FDA 510(k) Premarket Notification

K031827 ULTRAFLEX VENOUS CANNULA, MODEL 970XX AND CB970XX

CLEARED

Jul 2, 2003
19 days to decision

K031827 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k031827/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Jun 13, 2003
Decision date	Jul 2, 2003
Days to decision	19 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronic Perfusion Systems**
Location	Minneapolis, MN, US
Contact	PREETI JAIN
510(k) history	29 submissions · 29 cleared · 2000-2024

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k031827/ 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 14, 2026