

**K013581 CORDIS AVIATOR PERIPHERAL DILATATION
CATHETER**Nov 28, 2001
30 days to decisionK013581 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k013581/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Oct 29, 2001
Decision date	Nov 28, 2001
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cordis, A Johnson & Johnson Co.
Location	Warren, NJ, US
Contact	CHUCK RYAN
510(k) history	4 submissions · 2 cleared · 1999-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k013581/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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