

K130056 PATIENT MONITORApr 11, 2013
92 days to decisionK130056 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k130056/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 19, 2026