

**K883696 FLO-RESTER**Nov 30, 1988  
92 days to decisionK883696 · Product code: **DXC** · Cardiovascular  
Source: <https://www.510kdatabase.net/k883696/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Vascular (DXC)
Date received	Aug 30, 1988
Decision date	Nov 30, 1988
Days to decision	92 days
Third-party review	No

**APPLICANT**

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Company	<b>Bio-Vascular, Inc.</b>
Location	St. Paul, MN, US
Contact	SHERRY A KRATTLEY
510(k) history	26 submissions · 25 cleared · 1986-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883696/>; 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 20, 2026