

**K842007 CENTRIMED SYSTEM 1**Aug 3, 1984  
78 days to decisionK842007 · Product code: **KFM** · CardiovascularSource: <https://www.510kdatabase.net/k842007/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Blood, Cardiopulmonary Bypass, Non-roller Type (KFM)
Date received	May 17, 1984
Decision date	Aug 3, 1984
Days to decision	78 days
Third-party review	No

**APPLICANT**

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Company	<b>Centrimed Corp.</b>
Location	Walker, MI, US
510(k) history	4 submissions · 4 cleared · 1984-1985

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

Device record: <https://www.510kdatabase.net/k842007/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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