

**K110193 FEMOSTOP PLYS, FEMOSTOP HD, FEMOSTOP  
COMPRESSION ARCH**Feb 24, 2011  
31 days to decisionK110193 · Product code: **DXC** · Cardiovascular  
Source: <https://www.510kdatabase.net/k110193/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Clamp, Vascular (DXC)
Date received	Jan 24, 2011
Decision date	Feb 24, 2011
Days to decision	31 days
Third-party review	No
Summary / Statement	Summary
Other names	FEMOSTOP II PLUS SET; FEMOSTOP II PLUS ARCH, BILATERAL ADAPTER; F

**APPLICANT**

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Company	<b>St. Jude Medical Systems AB</b>
Location	Uppsala, SE
Contact	KATRIN SVENSSON
510(k) history	5 submissions · 5 cleared · 2011-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k110193/> 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 June 5, 2026