

**K812130 SPIKE GUARD**Sep 9, 1981  
44 days to decisionK812130 · Product code: **KDJ** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k812130/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, For Peritoneal Dialysis, Disposable (KDJ)
Date received	Jul 27, 1981
Decision date	Sep 9, 1981
Days to decision	44 days
Third-party review	No

**APPLICANT**

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Company	<b>Clinipad Corp.</b>
Location	Mchenry, IL, US
510(k) history	16 submissions · 12 cleared · 1980-1994

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

Device record: <https://www.510kdatabase.net/k812130/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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