

**K770068 CANNULA, HUNTER-SESSIONS, SELF-RETAIN.**Mar 1, 1977  
48 days to decisionK770068 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k770068/>**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	Jan 12, 1977
Decision date	Mar 1, 1977
Days to decision	48 days
Third-party review	No

**APPLICANT**

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Company	<b>Barlow Mfg., Corp.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1977-1977

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k770068/> 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 July 10, 2026