

**K203803 VASOBand Vascular Compression Device**Jan 26, 2021  
29 days to decisionK203803 · Product code: **DXC** · Cardiovascular  
Source: <https://www.510kdatabase.net/k203803/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Vascular (DXC)
Date received	Dec 28, 2020
Decision date	Jan 26, 2021
Days to decision	29 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vasoinnovations, Inc.</b>
Location	South Pasadena, CA, US
Contact	Rajendra Gurudas Sardesai
510(k) history	2 submissions · 2 cleared · 2019-2021

**REGULATORY CONSULTANT**

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Consulting firm	<b>Regulatory Technology Services, LLC</b>
Contact	Prithul Bom

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k203803/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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