

**K792202 TROCAR CATHETER/THORACIC CATHETER**Nov 20, 1979  
18 days to decisionK792202 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k792202/>**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 2, 1979
Decision date	Nov 20, 1979
Days to decision	18 days
Third-party review	No

**APPLICANT**

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Company	<b>Portex, Inc.</b>
Location	Walker, MI, US
510(k) history	20 submissions · 20 cleared · 1977-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k792202/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 25, 2026