

**K944875 ANGIOMAT 3000, 6000, CT INJECTOR**Jan 4, 1995  
93 days to decisionK944875 · Product code: **DXT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k944875/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Oct 3, 1994
Decision date	Jan 4, 1995
Days to decision	93 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Lieval-Flarsheim Co.</b>
Location	Oh, OH, US
Contact	M L FRIEDMAN
510(k) history	1 submissions · 1 cleared · 1995-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k944875/> 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 July 6, 2026