

K213004 Needle Free ConnectorMay 10, 2022
232 days to decisionK213004 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k213004/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 20, 2021
Decision date	May 10, 2022
Days to decision	232 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Medcaptain Life Science Co., Ltd.
Location	Shenzhen, CN
Contact	David Xia
510(k) history	5 submissions · 5 cleared · 2021-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k213004/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026