FDA 510(k) Premarket Notification

K854910 ANGIOFLOW

CLEARED

Jul 17, 1986
220 days to decision

K854910 · Product code: **DQO** · Cardiovascular
Source: 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