

**K810645 SELECT-A-FLOW COLLECTOR**May 5, 1981  
56 days to decisionK810645 · Product code: **KNX** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k810645/>**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	Mar 10, 1981
Decision date	May 5, 1981
Days to decision	56 days
Third-party review	No

**APPLICANT**

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Company	<b>Wenmar Intl. Corp.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1981

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

Device record: <https://www.510kdatabase.net/k810645/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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