FDA 510(k) Premarket Notification

CLEARED

K151880 Wingman Crossing Catheter

Aug 7, 2015
29 days to decision

K151880 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k151880/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Jul 9, 2015
Decision date	Aug 7, 2015
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Reflow Medical**
Location	San Diego, CA, US
Contact	REBECCA K PINE
510(k) history	10 submissions · 10 cleared · 2010-2021

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k151880/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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