FDA 510(k) Premarket Notification

K032439 ZOLL M SERIES WITH RECTILINEAR BI-PHASIC OPTION (M SERIES BI-PHASIC)

CLEARED

Nov 5, 2003
90 days to decision

K032439 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k032439/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Aug 7, 2003
Decision date	Nov 5, 2003
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Zoll Medical Corp**
Location	Woburn, MA, US
Contact	SCOTT AUGUST
510(k) history	33 submissions · 27 cleared · 1993-2014