

**K120193 ANY VIEW PATIENT MONITORS**Jun 29, 2012  
158 days to decisionK120193 · Product code: **MWI** · CardiovascularSource: <https://www.510kdatabase.net/k120193/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Jan 23, 2012
Decision date	Jun 29, 2012
Days to decision	158 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120193/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 18, 2026