

K854910 ANGIOFLOWJul 17, 1986
220 days to decisionK854910 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k854910/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Dec 9, 1985
Decision date	Jul 17, 1986
Days to decision	220 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k854910/>, 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