

K951603 CARDIMA, INC. FORERUNNEROct 5, 1995
181 days to decisionK951603 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k951603/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Apr 7, 1995
Decision date	Oct 5, 1995
Days to decision	181 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cardima, Inc.
Location	Fremont, CA, US
Contact	J. D STEVENS
510(k) history	12 submissions · 12 cleared · 1993-2006

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