

**K831766 MICROSTAR VCI**Oct 14, 1983  
135 days to decisionK831766 · Product code: **FKX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k831766/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Jun 1, 1983
Decision date	Oct 14, 1983
Days to decision	135 days
Third-party review	No

**APPLICANT**

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Company	<b>Medionics International , Ltd.</b>
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1982-1992

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

Device record: <https://www.510kdatabase.net/k831766/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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