FDA 510(k) Premarket Notification

K120834 CORPATH 200 SYSTEM

CLEARED

Jul 19, 2012
122 days to decision

K120834 · Product code: **DXX** · Cardiovascular
Source: https://www.510kdatabase.net/k120834/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Catheter Control, Steerable (DXX)
Date received	Mar 19, 2012
Decision date	Jul 19, 2012
Days to decision	122 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Corindus, Inc.**
Location	Baltimore, MD, US
Contact	MONA ADVANI
510(k) history	9 submissions · 9 cleared · 2012-2022