

**K110503 AIRSTRIP REMOTE PATIENT MONITORING (RPM)**Mar 10, 2011  
16 days to decisionK110503 · Product code: **MWI** · CardiovascularSource: <https://www.510kdatabase.net/k110503/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Feb 22, 2011
Decision date	Mar 10, 2011
Days to decision	16 days
Third-party review	Yes
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Airstrip Technologies, LP</b>
Location	San Antonio, TX, US
Contact	ANDY MILLER
510(k) history	10 submissions · 10 cleared · 2009-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k110503/> 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 June 7, 2026