

**K093134 CERNER CAREAWARE IBUS**Nov 27, 2009  
53 days to decisionK093134 · Product code: **DRG** · Cardiovascular  
Source: <https://www.510kdatabase.net/k093134/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Oct 5, 2009
Decision date	Nov 27, 2009
Days to decision	53 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Cerner Corp.</b>
Location	Kansas City, MO, US
Contact	SHELLEY S LOOBY
510(k) history	7 submissions · 7 cleared · 1996-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093134/> 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 17, 2026