FDA 510(k) Premarket Notification

K152999 CorPath 200 System

CLEARED

Mar 18, 2016
157 days to decision

K152999 · Product code: **DXX** · Cardiovascular
Source: https://www.510kdatabase.net/k152999/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Catheter Control, Steerable (DXX)
Date received	Oct 13, 2015
Decision date	Mar 18, 2016
Days to decision	157 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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