

**K830881 AMPERCIDE DISINFECTANT ACCESSORY TO -**Jun 2, 1983  
76 days to decisionK830881 · Product code: **FKX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k830881/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Mar 18, 1983
Decision date	Jun 2, 1983
Days to decision	76 days
Third-party review	No

**APPLICANT**

---

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

---

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

Device record: <https://www.510kdatabase.net/k830881/> 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 29, 2026