

**K970182 DINAMAP COMPACT MONITOR**Aug 18, 1997  
213 days to decisionK970182 · Product code: **MHX** · CardiovascularSource: <https://www.510kdatabase.net/k970182/>**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	Jan 17, 1997
Decision date	Aug 18, 1997
Days to decision	213 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Johnson &amp; Johnson Medical, Inc.</b>
Location	Arlington, TX, US
Contact	DARLENE T KORAB
510(k) history	53 submissions · 50 cleared · 1990-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k970182/> 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 28, 2026