

**K972579 HIGH DOSE DISCONNECT CAP AND MINICAP WITH
POVIDONE-IODINE, HIGH DOSE DISCONNECT CAP WITH
POVIDONE-IODINE**

Oct 1, 1997
83 days to decision

K972579 · Product code: **KDJ** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k972579/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, For Peritoneal Dialysis, Disposable (KDJ)
Date received	Jul 10, 1997
Decision date	Oct 1, 1997
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Baxter Healthcare Corp
Location	Mchenry, IL, US
Contact	DAVID C ROSS
510(k) history	505 submissions · 496 cleared · 1977-2019

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Device record: <https://www.510kdatabase.net/k972579/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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