

K811266 QUINTON CATHETER FINGER GRIPJun 18, 1981
37 days to decisionK811266 · Product code: **FKO** · Gastroenterology & UrologySource: <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

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Device record: <https://www.510kdatabase.net/k811266/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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