

K913328 .010 HI-TORQUE PLUS(TM) GUIDE WIREOct 23, 1991
90 days to decisionK913328 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k913328/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jul 25, 1991
Decision date	Oct 23, 1991
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Advanced Cardiovascular Systems, Inc.
Location	Santa Clara, CA, US
Contact	JANE E BEGGS
510(k) history	103 submissions · 100 cleared · 1982-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k913328/> 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 July 1, 2026