

K771420 BETA-CAPNov 28, 1977
119 days to decisionK771420 · 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

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Device record: <https://www.510kdatabase.net/k771420/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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