

**K090485 COAXIAL ACCORDION ERCP DEVICE, MODELS  
COAC19001, COAC20005, COAC22003**Jul 22, 2009  
147 days to decisionK090485 · Product code: LQR · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k090485/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dislodger, Stone, Biliary (LQR)
Date received	Feb 25, 2009
Decision date	Jul 22, 2009
Days to decision	147 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/k090485/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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