

**K100155 TRUVUE WIRELESS AMBULATORY ECG  
MONITORING SYSTEM**Apr 20, 2010  
90 days to decisionK100155 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k100155/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jan 20, 2010
Decision date	Apr 20, 2010
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Biomedical Systems Corp.</b>
Location	St. Louis, MO, US
Contact	MICHAEL KROEHNKE
510(k) history	2 submissions · 2 cleared · 2003-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k100155/> Data sourced from FDA 510(k) public records (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 June 21, 2026