

**K133599 CORGRIP**Mar 26, 2014  
121 days to decisionK133599 · Product code: **KNT** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k133599/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Nov 25, 2013
Decision date	Mar 26, 2014
Days to decision	121 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Corpak Medsystems</b>
Location	Buffalo Grove, IL, US
Contact	STEPHANIE WASIELEWSKI
510(k) history	15 submissions · 15 cleared · 1982-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133599/> 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 5, 2026