

**K090308 RESTEP DVY SYSTEM, MODEL RSP-101**May 1, 2009  
84 days to decisionK090308 · Product code: **JOW** · Cardiovascular  
Source: <https://www.510kdatabase.net/k090308/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sleeve, Limb, Compressible (JOW)
Date received	Feb 6, 2009
Decision date	May 1, 2009
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stortford Medical, LLC</b>
Location	West Windsor, NJ, US
Contact	CHRIS DAUGHTERY
510(k) history	2 submissions · 2 cleared · 2009-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090308/> 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 June 1, 2026