

**K984084 BARDEX LUBRI-SIL FOLEY CATHETER**Feb 1, 1999  
77 days to decisionK984084 · Product code: **EZL** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k984084/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Retention Type, Balloon (EZL)
Date received	Nov 16, 1998
Decision date	Feb 1, 1999
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bard Medical Div.</b>
Location	Covington, GA, US
Contact	DONNA J WILSON
510(k) history	3 submissions · 3 cleared · 1997-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k984084/> 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 July 1, 2026