

K003880 CONCENTROC HD GUIDE CATHETER, MODELS 90017 (4F), 90019 (5F), 90021 (6F), 90022 (7F), 90023 (8F), 90024 (9F)May 16, 2001
152 days to decisionK003880 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k003880/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Catheter, Percutaneous (DQY)
Date received	Dec 15, 2000
Decision date	May 16, 2001
Days to decision	152 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Concentric Medical, Inc.
Location	Moutian View, CA, US
Contact	SAM LAZZARA
510(k) history	45 submissions · 44 cleared · 2001-2018

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k003880/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026