

K172221 VTRUST Vital Signs Monitor, Model: TD-2300Jun 6, 2018
317 days to decisionK172221 · Product code: **MWI** · Cardiovascular
Source: <https://www.510kdatabase.net/k172221/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Jul 24, 2017
Decision date	Jun 6, 2018
Days to decision	317 days
Third-party review	No
Summary / Statement	Summary
Other names	FORA Vital Signs Monitor, Model: VSM100

APPLICANT

Company	Taidoc Technology Corporation
Location	Taipei, County, TW
Contact	Anne Kuo
510(k) history	123 submissions · 123 cleared · 2004-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k172221/> 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 30, 2026