

**K934016 POWERCHARGER**Oct 18, 1994  
426 days to decisionK934016 · Product code: **DTE** · Cardiovascular  
Source: <https://www.510kdatabase.net/k934016/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pulse-generator, Pacemaker, External (DTE)
Date received	Aug 18, 1993
Decision date	Oct 18, 1994
Days to decision	426 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Zoll Medical Corp</b>
Location	Woburn, MA, US
Contact	FREDERICK W FALLER
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/k934016/> 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 6, 2026