

**K132026 V SERIES MONITORING SYSTEM, 12.1 DCU/V12 AND
21.0 DCU/V21**Dec 20, 2013
172 days to decisionK132026 · Product code: **MHX** · Neurology
Source: <https://www.510kdatabase.net/k132026/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jul 1, 2013
Decision date	Dec 20, 2013
Days to decision	172 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Mindray DS USA, Inc.
Location	Mahwh, NJ, US
Contact	RUSSELL OLSEN
510(k) history	3 submissions · 3 cleared · 2010-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k132026/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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