FDA 510(k) Premarket Notification

K870664 GUIDE WIRE

CLEARED

Mar 25, 1987
34 days to decision

K870664 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k870664/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 19, 1987
Decision date	Mar 25, 1987
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Datascope Corp.**
Location	Mchenry, IL, US
Contact	SCHNEIDER, PHD
510(k) history	136 submissions · 135 cleared · 1976-2019