

**K121442 POWERFLEX PRO PERCUTANEOUS TRANSLUMINAL ANGIOPLASTY CATHETER**Jun 14, 2012  
30 days to decisionK121442 · Product code: LIT · Cardiovascular  
Source: <https://www.510kdatabase.net/k121442/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Angioplasty, Peripheral, Transluminal (LIT)
Date received	May 15, 2012
Decision date	Jun 14, 2012
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cordis Corporation</b>
Location	Warren, NJ, US
Contact	DONNA MARSHALL
510(k) history	13 submissions · 12 cleared · 2004-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121442/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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