

**K832124 CPC, COUBLE LUMEN CARDIOPLEGIA-AIR ASPI**Aug 12, 1983  
43 days to decisionK832124 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k832124/>**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	Jun 30, 1983
Decision date	Aug 12, 1983
Days to decision	43 days
Third-party review	No

**APPLICANT**

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Company	<b>Med-West, Inc.</b>
Location	Mchenry, IL, US
510(k) history	11 submissions · 8 cleared · 1983-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k832124/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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