

K080360 ENDOLOGIX VISIFLEX DUAL LUMEN CATHETERApr 8, 2008
57 days to decisionK080360 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k080360/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Feb 11, 2008
Decision date	Apr 8, 2008
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Endologix, Inc.
Location	Irvine, CA, US
Contact	ANNALIZA VICTORIA
Website	http://www.endologix.com/
510(k) history	5 submissions · 5 cleared · 1999-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k080360/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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