FDA 510(k) Premarket Notification

K151896 Houdini Catheter

CLEARED

Dec 7, 2015
150 days to decision

K151896 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k151896/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jul 10, 2015
Decision date	Dec 7, 2015
Days to decision	150 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cruzar Medsystems, Inc.**
Location	Braintree, MA, US
Contact	Tom Kramer
510(k) history	2 submissions · 2 cleared · 2015-2022