FDA 510(k) Premarket Notification

K885221 MULTILUMEN CENTRAL VENOUS CATHETER

CLEARED

Dec 28, 1989
373 days to decision

K885221 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k885221/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Dec 20, 1988
Decision date	Dec 28, 1989
Days to decision	373 days
Third-party review	No

APPLICANT

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