

**K141809 RMU-1000 AUTOMATED CHEST COMPRESSION SYSTEM**Oct 27, 2014  
112 days to decisionK141809 · Product code: **DRM** · Cardiovascular  
Source: <https://www.510kdatabase.net/k141809/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Compressor, Cardiac, External (DRM)
Date received	Jul 7, 2014
Decision date	Oct 27, 2014
Days to decision	112 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Defibtech, LLC</b>
Location	Guilford, CT, US
Contact	Ed Horton
510(k) history	9 submissions · 8 cleared · 2002-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k141809/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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