FDA 510(k) Premarket Notification

K990762 ZOLL M SERIES WITH RECTILINEAR BI-PHASIC OPTION (M SERIES BI-PHASIC)

CLEARED

Sep 3, 1999
179 days to decision

K990762 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k990762/

SUBMISSION DETAILS

Decision	Substantially Equivalent - ST
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Mar 8, 1999
Decision date	Sep 3, 1999
Days to decision	179 days
Third-party review	No

APPLICANT

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