

**K243193 Cross Wise™ Multi-Use RF Adapter Cable**Nov 26, 2024  
57 days to decisionK243193 · Product code: **DXF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k243193/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Septostomy (DXF)
Date received	Sep 30, 2024
Decision date	Nov 26, 2024
Days to decision	57 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Circa Scientific, Inc.</b>
Location	Englewood, CO, US
Contact	Jennifer Willner
510(k) history	6 submissions · 6 cleared · 2023-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>JW Regulatory Consulting, LLC</b>
Contact	Jennifer Willner

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k243193/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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