FDA 510(k) Premarket Notification

K951603 CARDIMA, INC. FORERUNNER

CLEARED

Oct 5, 1995
181 days to decision

K951603 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k951603/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Apr 7, 1995
Decision date	Oct 5, 1995
Days to decision	181 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cardima, Inc.**
Location	Fremont, CA, US
Contact	J. D STEVENS
510(k) history	12 submissions · 12 cleared · 1993-2006