FDA 510(k) Premarket Notification

K081081 ZOLL E SERIES WITH BLUETOOTH DIAL UP NETWORKING

CLEARED

Mar 9, 2009
327 days to decision

K081081 · 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