

**K863739 BESCURE**Oct 24, 1986  
31 days to decisionK863739 · Product code: **KNX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k863739/>**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	Sep 23, 1986
Decision date	Oct 24, 1986
Days to decision	31 days
Third-party review	No

**APPLICANT**

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Company	<b>U-T, Inc.</b>
Location	Chagrin Falls, OH, US
Contact	RICHARD G BAERMAN
510(k) history	1 submissions · 1 cleared · 1986-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k863739/>; 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