

**K781422 KWIK-FLO URINAL SYSTEM**Sep 7, 1978  
21 days to decisionK781422 · Product code: **KNX** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k781422/>**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	Aug 17, 1978
Decision date	Sep 7, 1978
Days to decision	21 days
Third-party review	No

**APPLICANT**

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Company	<b>The Grace Mfg. Co.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978

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

Device record: <https://www.510kdatabase.net/k781422/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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