FDA 510(k) Premarket Notification

K951088 QUEST RETROGRADE CARDIPLEGIA CANNULA-RCCS

CLEARED

Jun 14, 1995
97 days to decision

K951088 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k951088/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 9, 1995
Decision date	Jun 14, 1995
Days to decision	97 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Quest Medical, Inc.**
Location	Walker, MI, US
Contact	DREW JOHNSON
510(k) history	39 submissions · 39 cleared · 1980-2024