

**K884354 IMAGER FLUSH CATHETERS**Apr 21, 1989  
186 days to decisionK884354 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k884354/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Oct 17, 1988
Decision date	Apr 21, 1989
Days to decision	186 days
Third-party review	No

**APPLICANT**

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Company	<b>Medi-Tech, Inc.</b>
Location	Mchenry, IL, US
Contact	ALBERT P SEPRINSKI
510(k) history	36 submissions · 35 cleared · 1978-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k884354/>; 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 27, 2026