

**K822547 CAPD BELT**Sep 9, 1982  
16 days to decisionK822547 · Product code: **FKX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k822547/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Aug 24, 1982
Decision date	Sep 9, 1982
Days to decision	16 days
Third-party review	No

**APPLICANT**

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Company	<b>American Medical Products, Inc.</b>
Location	Mchenry, IL, US
510(k) history	30 submissions · 30 cleared · 1978-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k822547/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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