

**K072008 BRAEMAR ER900 WIRELESS SERIES ARRHYTHMIA  
EVENT RECORDER**Oct 3, 2007  
72 days to decisionK072008 · Product code: **DRG** · Cardiovascular  
Source: <https://www.510kdatabase.net/k072008/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Jul 23, 2007
Decision date	Oct 3, 2007
Days to decision	72 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Braemar, Inc.</b>
Location	Burnsville, MN, US
Contact	DARREN DERSHEM
510(k) history	5 submissions · 5 cleared · 2004-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072008/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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