FDA 510(k) Premarket Notification

K150894 VIA 21 Microcatheter

CLEARED

Aug 28, 2015
148 days to decision

K150894 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k150894/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Apr 2, 2015
Decision date	Aug 28, 2015
Days to decision	148 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Sequent Medical, Inc.**
Location	Aliso Viejo, CA, US
Contact	Bethany Barrett
510(k) history	4 submissions · 4 cleared · 2013-2017