

K830208 BETA-CAP D ADAPTERMar 7, 1983
45 days to decisionK830208 · Product code: **FKX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k830208/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Jan 21, 1983
Decision date	Mar 7, 1983
Days to decision	45 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/k830208/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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