FDA 510(k) Premarket Notification

K912293 REMOVE LABELING RESTRICT FROM GUIDE WIRES

CLEARED

Dec 5, 1991
196 days to decision

K912293 · 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