

K875171 J-CATH(TM)Apr 6, 1988
111 days to decisionK875171 · Product code: **KNT** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k875171/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Dec 17, 1987
Decision date	Apr 6, 1988
Days to decision	111 days
Third-party review	No

APPLICANT

Company	Hdc Corp.
Location	Walker, MI, US
Contact	RONALD L COLEMAN
510(k) history	30 submissions · 29 cleared · 1983-2007

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k875171/>, Data sourced from FDA 510(k) public records (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. - Generated May 25, 2026