

K912293 REMOVE LABELING RESTRICT FROM GUIDE WIRESDec 5, 1991
196 days to decisionK912293 · Product code: **DQX** · Cardiovascular
Source: <https://www.510kdatabase.net/k912293/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	May 23, 1991
Decision date	Dec 5, 1991
Days to decision	196 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Target Therapeutics
Location	Los Angeles, CA, US
Contact	ALEX BALL
510(k) history	70 submissions · 70 cleared · 1985-1998

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