FDA 510(k) Premarket Notification

K091825 HI-TORQUE PROGRESS GUIDE WIRE FAMILY

CLEARED

Sep 25, 2009
98 days to decision

K091825 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k091825/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 19, 2009
Decision date	Sep 25, 2009
Days to decision	98 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Abbott Vascular, Inc.**
Location	Redwood, CA, US
Contact	MICHELE WALZ
510(k) history	20 submissions · 17 cleared · 2000-2014