

**K033881 MEDRAD STELLANT CT INJECTOR SYSTEM WITH  
IMAGING SYSTEM INTERFACE MODULE**Mar 2, 2004  
78 days to decisionK033881 · Product code: **DXT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k033881/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Dec 15, 2003
Decision date	Mar 2, 2004
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medrad, Inc.</b>
Location	Mchenry, IL, US
Contact	TROY A JACK
Website	<a href="http://www.medrad.com/">http://www.medrad.com/</a>
510(k) history	105 submissions · 105 cleared · 1976-2014

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k033881/> 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 May 17, 2026