FDA 510(k) Premarket Notification

K854912 ANGIOMED GUIED WIRES

CLEARED

Apr 16, 1986
128 days to decision

K854912 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k854912/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Dec 9, 1985
Decision date	Apr 16, 1986
Days to decision	128 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