FDA 510(k) Premarket Notification

K770431 CATHETER, TWO STAGE VENOUS RETURN

CLEARED

Mar 16, 1977
9 days to decision

K770431 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k770431/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Mar 7, 1977
Decision date	Mar 16, 1977
Days to decision	9 days
Third-party review	No

APPLICANT

Company	**3M Health Care, Sarns**
Location	Mchenry, IL, US
510(k) history	76 submissions · 76 cleared · 1976-1996