

**K931968 CDI 100 EXTRACORPOREAL HEMATOCRIT/OXYGEN SATURATION**

Nov 10, 1993  
203 days to decision

K931968 · Product code: **DRY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k931968/>

**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	Apr 21, 1993
Decision date	Nov 10, 1993
Days to decision	203 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>3M Health Care, Ltd.</b>
Location	Irvine, CA, US
Contact	STEVEN ARICK
510(k) history	49 submissions · 49 cleared · 1990-1996

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

Device record: <https://www.510kdatabase.net/k931968/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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