

K222376 Lianfa Safety Lancet (Five Models: NPA, PA, PA2, TPA, and APA)Oct 3, 2022
59 days to decisionK222376 · Product code: **FMK** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k222376/>**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	Aug 5, 2022
Decision date	Oct 3, 2022
Days to decision	59 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Shandong Lianfa Medical Plastic Products Co. , Ltd.
Location	Jinan, CN
Contact	Charles Shen
510(k) history	2 submissions · 2 cleared · 2022-2022

REGULATORY CONSULTANT

Consulting firm	Manton Business and Technology Services
Contact	Charles Shen

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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