

**K880401 DENVER PERITONEO-VEIN SHUNT**May 31, 1988  
123 days to decisionK880401 · Product code: **KPM** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k880401/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shunt, Peritoneal (KPM)
Date received	Jan 29, 1988
Decision date	May 31, 1988
Days to decision	123 days
Third-party review	No

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880401/>, 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