

**K860806 REGULATED PRESSURE INJECTOR**Mar 31, 1986  
27 days to decisionK860806 · Product code: **DXF** · CardiovascularSource: <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	<b>Target Therapeutics</b>
Location	Los Angeles, CA, US
Contact	MARIE DANIELS
510(k) history	70 submissions · 70 cleared · 1985-1998

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860806/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 30, 2026