FDA 510(k) Premarket Notification

K072460 HI-TORQUE BALANCE MIDDLEWEIGHT UNIVERSAL II GUIDE WIRE

CLEARED

Apr 11, 2008
224 days to decision

K072460 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k072460/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Aug 31, 2007
Decision date	Apr 11, 2008
Days to decision	224 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