FDA 510(k) Premarket Notification

# K092598 ZOLL E SERIES WITH SPCO/SPMET OPTION

**CLEARED**

Dec 3, 2009
101 days to decision

K092598 · 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 | **Zoll Medical Corporation, World Wide Headquarters** |
| Location | Chelmsford, MA, US |
| Contact | EILEEN M BOYLE |
| 510(k) history | 21 submissions · 21 cleared · 2007-2015 |