

**K200142 SpeediCath Soft**Jul 30, 2020  
190 days to decisionK200142 · Product code: **GBM** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k200142/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urethral (GBM)
Date received	Jan 22, 2020
Decision date	Jul 30, 2020
Days to decision	190 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Coloplast</b>
Location	Plymouth, MN, US
Contact	Delaney McDougal
Website	<a href="http://www.coloplast.com/">http://www.coloplast.com/</a>
510(k) history	15 submissions · 14 cleared · 2018-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k200142/> 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 25, 2026