

**K090960 VAPRO INTERMITTENT CATHETER, MODEL 72062,  
72082,72102, 72122, 72142, 72064, 72084, 72104, 72124,72144,  
72164, 72184,**Aug 20, 2009  
136 days to decisionK090960 · Product code: **GBM** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k090960/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urethral (GBM)
Date received	Apr 6, 2009
Decision date	Aug 20, 2009
Days to decision	136 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hollister, Inc.</b>
Location	Mchenry, IL, US
Contact	CHRIS STUKEL
510(k) history	85 submissions · 78 cleared · 1977-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090960/> 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 22, 2026