FDA 510(k) Premarket Notification

K102174 ZOLL PROPAQ M

CLEARED

Sep 15, 2010
44 days to decision

K102174 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k102174/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Aug 2, 2010
Decision date	Sep 15, 2010
Days to decision	44 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Zoll Medical Corporation, World Wide Headquarters**
Location	Chelmsford, MA, US
Contact	PAUL DIAS
510(k) history	21 submissions · 21 cleared · 2007-2015

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k102174/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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