FDA 510(k) Premarket Notification

K853999 GUIDE WIRE

CLEARED

Nov 5, 1985
36 days to decision

K853999 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k853999/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Sep 30, 1985
Decision date	Nov 5, 1985
Days to decision	36 days
Third-party review	No

APPLICANT

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