

K253142 Arm Blood Pressure Monitor (MJ1D,MJ1DS,MJ3D,MJ5D,MJ6D,MJ8D,RN3D,MJ4D,RN1D,RN2D)Jan 9, 2026
106 days to decisionK253142 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k253142/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Sep 25, 2025
Decision date	Jan 9, 2026
Days to decision	106 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Ningbo Ranor Medical Technology Co., Ltd.
Location	Ningbo, CN
Contact	Jia'nan Xu
510(k) history	1 submissions · 1 cleared · 2026-2026

REGULATORY CONSULTANT

Consulting firm	Shanghai Truthful Information Technology Co., Ltd.
Contact	Boyle Wang

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k253142/> 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 13, 2026