FDA 510(k) Premarket Notification

K141175 RENOVOCATH RC120

CLEARED

Oct 24, 2014
171 days to decision

K141175 · Product code: **MJN** · Cardiovascular
Source: https://www.510kdatabase.net/k141175/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular Occluding, Temporary (MJN)
Date received	May 6, 2014
Decision date	Oct 24, 2014
Days to decision	171 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Renovorx, Inc.**
Location	Sunnyvale, CA, US
Contact	RON S WARREN
510(k) history	4 submissions · 4 cleared · 2014-2021