FDA 510(k) Premarket Notification

K233637 Cordis BRITECROSS Support Catheter

CLEARED

Jun 28, 2024
228 days to decision

K233637 · Product code: **DQY** · Cardiovascular
Source: https://www.510kdatabase.net/k233637/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Nov 13, 2023
Decision date	Jun 28, 2024
Days to decision	228 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Cordis US Corp**
Location	Miami Lakes, FL, US
Contact	Linda Ruedy
510(k) history	4 submissions · 4 cleared · 2022-2024