FDA 510(k) Premarket Notification

K081574 ZOLL R SERIES

CLEARED

Sep 18, 2008
105 days to decision

K081574 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k081574/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jun 5, 2008
Decision date	Sep 18, 2008
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**ZOLL Medical Corporation**
Location	Chelmsford, MA, US
Contact	EILEEN M BOYLE
510(k) history	30 submissions · 30 cleared · 2005-2024