

K100217 LUCON M-SERIES PATIENT MONITOR, MODELS M20 AND M30

Jun 8, 2010
134 days to decision

K100217 · Product code: **MHX** · Cardiovascular
Source: <https://www.510kdatabase.net/k100217/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jan 25, 2010
Decision date	Jun 8, 2010
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Mediana Co., Ltd.
Location	Flintville, TN, US
Contact	CHARLIE MACK
510(k) history	10 submissions · 10 cleared · 2005-2020

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k100217/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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