FDA 510(k) Premarket Notification

K812615 TM-CAPD

CLEARED

Jan 5, 1982
112 days to decision

K812615 · Product code: **KPP** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k812615/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Peritoneal Dialysate Filter (KPP)
Date received	Sep 15, 1981
Decision date	Jan 5, 1982
Days to decision	112 days
Third-party review	No

APPLICANT

Company	**Tri-Med, Inc.**
Location	Mchenry, IL, US
510(k) history	29 submissions · 29 cleared · 1977-2004