

**K192666 Promisemed Blood Lancet, VeriFine Safety Lancet, VeriFine Mini-Safety Lancet**

Dec 2, 2019  
68 days to decision

K192666 · Product code: **FMK** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k192666/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Single Use Only Blood Lancet With An Integral Sharps Injury Prevention Feature (FMK)
Date received	Sep 25, 2019
Decision date	Dec 2, 2019
Days to decision	68 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Promisemed Hangzhou Meditech Co., Ltd.</b>
Location	Hangzhou, CN
Contact	Zearou Yang
510(k) history	34 submissions · 34 cleared · 2017-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k192666/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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