

**K032737 CORDIS POWERFLEX EXTREME PTA BALLOON  
CATHETER, CORDIS POWERFLEX PLUS PTA BALLOON  
CATHETER, CORDIS POWERFLEX P3 PTA**Oct 2, 2003  
28 days to decisionK032737 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k032737/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Sep 4, 2003
Decision date	Oct 2, 2003
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cordis Europa, N.V.</b>
Location	Warren, NJ, US
Contact	KAREN WILK
510(k) history	7 submissions · 5 cleared · 2001-2007

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