

K202682 Promisemed Triple Safety Pen NeedleNov 24, 2020
70 days to decisionK202682 · Product code: **FMI** · General Hospital
Source: <https://www.510kdatabase.net/k202682/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Sep 15, 2020
Decision date	Nov 24, 2020
Days to decision	70 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Promisemed Hangzhou Meditech Co., Ltd.
Location	Hangzhou, CN
Contact	Zearou Yang
510(k) history	35 submissions · 35 cleared · 2017-2026

REGULATORY CONSULTANT

Consulting firm	Vee Care (Asia) Limited
Contact	Wei Shan Hsu

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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