

**K060058 STELLAR 404T**Jan 20, 2006  
11 days to decisionK060058 · Product code: **MWI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k060058/>**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 9, 2006
Decision date	Jan 20, 2006
Days to decision	11 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Larsen &amp; Toubro Limited</b>
Location	Crofton, MD, US
Contact	HARRY GUGNANI
510(k) history	18 submissions · 18 cleared · 2003-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k060058/> 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 18, 2026