

**K882238 VESSEL DILATOR AND INTRODUCER SHEATH**Aug 25, 1988  
90 days to decisionK882238 · Product code: **DRE** · CardiovascularSource: <https://www.510kdatabase.net/k882238/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	May 27, 1988
Decision date	Aug 25, 1988
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>Ideal Medical, Inc.</b>
Location	Mchenry, IL, US
Contact	LORNA K LINVILLE
510(k) history	16 submissions · 14 cleared · 1981-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882238/>; 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 28, 2026