

**K102004 ENDEAVOUR MONITORING SYSTEM**Sep 10, 2010  
57 days to decisionK102004 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k102004/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Jul 15, 2010
Decision date	Sep 10, 2010
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k102004/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 24, 2026