

K823827 AMP 80/2 OUTFLOW MONITOR MODIFICATIONMar 17, 1983
87 days to decisionK823827 · Product code: **FKX** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k823827/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Dec 20, 1982
Decision date	Mar 17, 1983
Days to decision	87 days
Third-party review	No

APPLICANT

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

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