

**K771986 CATHETER, MEDIASTINAL&PERICARDIAL**Oct 28, 1977  
9 days to decisionK771986 · Product code: **FHF** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k771986/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Evacuator, Fluid (FHF)
Date received	Oct 19, 1977
Decision date	Oct 28, 1977
Days to decision	9 days
Third-party review	No

**APPLICANT**

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Company	<b>The Minnesota Back Brace Co.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1977-1977

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k771986/> 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 July 4, 2026