

**K874309 URETHAL CATHETER**Dec 7, 1987  
48 days to decisionK874309 · Product code: **KNX** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k874309/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Oct 20, 1987
Decision date	Dec 7, 1987
Days to decision	48 days
Third-party review	No

**APPLICANT**

---

Company	<b>Medovations, Inc.</b>
Location	Mchenry, IL, US
Contact	LARRY R PIERCE
510(k) history	31 submissions · 31 cleared · 1983-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874309/>; 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 May 14, 2026