

**K981061 INTRAVASCULAR RETRIEVAL DEVICE**May 26, 1998  
64 days to decisionK981061 · Product code: **GCC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k981061/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Catheter (GCC)
Date received	Mar 23, 1998
Decision date	May 26, 1998
Days to decision	64 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 FEARNOT
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/k981061/> 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 19, 2026