FDA 510(k) Premarket Notification

K821190 GUIDE WIRE

CLEARED

Aug 31, 1982
126 days to decision

K821190 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k821190/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Apr 27, 1982
Decision date	Aug 31, 1982
Days to decision	126 days
Third-party review	No

APPLICANT

Company	**Advanced Cardiovascular Systems, Inc.**
Location	Santa Clara, CA, US
510(k) history	103 submissions · 100 cleared · 1982-2002