

**K953821 BIOCOR 200 HARDSHELL VENOUS RESERVIOR**Mar 7, 1996  
205 days to decisionK953821 · Product code: **DTN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k953821/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reservoir, Blood, Cardiopulmonary Bypass (DTN)
Date received	Aug 15, 1995
Decision date	Mar 7, 1996
Days to decision	205 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Minntech Corp.</b>
Location	Minneapolis, MN, US
Contact	LEROY J FISCHBACH
510(k) history	33 submissions · 33 cleared · 1987-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k953821/> 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 June 13, 2026