FDA 510(k) Premarket Notification

K072923 ZOLL E SERIES DEFIBRILLATOR

CLEARED

Apr 24, 2008
192 days to decision

K072923 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k072923/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Oct 15, 2007
Decision date	Apr 24, 2008
Days to decision	192 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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