K032737 CORDIS POWERFLEX EXTREME PTA BALLOON CATHETER, CORDIS POWERFLEX PLUS PTA BALLOON CATHETER, CORDIS POWERFLEX P3 PTA

CLEARED

Oct 2, 2003
28 days to decision

K032737 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k032737/

SUBMISSION DETAILS

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

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