FDA 510(k) Premarket Notification

K895214 HEMOSTASIS VALVE

CLEARED

Nov 14, 1989
88 days to decision

K895214 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k895214/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Aug 18, 1989
Decision date	Nov 14, 1989
Days to decision	88 days
Third-party review	No

APPLICANT

Company	**Applied Vascular Devices, Inc.**
Location	Santa Ana, CA, US
Contact	ROBERT P COOPER
510(k) history	21 submissions · 21 cleared · 1988-1991