

**K812717 NON-STERILE SHADOW-STRIPE CATHETER**Oct 19, 1981  
24 days to decisionK812717 · Product code: **FKO** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k812717/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Peritoneal Dialysis, Single Use (FKO)
Date received	Sep 25, 1981
Decision date	Oct 19, 1981
Days to decision	24 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k812717/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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