

**K842829 HIGH PRESSURE, 1 & 3 WAY STERILE DISPOS**Oct 18, 1984  
90 days to decisionK842829 · Product code: **DTL** · Cardiovascular  
Source: <https://www.510kdatabase.net/k842829/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Adaptor, Stopcock, Manifold, Fitting, Cardiopulmonary Bypass (DTL)
Date received	Jul 20, 1984
Decision date	Oct 18, 1984
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>North American Instrument Corp.</b>
Location	Walker, MI, US
Contact	PHILLIP H MORSE
510(k) history	28 submissions · 28 cleared · 1983-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k842829/>; 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 21, 2026