

K883933 INTRODUCING CATHETER

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

510k Database - www.510kdatabase.net
Device record: <https://www.510kdatabase.net/k883933/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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