FDA 510(k) Premarket Notification

K780957 SET, ADMINISTRATION, MANUAL, DIALYSIS

CLEARED

Jul 6, 1978
24 days to decision

K780957 · Product code: **KDJ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k780957/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, For Peritoneal Dialysis, Disposable (KDJ)
Date received	Jun 12, 1978
Decision date	Jul 6, 1978
Days to decision	24 days
Third-party review	No

APPLICANT

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