

**K123711 PATIENT MONITOR**Oct 1, 2013  
301 days to decisionK123711 · Product code: **MWI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k123711/>**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	Dec 4, 2012
Decision date	Oct 1, 2013
Days to decision	301 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Shenzhen Creative Industry Co., Ltd.</b>
Location	Flintville, TN, US
Contact	CHARLIE MACK
510(k) history	11 submissions · 11 cleared · 2007-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k123711/> 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 May 27, 2026