

K141782 HI-TORQUE VERSATURN GUIDE WIREAug 7, 2014
36 days to decisionK141782 · Product code: **DQX** · CardiovascularSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k141782/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026