

**K011862 DENVER ASCITES SHUNT AND PERCUTANEOUS  
ACCESS KIT WITH ASCITIES SHUNT**Jul 12, 2001  
28 days to decisionK011862 · Product code: **KPM** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k011862/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Shunt, Peritoneal (KPM)
Date received	Jun 14, 2001
Decision date	Jul 12, 2001
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Denver Biomedical, Inc.</b>
Location	Golden, CO, US
Contact	BONNIE B VIVIAN
510(k) history	5 submissions · 5 cleared · 2001-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k011862/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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