

**K152268 VAPRO PLUS POCKET Intermittent Catheter**Sep 9, 2015  
29 days to decisionK152268 · Product code: **GBM** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k152268/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Urethral (GBM)
Date received	Aug 11, 2015
Decision date	Sep 9, 2015
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hollister Incorporated</b>
Location	Libertyville, IL, US
Contact	JEANNE LEE
Website	<a href="http://www.hollister.com/">http://www.hollister.com/</a>
510(k) history	14 submissions · 14 cleared · 2011-2025

Hollister Incorporated specializes in ostomy, continence, and critical care products with a manufacturing facility in Libertyville, US. The company serves patients and healthcare professionals globally across multiple therapeutic areas. Hollister has received FDA 510(k) clearances from total submissions since 2011. The company's regulatory focus centers on Gastroenterology & Urology devices, with the most recent clearance in 2025. This demonstrates sustained innovation and active market engagement in continence care solutions. Recent cleared devices include intermittent c...

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