FDA 510(k) Premarket Notification

K771420 BETA-CAP

CLEARED

Nov 28, 1977
119 days to decision

K771420 · Product code: **FJS** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k771420/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Peritoneal, Long-term Indwelling (FJS)
Date received	Aug 1, 1977
Decision date	Nov 28, 1977
Days to decision	119 days
Third-party review	No

APPLICANT

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