

**K953375 NG SECURE**Sep 8, 1995  
52 days to decisionK953375 · Product code: **KNT** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k953375/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Jul 18, 1995
Decision date	Sep 8, 1995
Days to decision	52 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>M.C. Johnson Co., Inc.</b>
Location	Walker, MI, US
Contact	PHILIP BAKER
510(k) history	3 submissions · 3 cleared · 1983-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k953375/> 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 21, 2026