FDA 510(k) Premarket Notification

K896636 RMI ANTEPLEGIA CANNULA

CLEARED

Feb 15, 1990
85 days to decision

K896636 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k896636/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Nov 22, 1989
Decision date	Feb 15, 1990
Days to decision	85 days
Third-party review	No

APPLICANT

Company	**Research Medical, Inc.**
Location	Mchenry, IL, US
Contact	MICHAEL N KELLY
510(k) history	35 submissions · 35 cleared · 1984-1997