

K874771 ANGIOMED INTRODUCING SETSJan 12, 1988
54 days to decisionK874771 · Product code: **DWB** · CardiovascularSource: <https://www.510kdatabase.net/k874771/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pump, Blood, Cardiopulmonary Bypass, Roller Type (DWB)
Date received	Nov 19, 1987
Decision date	Jan 12, 1988
Days to decision	54 days
Third-party review	No

APPLICANT

Company	Angiomed U.S., Inc.
Location	Anaheim, CA, US
Contact	MICHAEL MCCALLUM
510(k) history	24 submissions · 24 cleared · 1986-1988

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k874771/>, 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 29, 2026