FDA 510(k) Premarket Notification

K140502 ZOLL E SERIES ALS

CLEARED

Nov 6, 2014
252 days to decision

K140502 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k140502/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Feb 27, 2014
Decision date	Nov 6, 2014
Days to decision	252 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**ZOLL Medical Corporation**
Location	Chelmsford, MA, US
Contact	TANMAY B SHUKLA
510(k) history	30 submissions · 30 cleared · 2005-2024