

**K800186 LOGO-NU-WAY**Feb 26, 1980  
28 days to decisionK800186 · Product code: **KNX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k800186/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Jan 29, 1980
Decision date	Feb 26, 1980
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Gary Giancarlo</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980

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

Device record: <https://www.510kdatabase.net/k800186/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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