

**K020494 MODIFICATION TO M2376A DEVICELINK SYSTEM**Feb 21, 2002  
8 days to decisionK020494 · Product code: **MWI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k020494/>**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 13, 2002
Decision date	Feb 21, 2002
Days to decision	8 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Philips Medical Systems, Inc.</b>
Location	Mchenry, IL, US
Contact	MIKE HUDON
510(k) history	111 submissions · 110 cleared · 1983-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020494/> 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 17, 2026