

K071733 DL900 SERIES HOLTER RECORDERJul 24, 2007
28 days to decisionK071733 · Product code: **MWJ** · CardiovascularSource: <https://www.510kdatabase.net/k071733/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrocardiograph, Ambulatory (without Analysis) (MWJ)
Date received	Jun 26, 2007
Decision date	Jul 24, 2007
Days to decision	28 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

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