

**K811183 DENVER PERITONEO-VEIN SHUNT**Jul 1, 1981  
63 days to decisionK811183 · Product code: **KPM** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k811183/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Shunt, Peritoneal (KPM)
Date received	Apr 29, 1981
Decision date	Jul 1, 1981
Days to decision	63 days
Third-party review	No

**APPLICANT**

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Company	<b>Denver Biomedicals, Inc.</b>
Location	Englewood, CO, US
Contact	JOHN B NEWKIRK
510(k) history	10 submissions · 10 cleared · 1981-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k811183/>; 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 25, 2026