

K821190 GUIDE WIREAug 31, 1982
126 days to decisionK821190 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k821190/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Apr 27, 1982
Decision date	Aug 31, 1982
Days to decision	126 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k821190/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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