K121442 POWERFLEX PRO PERCUTANEOUS TRANSLUMINAL ANGIOPLASTY CATHETER

CLEARED

Jun 14, 2012
30 days to decision

K121442 · Product code: **LIT** · Cardiovascular
Source: https://www.510kdatabase.net/k121442/

SUBMISSION DETAILS

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

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