

K801679 BETA-CAP IIOct 3, 1980
73 days to decisionK801679 · 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

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Device record: <https://www.510kdatabase.net/k801679/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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