FDA 510(k) Premarket Notification

K110168 ZOLL E SERIES WITH 2010 AHA GUIDLINES SOFTWARE UPDATE

CLEARED

Feb 17, 2011
28 days to decision

K110168 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k110168/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jan 20, 2011
Decision date	Feb 17, 2011
Days to decision	28 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