FDA 510(k) Premarket Notification

K883579 CORONARY GUIDE WIRE

CLEARED

Nov 4, 1988
74 days to decision

K883579 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k883579/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Aug 22, 1988
Decision date	Nov 4, 1988
Days to decision	74 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