

**K920178 CARDIOTOMY RESERVOIR W/20 MICRON FILITER**May 13, 1993  
485 days to decisionK920178 · Product code: **DTN** · CardiovascularSource: <https://www.510kdatabase.net/k920178/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reservoir, Blood, Cardiopulmonary Bypass (DTN)
Date received	Jan 14, 1992
Decision date	May 13, 1993
Days to decision	485 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lifestream Int&amp;apos;L, Inc.</b>
Location	Sugar Land, TX, US
Contact	RUSSELL JONES
510(k) history	23 submissions · 23 cleared · 1984-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920178/> 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 22, 2026