

K233637 Cordis BRITECROSS Support CatheterJun 28, 2024
228 days to decisionK233637 · Product code: **DQY** · CardiovascularSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k233637/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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