

**K092298 MODIFICATION TO MODIFIED HD GUIDE CATHETER**

Nov 5, 2009  
 99 days to decision

K092298 · Product code: **DQY** · Cardiovascular  
 Source: <https://www.510kdatabase.net/k092298/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Jul 29, 2009
Decision date	Nov 5, 2009
Days to decision	99 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Concentric Medical, Inc.</b>
Location	Moutian View, CA, US
Contact	KIRSTEN VALLEY
510(k) history	45 submissions · 44 cleared · 2001-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k092298/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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