FDA 510(k) Premarket Notification

CLEARED

K072008 BRAEMAR ER900 WIRELESS SERIES ARRHYTHMIA EVENT RECORDER

Oct 3, 2007
72 days to decision

K072008 · Product code: **DRG** · Cardiovascular
Source: https://www.510kdatabase.net/k072008/

SUBMISSION DETAILS

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

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