

**K183328 Pulse Rx 2.0, Pulse Pro Rx 2.0**Dec 21, 2018  
21 days to decisionK183328 · Product code: **JOW** · CardiovascularSource: <https://www.510kdatabase.net/k183328/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sleeve, Limb, Compressible (JOW)
Date received	Nov 30, 2018
Decision date	Dec 21, 2018
Days to decision	21 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>NormaTec Industries, LP</b>
Location	Washington, DC, US
Contact	Albert G. Sanford
510(k) history	9 submissions · 9 cleared · 2012-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k183328/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 26, 2026