FDA 510(k) Premarket Notification

K141782 HI-TORQUE VERSATURN GUIDE WIRE

CLEARED

Aug 7, 2014
36 days to decision

K141782 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k141782/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jul 2, 2014
Decision date	Aug 7, 2014
Days to decision	36 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Abbott Vascular**
Location	S,Mta Clara, CA, US
Contact	VIVEK THAKKAR
510(k) history	30 submissions · 30 cleared · 2009-2022