

**K011832 VANGUARD REPROCESSED FEMORAL  
COMPRESSION DEVICE**Dec 21, 2001  
192 days to decisionK011832 · Product code: **NMF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k011832/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Vascular, Reprocessed (NMF)
Date received	Jun 12, 2001
Decision date	Dec 21, 2001
Days to decision	192 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vanguard Medical Concepts, Inc.</b>
Location	Plant City, FL, US
Contact	MIKE SAMMON
510(k) history	33 submissions · 33 cleared · 1991-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k011832/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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