FDA 510(k) Premarket Notification

K172221 VTRUST Vital Signs Monitor, Model: TD-2300

CLEARED

Jun 6, 2018
317 days to decision

K172221 · 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