

**K920377 MORTARA INSTRUMENT MODEL ECG PREVU**Apr 28, 1992  
90 days to decisionK920377 · Product code: **LOS** · CardiovascularSource: <https://www.510kdatabase.net/k920377/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Jan 29, 1992
Decision date	Apr 28, 1992
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Mortara Instrument, Inc.</b>
Location	Walker, MI, US
Contact	JOHN PRIMOZICH
510(k) history	51 submissions · 51 cleared · 1983-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920377/>, 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 19, 2026