FDA 510(k) Premarket Notification

K013581 CORDIS AVIATOR PERIPHERAL DILATATION CATHETER

CLEARED

Nov 28, 2001
30 days to decision

K013581 · 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