

K150201 Quick-Cross Extreme Support Catheter, Quick-Cross Select Support CatheterFeb 27, 2015
29 days to decisionK150201 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k150201/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Jan 29, 2015
Decision date	Feb 27, 2015
Days to decision	29 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Spectranetics, Inc.
Location	Colorado Springs, CO, US
Contact	Christine Godleski
510(k) history	17 submissions · 16 cleared · 2014-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k150201/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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