FDA 510(k) Premarket Notification

K883916 ROTATING HEMOSTATIC VALVE

CLEARED

Oct 28, 1988
42 days to decision

K883916 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k883916/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Sep 16, 1988
Decision date	Oct 28, 1988
Days to decision	42 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/k883916/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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