

K210685 AcQCross Qx Integrated Transseptal Dilator/NeedleApr 6, 2021
29 days to decisionK210685 · Product code: **DYB** · Cardiovascular
Source: <https://www.510kdatabase.net/k210685/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Mar 8, 2021
Decision date	Apr 6, 2021
Days to decision	29 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Acutus Medical, Inc.
Location	Carlsbad, CA, US
Contact	Sarah Clay
510(k) history	24 submissions · 24 cleared · 2017-2023

REGULATORY CONSULTANT

Consulting firm	Regulatory Technology Services, LLC
Contact	Prithul Bom

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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