

**K003085 CONCENTRIC GUIDE CATHETER, MODELS 90018,  
90020**Mar 7, 2001  
155 days to decisionK003085 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k003085/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Oct 3, 2000
Decision date	Mar 7, 2001
Days to decision	155 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	LINDA BRADLEY
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/k003085/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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