FDA 510(k) Premarket Notification

K853998 GUIDING CATHETER

CLEARED

Nov 6, 1985
37 days to decision

K853998 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k853998/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Sep 30, 1985
Decision date	Nov 6, 1985
Days to decision	37 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