

**K092970 GENERAL PATIENT MONITORS, MODELS: G3C/ G3D/  
G3F/ G3G/ G3H**Feb 2, 2010  
130 days to decisionK092970 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k092970/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Sep 25, 2009
Decision date	Feb 2, 2010
Days to decision	130 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>General Meditech, Inc.</b>
Location	Shanghai, CN
Contact	LEE FU
510(k) history	2 submissions · 2 cleared · 2009-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092970/>. 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 24, 2026