

**K092598 ZOLL E SERIES WITH SPCO/SPMET OPTION**Dec 3, 2009  
101 days to decisionK092598 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k092598/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Aug 24, 2009
Decision date	Dec 3, 2009
Days to decision	101 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Zoll Medical Corporation, World Wide Headquarters</b>
Location	Chelmsford, MA, US
Contact	EILEEN M BOYLE
510(k) history	21 submissions · 21 cleared · 2007-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092598/> 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 5, 2026