FDA 510(k) Premarket Notification

K811266 QUINTON CATHETER FINGER GRIP

CLEARED

Jun 18, 1981
37 days to decision

K811266 · Product code: **FKO** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k811266/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Peritoneal Dialysis, Single Use (FKO)
Date received	May 12, 1981
Decision date	Jun 18, 1981
Days to decision	37 days
Third-party review	No

APPLICANT

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