

**K061766 EXPRESSWAY INTERMITTENT CATHETER**Mar 9, 2007  
260 days to decisionK061766 · Product code: **EZD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k061766/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Straight (EZD)
Date received	Jun 22, 2006
Decision date	Mar 9, 2007
Days to decision	260 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Percutaneous Systems, Incorporated</b>
Location	Mountain View, CA, US
Contact	THOMAS LAWSON
510(k) history	5 submissions · 5 cleared · 2005-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k061766/> 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 June 22, 2026