

**K052633 DUAL SHOT - CONTRAST DELIVERY SYSTEM**Oct 17, 2005  
21 days to decisionK052633 · Product code: **DXT** · CardiovascularSource: <https://www.510kdatabase.net/k052633/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Sep 26, 2005
Decision date	Oct 17, 2005
Days to decision	21 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nemoto Kyorindo Co., Ltd.</b>
Location	Tokyo, JP
Contact	TOSHIO KANETAKA
510(k) history	8 submissions · 8 cleared · 2005-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k052633/> 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 25, 2026