

**K871221 CDI(TM) EXTRACORPOREAL BLOOD GAS SYSTEM
300**Jun 5, 1987
70 days to decisionK871221 · Product code: **DRY** · Cardiovascular
Source: <https://www.510kdatabase.net/k871221/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Blood-gas, On-line, Cardiopulmonary Bypass (DRY)
Date received	Mar 27, 1987
Decision date	Jun 5, 1987
Days to decision	70 days
Third-party review	No

APPLICANT

Company	Cardiovascular Devices, Inc.
Location	Walker, MI, US
Contact	STEVEN ARICK
510(k) history	8 submissions · 8 cleared · 1982-1989

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k871221/> Data sourced from FDA 510(k) public records (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. - Generated May 27, 2026