

**K132281 ReVive PV (Peripheral Vasculature) Thrombectomy Device**Aug 30, 2013  
38 days to decisionK132281 · Product code: **QEW** · Cardiovascular  
Source: <https://www.510kdatabase.net/k132281/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Peripheral Mechanical Thrombectomy With Aspiration (QEW)
Date received	Jul 23, 2013
Decision date	Aug 30, 2013
Days to decision	38 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Codman &amp; Shurtleff, Inc.</b>
Location	Mchenry, IL, US
Contact	Amarilys Machado
510(k) history	152 submissions · 151 cleared · 1976-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k132281/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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