

**K091971 AVIVO MOBILE PATIENT MANAGEMENT SYSTEM,
NUVANT MOBILE CARDIAC TELEMETRY (MCT) SYSTEM**

Aug 27, 2009
57 days to decision

K091971 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k091971/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 1, 2009
Decision date	Aug 27, 2009
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Corventis, Inc.
Location	San Jose, CA, US
Contact	MADHURI BHAT
510(k) history	7 submissions · 7 cleared · 2009-2014

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k091971/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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