

K140502 ZOLL E SERIES ALSNov 6, 2014
252 days to decisionK140502 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k140502/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 5, 2026