23, 2023
87 days to decisionK223313 · Product code: **FMK** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k223313/>**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	Oct 28, 2022
Decision date	Jan 23, 2023
Days to decision	87 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Huaian Hening Medical Instruments Co., Ltd.
Location	Huaian, CN
Contact	Zhengcan Da
510(k) history	2 submissions · 2 cleared · 2023-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k223313/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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