

**K823330 DIALATORS PC-203 & 202**Jan 26, 1983  
79 days to decisionK823330 · Product code: **FKX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k823330/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Nov 8, 1982
Decision date	Jan 26, 1983
Days to decision	79 days
Third-party review	No

**APPLICANT**

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Company	<b>Medigroup</b>
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1983-1983

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k823330/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 30, 2026