

**K790279 CATHETER, MMG LATEX EXTERNAL**Mar 6, 1979  
21 days to decisionK790279 · Product code: **KNX** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k790279/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Collector, Urine, (and Accessories) For Indwelling Catheter (KNX)
Date received	Feb 13, 1979
Decision date	Mar 6, 1979
Days to decision	21 days
Third-party review	No

**APPLICANT**

---

Company	<b>Medical Marketing Group</b>
Location	Decatur, GA, US
Contact	TIMOTHY GOLDEN
510(k) history	11 submissions · 10 cleared · 1979-1991

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k790279/>; 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 27, 2026