FDA 510(k) Premarket Notification

K041892 ZOLL AED PRO EXTERNAL DEFIBRILLATOR

CLEARED

Feb 4, 2005
206 days to decision

K041892 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k041892/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jul 13, 2004
Decision date	Feb 4, 2005
Days to decision	206 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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