FDA 510(k) Premarket Notification

K801679 BETA-CAP II

CLEARED

Oct 3, 1980
73 days to decision

K801679 · Product code: **FJS** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k801679/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Peritoneal, Long-term Indwelling (FJS)
Date received	Jul 22, 1980
Decision date	Oct 3, 1980
Days to decision	73 days
Third-party review	No

APPLICANT

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