

K131898 A SERIES AND Q SERIES PATIENT MONITORSFeb 18, 2014
238 days to decisionK131898 · Product code: **MWI** · Cardiovascular
Source: <https://www.510kdatabase.net/k131898/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Jun 25, 2013
Decision date	Feb 18, 2014
Days to decision	238 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Guangdong Biolight Meditech Co., Ltd.
Location	Shanghai, CN
Contact	Diana Hong
510(k) history	21 submissions · 21 cleared · 2008-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k131898/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 18, 2026