

**K253671 Dual Stage Venous Cannulae**Jan 30, 2026  
70 days to decisionK253671 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k253671/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Nov 21, 2025
Decision date	Jan 30, 2026
Days to decision	70 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sorin Group Italia S.R.L.</b>
Location	Mirandola, IT
Contact	Luigi Vecchi
510(k) history	61 submissions · 61 cleared · 1995-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k253671/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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