

**K103686 STAR 50 N**Mar 15, 2011  
88 days to decisionK103686 · Product code: **MWI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k103686/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient (without Arrhythmia Detection Or Alarms) (MWI)
Date received	Dec 17, 2010
Decision date	Mar 15, 2011
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k103686/> 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