

**K914828 010 HI-TOR INTER/STAND GUIDWIRE PRESHAPE
TIP/MODIF**Jan 8, 1992
76 days to decisionK914828 · Product code: **DQX** · Cardiovascular
Source: <https://www.510kdatabase.net/k914828/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Oct 24, 1991
Decision date	Jan 8, 1992
Days to decision	76 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/k914828/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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