FDA 510(k) Premarket Notification

K860136 BARD-PARKER CAROTID SHUNT

CLEARED

Jun 25, 1986
162 days to decision

K860136 · Product code: **DWF** · Cardiovascular
Source: https://www.510kdatabase.net/k860136/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Jan 14, 1986
Decision date	Jun 25, 1986
Days to decision	162 days
Third-party review	No

APPLICANT

Company	**Bd Becton Dickinson Vacutainer Systems Preanalytic**
Location	Washington, DC, US
Contact	RUSSELL ARNSBERGER
510(k) history	632 submissions · 625 cleared · 1976-2001