FDA 510(k) Premarket Notification

K123711 PATIENT MONITOR

CLEARED

Oct 1, 2013
301 days to decision

K123711 · Product code: **MWI** · Cardiovascular
Source: https://www.510kdatabase.net/k123711/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Dec 4, 2012
Decision date	Oct 1, 2013
Days to decision	301 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Shenzhen Creative Industry Co., Ltd.**
Location	Flintville, TN, US
Contact	CHARLIE MACK
510(k) history	11 submissions · 11 cleared · 2007-2021