

**K990762 ZOLL M SERIES WITH RECTILINEAR BI-PHASIC
OPTION (M SERIES BI-PHASIC)**Sep 3, 1999
179 days to decisionK990762 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k990762/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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