

**K860842 ANGIOJECT**Apr 23, 1986  
50 days to decisionK860842 · Product code: **DXT** · CardiovascularSource: <https://www.510kdatabase.net/k860842/>**SUBMISSION DETAILS**

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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**

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Company	<b>Target Therapeutics</b>
Location	Los Angeles, CA, US
Contact	MARIE DANIELS
510(k) history	70 submissions · 70 cleared · 1985-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k860842/>; 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