FDA 510(k) Premarket Notification

K150869 Checkme Pro Health Monitor

CLEARED

Dec 10, 2015
253 days to decision

K150869 · Product code: **MWI** · Cardiovascular
Source: https://www.510kdatabase.net/k150869/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Apr 1, 2015
Decision date	Dec 10, 2015
Days to decision	253 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Viatom Technology Co., Ltd.**
Location	Nanshan, Shenzhen, CN
Contact	Zhou Saixin
510(k) history	1 submissions · 1 cleared · 2015-2015