

**K792312 HEART RATE MONITOR MD-1**Nov 21, 1979  
5 days to decisionK792312 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k792312/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Nov 16, 1979
Decision date	Nov 21, 1979
Days to decision	5 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Devices, Inc.</b>
Location	Mchenry, IL, US
510(k) history	49 submissions · 47 cleared · 1977-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k792312/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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