FDA 510(k) Premarket Notification

K121367 ZOLL PROPAQ XM

CLEARED

Jun 21, 2012
45 days to decision

K121367 · Product code: **MHX** · Cardiovascular
Source: https://www.510kdatabase.net/k121367/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	May 7, 2012
Decision date	Jun 21, 2012
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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