

**K024283 CARDIODIRECT, MODELS 12M, 12I, 12S**Aug 1, 2003  
221 days to decisionK024283 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k024283/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Dec 23, 2002
Decision date	Aug 1, 2003
Days to decision	221 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Reynolds Medical , Ltd.</b>
Location	Mchenry, IL, US
Contact	GEORGE MYERS
510(k) history	12 submissions · 12 cleared · 1982-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k024283/> 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 July 3, 2026