

**K894105 TARGET THERAPEUTICS TRACKER LF CATHETER**Sep 28, 1989  
111 days to decisionK894105 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k894105/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jun 9, 1989
Decision date	Sep 28, 1989
Days to decision	111 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/k894105/>; 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