

**K080206 MODIFICATION TO FEMOSTOP FEMORAL
COMPRESSION SYSTEM**Feb 26, 2008
29 days to decisionK080206 · Product code: **DXC** · Cardiovascular
Source: <https://www.510kdatabase.net/k080206/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Clamp, Vascular (DXC)
Date received	Jan 28, 2008
Decision date	Feb 26, 2008
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Radi Medical Systems AB
Location	Uppsala, SE
Contact	HELENE EKSTRAND
510(k) history	24 submissions · 24 cleared · 1990-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k080206/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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