

**K121367 ZOLL PROPAQ XM**Jun 21, 2012  
45 days to decisionK121367 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k121367/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	May 7, 2012
Decision date	Jun 21, 2012
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>ZOLL Medical Corporation</b>
Location	Chelmsford, MA, US
Contact	CHARLES W KOLIFRATH
510(k) history	30 submissions · 30 cleared · 2005-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121367/> 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 5, 2026