

**K881073 SHILEY FEMORAL CANNULAE INTRODUCER KIT**Jun 9, 1988  
87 days to decisionK881073 · Product code: **DRE** · CardiovascularSource: <https://www.510kdatabase.net/k881073/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, For Percutaneous Catheterization (DRE)
Date received	Mar 14, 1988
Decision date	Jun 9, 1988
Days to decision	87 days
Third-party review	No

**APPLICANT**

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Company	<b>Shiley, Inc.</b>
Location	Mchenry, IL, US
Contact	RADINE POBUDA
510(k) history	174 submissions · 174 cleared · 1976-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881073/>; 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 26, 2026