

**K003940 2010 PLUS HOLTER FOR WINDOWS**Feb 8, 2001  
49 days to decisionK003940 · Product code: **MLO** · Cardiovascular  
Source: <https://www.510kdatabase.net/k003940/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrocardiograph, Ambulatory, With Analysis Algorithm (MLO)
Date received	Dec 21, 2000
Decision date	Feb 8, 2001
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Agilent Technologies</b>
Location	Walker, MI, US
Contact	GRETEL LUMLEY
Website	<a href="http://www.agilent.com">http://www.agilent.com</a>
510(k) history	13 submissions · 13 cleared · 1982-2001

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k003940/> 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 25, 2026