FDA 510(k) Premarket Notification

K860806 REGULATED PRESSURE INJECTOR

CLEARED

Mar 31, 1986
27 days to decision

K860806 · Product code: **DXF** · Cardiovascular
Source: https://www.510kdatabase.net/k860806/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Septostomy (DXF)
Date received	Mar 4, 1986
Decision date	Mar 31, 1986
Days to decision	27 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