FDA 510(k) Premarket Notification

K100654 ZOLL PROPAQ MD

CLEARED

Jul 29, 2010
143 days to decision

K100654 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k100654/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Mar 8, 2010
Decision date	Jul 29, 2010
Days to decision	143 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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