FDA 510(k) Premarket Notification

# K830881 AMPERCIDE DISINFECTANT ACCESSORY TO -

**CLEARED**

Jun 2, 1983
76 days to decision

K830881 · Product code: **FKX** · Gastroenterology & Urology
Source: 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 | **American Medical Products, Inc.** |
| Location | Mchenry, IL, US |
| 510(k) history | 30 submissions · 30 cleared · 1978-1992 |