

**K102582 ARTERIOFLOW MODEL 7500 SEQUENTIAL
COMPRESSION DEVICE**Oct 7, 2010
29 days to decisionK102582 · Product code: **JOW** · Cardiovascular
Source: <https://www.510kdatabase.net/k102582/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sleeve, Limb, Compressible (JOW)
Date received	Sep 8, 2010
Decision date	Oct 7, 2010
Days to decision	29 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Devon Medical, Inc.
Location	King Of Prussia, PA, US
Contact	RUTH WU
510(k) history	11 submissions · 11 cleared · 2009-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k102582/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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