

**K102887 ACCORDION STONE MANAGEMENT DEVICE MODEL  
AC281205 AND AC3814510**May 16, 2011  
228 days to decisionK102887 · Product code: **FFL** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k102887/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dislodger, Stone, Basket, Ureteral, Metal (FFL)
Date received	Sep 30, 2010
Decision date	May 16, 2011
Days to decision	228 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Percutaneous Systems, Inc.</b>
Location	Washington, Dc, DC, US
Contact	THOMAS LAWSON, PHD
510(k) history	6 submissions · 6 cleared · 2004-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k102887/> 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 19, 2026