

K081081 ZOLL E SERIES WITH BLUETOOTH DIAL UP NETWORKINGMar 9, 2009
327 days to decisionK081081 · Product code: **MKJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k081081/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Apr 16, 2008
Decision date	Mar 9, 2009
Days to decision	327 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k081081/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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