FDA 510(k) Premarket Notification

K822547 CAPD BELT

CLEARED

Sep 9, 1982
16 days to decision

K822547 · Product code: **FKX** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k822547/

SUBMISSION DETAILS

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

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