

**K780957 SET, ADMINISTRATION, MANUAL, DIALYSIS**Jul 6, 1978  
24 days to decisionK780957 · Product code: **KDJ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k780957/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, For Peritoneal Dialysis, Disposable (KDJ)
Date received	Jun 12, 1978
Decision date	Jul 6, 1978
Days to decision	24 days
Third-party review	No

**APPLICANT**

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Company	<b>American Medical Products, Inc.</b>
Location	Mchenry, IL, US
510(k) history	30 submissions · 30 cleared · 1978-1992

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

Device record: <https://www.510kdatabase.net/k780957/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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