FDA 510(k) Premarket Notification

K042388 RADIANT MEDICAL ENDOCATHETER TEMPERATURE PROBE

CLEARED

Oct 28, 2004
56 days to decision

K042388 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k042388/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Sep 2, 2004
Decision date	Oct 28, 2004
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Radiant Medical**
Location	Redwood City, CA, US
Contact	ANDREW CLEELAND
510(k) history	6 submissions · 5 cleared · 2002-2007