

**K914555 WORKSTATION FEMORAL INTRAVASCULAR  
RETRIEVAL (MODI)**Jan 3, 1992  
88 days to decisionK914555 · Product code: **DRE** · Cardiovascular  
Source: <https://www.510kdatabase.net/k914555/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	Oct 7, 1991
Decision date	Jan 3, 1992
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Med Institute, Inc.</b>
Location	West Lafayette, IN, US
Contact	NEAL E GEARNOT
510(k) history	26 submissions · 23 cleared · 1990-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k914555/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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