

K895214 HEMOSTASIS VALVENov 14, 1989
88 days to decisionK895214 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895214/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026