

**K972962 3M CDI BLOOD PARAMETER MONITORING SYSTEM  
500**Nov 6, 1997  
87 days to decisionK972962 · Product code: **DRY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k972962/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Blood-gas, On-line, Cardiopulmonary Bypass (DRY)
Date received	Aug 11, 1997
Decision date	Nov 6, 1997
Days to decision	87 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>3M Company</b>
Location	White City, OR, US
Contact	ANNE BUTEYN
Website	<a href="http://www.3m.com/">http://www.3m.com/</a>
510(k) history	331 submissions · 322 cleared · 1976-2025

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