FDA 510(k) Premarket Notification

K102582 ARTERIOFLOW MODEL 7500 SEQUENTIAL COMPRESSION DEVICE

CLEARED

Oct 7, 2010
29 days to decision

K102582 · 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