

K823857 IMPRA PERITONEAL CATHETERJan 26, 1983
35 days to decisionK823857 · Product code: **FKX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k823857/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Dec 22, 1982
Decision date	Jan 26, 1983
Days to decision	35 days
Third-party review	No

APPLICANT

Company	Impra, Inc.
Location	Mchenry, IL, US
510(k) history	29 submissions · 24 cleared · 1979-2001

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Device record: <https://www.510kdatabase.net/k823857/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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