

**K090172 STAR 55 MODEL 100**Jun 5, 2009  
133 days to decisionK090172 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k090172/>**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	Jan 23, 2009
Decision date	Jun 5, 2009
Days to decision	133 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Larsen &amp; Toubro Limited</b>
Location	Crofton, MD, US
Contact	S.B. BHOSALE
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/k090172/> 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