FDA 510(k) Premarket Notification

K930961 CATHETER GUIDEWIRE (PERIPHERAL USE)

CLEARED

May 26, 1993
91 days to decision

K930961 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k930961/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 24, 1993
Decision date	May 26, 1993
Days to decision	91 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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