

**K120182 NEOMED GASTROINTESTINAL TUBE AND ACCESSORIES**Apr 19, 2012  
90 days to decisionK120182 · Product code: **KNT** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k120182/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Jan 20, 2012
Decision date	Apr 19, 2012
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Neomed, Inc.</b>
Location	Swanee, GA, US
Contact	PENNY NORTHCUTT
510(k) history	13 submissions · 13 cleared · 2007-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120182/> 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](http://accessdata.fda.gov). - Generated June 20, 2026