FDA 510(k) Premarket Notification

K860842 ANGIOJECT

CLEARED

Apr 23, 1986
50 days to decision

K860842 · Product code: **DXT** · Cardiovascular
Source: https://www.510kdatabase.net/k860842/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Mar 4, 1986
Decision date	Apr 23, 1986
Days to decision	50 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