

**K082803 COAXIAL ACCORDION STONE MANAGEMENT
DEVICE, MODEL: COAC12005 - COA20015**Nov 13, 2008
50 days to decisionK082803 · Product code: FFL · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k082803/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dislodger, Stone, Basket, Ureteral, Metal (FFL)
Date received	Sep 24, 2008
Decision date	Nov 13, 2008
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082803/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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