

**K820565 MALE EXTERNAL CATHETER**Apr 1, 1982  
30 days to decisionK820565 · Product code: **KNX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k820565/>**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 2, 1982
Decision date	Apr 1, 1982
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>Urocare Products, Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k820565/> 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 July 3, 2026