

**K931805 ZOLL PD 5200C ADULT SOLID GEL MULTI-FUNCTION ELECT**Oct 8, 1993  
179 days to decisionK931805 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k931805/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Apr 12, 1993
Decision date	Oct 8, 1993
Days to decision	179 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Zoll Medical Corp</b>
Location	Woburn, MA, US
Contact	BLAKE A CERULLO
510(k) history	33 submissions · 27 cleared · 1993-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931805/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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