

K151638 Phenom 17 Catheter, Phenom 21 Catheter, Phenom 27 Catheter, Phenom Plus CatheterNov 13, 2015
149 days to decisionK151638 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k151638/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jun 17, 2015
Decision date	Nov 13, 2015
Days to decision	149 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cathera, Inc.
Location	Mountain View, CA, US
Contact	Victor Ham
510(k) history	2 submissions · 2 cleared · 2015-2018

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k151638/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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