FDA 510(k) Premarket Notification

K030617 PT2 GUIDE WIRE AND ADDWIRE EXTENSION WIRE

CLEARED

May 21, 2003
84 days to decision

K030617 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k030617/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 26, 2003
Decision date	May 21, 2003
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Boston Scientific Scimed, Inc.**
Location	Plymouth, MN, US
Contact	ANNE V ROSSI
510(k) history	35 submissions · 26 cleared · 1994-2004