FDA 510(k) Premarket Notification

K823827 AMP 80/2 OUTFLOW MONITOR MODIFICATION

CLEARED

Mar 17, 1983
87 days to decision

K823827 · 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