FDA 510(k) Premarket Notification

K130056 PATIENT MONITOR

CLEARED

Apr 11, 2013
92 days to decision

K130056 · Product code: **MWI** · Cardiovascular
Source: https://www.510kdatabase.net/k130056/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Jan 9, 2013
Decision date	Apr 11, 2013
Days to decision	92 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ld Technology, LLC**
Location	Miami, FL, US
Contact	ALBERT MAAREK
510(k) history	14 submissions · 14 cleared · 2009-2020