

**K852051 EXTERNAL MALE CATHETER**Aug 22, 1985  
105 days to decisionK852051 · Product code: **KNX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k852051/>**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	May 9, 1985
Decision date	Aug 22, 1985
Days to decision	105 days
Third-party review	No

**APPLICANT**

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Company	<b>Buffalo Medical Specialties Mfg., Inc.</b>
Location	Walker, MI, US
Contact	HANSEN
510(k) history	3 submissions · 3 cleared · 1983-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k852051/>, 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 June 29, 2026