FDA 510(k) Premarket Notification

K123067 PROGRESS GUIDE WIRE FAMILY, PILOT GUIDE WIRE FAMILY

CLEARED

Jan 29, 2013
120 days to decision

K123067 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k123067/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Oct 1, 2012
Decision date	Jan 29, 2013
Days to decision	120 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k123067/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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