FDA 510(k) Premarket Notification

K972962 3M CDI BLOOD PARAMETER MONITORING SYSTEM 500

CLEARED

Nov 6, 1997
87 days to decision

K972962 · Product code: **DRY** · Cardiovascular
Source: https://www.510kdatabase.net/k972962/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Blood-gas, On-line, Cardiopulmonary Bypass (DRY)
Date received	Aug 11, 1997
Decision date	Nov 6, 1997
Days to decision	87 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**3M Company**
Location	White City, OR, US
Contact	ANNE BUTEYN
Website	http://www.3m.com/
510(k) history	331 submissions · 322 cleared · 1976-2025