

**K062240 CAREVENT**Oct 16, 2006  
75 days to decisionK062240 · Product code: **KNY** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k062240/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessories, Catheter, G-u (KNY)
Date received	Aug 2, 2006
Decision date	Oct 16, 2006
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Chief Medical, LLC</b>
Location	Teton Village, WY, US
Contact	SCOTT HORN
510(k) history	1 submissions · 1 cleared · 2006-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062240/> 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