

K132033 BM3/BM3 PLUSDec 4, 2013
156 days to decisionK132033 · Product code: **MWI** · Cardiovascular
Source: <https://www.510kdatabase.net/k132033/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Jul 1, 2013
Decision date	Dec 4, 2013
Days to decision	156 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Noblitt & Rueland
Location	Irvine, CA, US
Contact	MARC GOODMAN
510(k) history	1 submissions · 1 cleared · 2013-2013

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