FDA 510(k) Premarket Notification

K032129 RADIUS 018 COUGAR WIRE

CLEARED

Jul 17, 2003
7 days to decision

K032129 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k032129/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jul 10, 2003
Decision date	Jul 17, 2003
Days to decision	7 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Radius Medical Technologies, Inc.**
Location	Hopkonton, MA, US
Contact	DEBBIE IAMPIETRO
510(k) history	9 submissions · 9 cleared · 1997-2007