FDA 510(k) Premarket Notification

K812717 NON-STERILE SHADOW-STRIPE CATHETER

CLEARED

Oct 19, 1981
24 days to decision

K812717 · Product code: **FKO** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k812717/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Peritoneal Dialysis, Single Use (FKO)
Date received	Sep 25, 1981
Decision date	Oct 19, 1981
Days to decision	24 days
Third-party review	No

APPLICANT

Company	**Quinton, Inc.**
Location	Mchenry, IL, US
510(k) history	164 submissions · 160 cleared · 1976-2003