FDA 510(k) Premarket Notification

K883933 INTRODUCING CATHETER

CLEARED

Dec 2, 1988
78 days to decision

K883933 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k883933/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Sep 15, 1988
Decision date	Dec 2, 1988
Days to decision	78 days
Third-party review	No

APPLICANT

Company	**Devices For Vascular Intervention, Inc.**
Location	Redwood City, CA, US
Contact	DIANE RUPPERT
510(k) history	14 submissions · 14 cleared · 1987-1994