

**K032871 PLANET MONITORING SYSTEM**Nov 10, 2003  
56 days to decisionK032871 · Product code: **MWI** · CardiovascularSource: <https://www.510kdatabase.net/k032871/>**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	Sep 15, 2003
Decision date	Nov 10, 2003
Days to decision	56 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	E.J. Smith
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/k032871/> 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