FDA 510(k) Premarket Notification

K170712 Accutorr 7/VS-900 Vital Signs Monitor

CLEARED

Aug 2, 2017
146 days to decision

K170712 · Product code: **MWI** · Cardiovascular
Source: https://www.510kdatabase.net/k170712/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Mar 9, 2017
Decision date	Aug 2, 2017
Days to decision	146 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Shenzhen Mindray Bio-Meidcal Electronics Co., Ltd.**
Location	Great Neck, NY, US
Contact	Yanhong Bai
510(k) history	9 submissions · 9 cleared · 2012-2022