

**K033657 MOBILE EXTERNAL COUNTER PULSATION SYSTEM  
ANGIONEW-V**Dec 19, 2003  
28 days to decisionK033657 · Product code: **DRN** · Cardiovascular  
Source: <https://www.510kdatabase.net/k033657/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Device, Counter-pulsating, External (DRN)
Date received	Nov 21, 2003
Decision date	Dec 19, 2003
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Living Data Technologies Corporation</b>
Location	Great Neck, NY, US
Contact	SUSAN D GOLDSTEIN-FALK
510(k) history	2 submissions · 2 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k033657/>. 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 July 3, 2026