

**K880570 DENVER PLEURO-PERITONEAL SHUNT**Jun 20, 1988  
131 days to decisionK880570 · Product code: **KPM** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k880570/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shunt, Peritoneal (KPM)
Date received	Feb 10, 1988
Decision date	Jun 20, 1988
Days to decision	131 days
Third-party review	No

**APPLICANT**

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Company	<b>Codman &amp; Shurtleff, Inc.</b>
Location	Mchenry, IL, US
Contact	G CLOUTIER
510(k) history	152 submissions · 151 cleared · 1976-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880570/>, 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 June 15, 2026