

**K062976 MODIFICATION TO CIC PRO CLINICAL INFORMATION CENTER**Mar 16, 2007  
168 days to decisionK062976 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k062976/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 29, 2006
Decision date	Mar 16, 2007
Days to decision	168 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ge Medical Systems Information Technologies</b>
Location	Mchenry, IL, US
Contact	LISA M BAUMHARDT
510(k) history	136 submissions · 132 cleared · 1978-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062976/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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