FDA 510(k) Premarket Notification

K953821 BIOCOR 200 HARDSHELL VENOUS RESERVIOR

CLEARED

Mar 7, 1996
205 days to decision

K953821 · Product code: **DTN** · Cardiovascular
Source: https://www.510kdatabase.net/k953821/

SUBMISSION DETAILS

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

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