FDA 510(k) Premarket Notification

K111594 ZOLL E SERIES

CLEARED

Aug 17, 2011
71 days to decision

K111594 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k111594/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jun 7, 2011
Decision date	Aug 17, 2011
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Zoll Medical Corporation, World Wide Headquarters**
Location	Chelmsford, MA, US
Contact	CHUCK KOLIFRATH
510(k) history	21 submissions · 21 cleared · 2007-2015