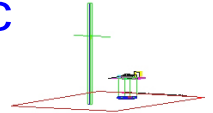


PCTEST Engineering Laboratory, Inc

6660-B Dobbin Road • Columbia, MD 21045 • U.S.A.

TEL (410) 290-6652 • FAX (410) 290-6654

<http://www.pctestlab.com>



CERTIFICATE OF COMPLIANCE

Manufacturer:

CARDIOMEMS INC.

75 Fifth Street, NW Room 205
Atlanta, GA 30308

Dates of Tests: February 6 2004

Test Report S/N: 15.240205064.CDM

Test Site: PCTEST Lab, MD U.S.A.

FCC ID:

XXXCS-A-000051

APPLICANT

CARDIOMEMS INC.

FCC Rule Part(s):

FCC Part 15.209

FCC Classification:

Low Power Transmitter

Equipment EUT Type:

CardioMEMS Wireless AAA Pressure Sensor

Trade Name:

CARDIOMEMS

TX Frequency:

30 - 37.5 MHz

The device bearing the FCC ID number specified above has been shown to comply with the applicable technical standards as indicated in the measurement report and has been tested in accordance with the applicable measurement procedures specified is ANSI C63.4-1992. (See Test Report if any modifications were made for compliance).

I attest to the accuracy of data. All measurements reported herein were performed by me or were made under my supervision and are correct to the best of my knowledge and belief. I assume full responsibility for the completeness of these measurements and vouch for the qualifications of all persons taking them.

NVLAP accreditation does not constitute any product endorsement by NVLAP or any agency of the United States Government
PCTEST certifies that no party to this application has been denied the FCC benefits pursuant to Section 5301 of the
Anti-Drug Abuse Act of 1988, 21 U.S.C. 862.


Randy Ortanez
President



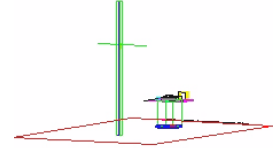
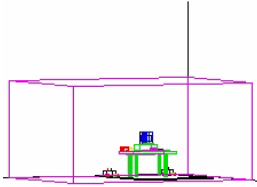
NVLAP
Lab Code 100431-0

TABLE OF CONTENTS

ATTACHMENT A:	COVER LETTER(S)	
ATTACHMENT B:	ATTESTATION STATEMENT(S)	
ATTACHMENT C:	TEST REPORT	
	SCOPE	1
	INTRODUCTION (SITE DESCRIPTION)	2
	PRODUCTION INFORMATION	3
	DESCRIPTION OF TESTS (CONDUCTED)	4
	DESCRIPTION OF TESTS (RADIATED)	5
	TEST EQUIPMENT	6
	TEST DATA (CONDUCTED)	7
	TEST DATA (RADIATED)	8
	EXTRAPOLATED TO 3 METER	9
	SAMPLE CALCULATIONS	10
	ACCURACY OF MEASUREMENT	11
	LIST OF TEST EQUIPMENT	12
	TEST SOFTWARE USED	13
	CONCLUSION	14
ATTACHMENT D:	TEST PLOTS	
ATTACHMENT E:	FCC ID LABEL & LOCATION	
ATTACHMENT F:	BLOCK DIAGRAM	
ATTACHMENT G:	SCHEMATICS	
ATTACHMENT H:	TEST SETUP PHOTOGRAPHS	
ATTACHMENT I:	EXTERNAL PHOTOS	
ATTACHMENT J:	INTERNAL PHOTOS	
ATTACHMENT K:	USERS MANUAL	

PCTEST™ PT. 15. REPORT	 Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CDM	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXXX	Page i of i

MEASUREMENT REPORT



Scope - Measurement and determination of electromagnetic emissions (EME) of radio frequency devices including intentional and/or unintentional radiators for compliance with the technical rules and regulations of the Federal Communications Commission.

Company Name: CARDIOMEMS INC.
Address: 75 Fifth Street, NW Room 205
Atlanta, GA 30308

- Trade Name(s): **CARDIOMEMS**
- FCC ID: **XXXCS-A-000051**
- EUT Type: CardioMEMS Wireless AAA Pressure Sensor
- FCC Rule Part(s): FCC Part 15.209
- Classification: Low Power Transmitter
- Test Procedure(s): ANSI C63.4 (1992)/EN55022: 1998(CISPR 22) w/A1)
- Dates of Tests: February 6, 2004
- Place of Tests: PCTEST Lab, Columbia, MD U.S.A.
- Test Report S/N: 15.240205064.CDM



PCTEST™ PT. 15 REPORT	Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CDM	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXX	Page 1 of 14

1.1 INTRODUCTION

The measurement procedure described in American National Standard for Methods of Measurement of Radio-Noise Emissions from Low-Voltage Electrical and Electronic Equipment in the Range of 9kHz to 40GHz (ANSI C63.4-1992) was used in determining radiated and conducted emissions emanating from **CardioMEMS Wireless AAA Pressure Sensor FCC ID: XXXCS-A-000051**.

These measurement tests were conducted at **PCTEST Engineering Laboratory, Inc.** facility in New Concept Business Park, Guilford Industrial Park, Columbia, Maryland. The site address is 6660-B Dobbin Road, Columbia, MD 21045. The test site is one of the highest points in the Columbia area with an elevation of 390 feet above mean sea level. The site coordinates are 39° 11'15" N latitude and 76° 49'38" W longitude. The facility is 1.5 miles North of the FCC laboratory, and the ambient signal and ambient signal strength are approximately equal to those of the FCC laboratory. There are no FM or TV transmitters within 15 miles of the site. The detailed description of the measurement facility was found to be in compliance with the requirements of § 2.948 according to ANSI C63.4 on October 19, 1992.

PCTEST™ PT. 15 REPORT	 Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXXX	Page 2 of 14

1.2 PCTEST Location

The map at right shows the location of the PCTEST Lab, its proximity to the FCC Lab, the Columbia vicinity area, the Baltimore-Washington International (BWI) airport, and the city of Baltimore, and the Washington, D.C. area. (see Figure1).

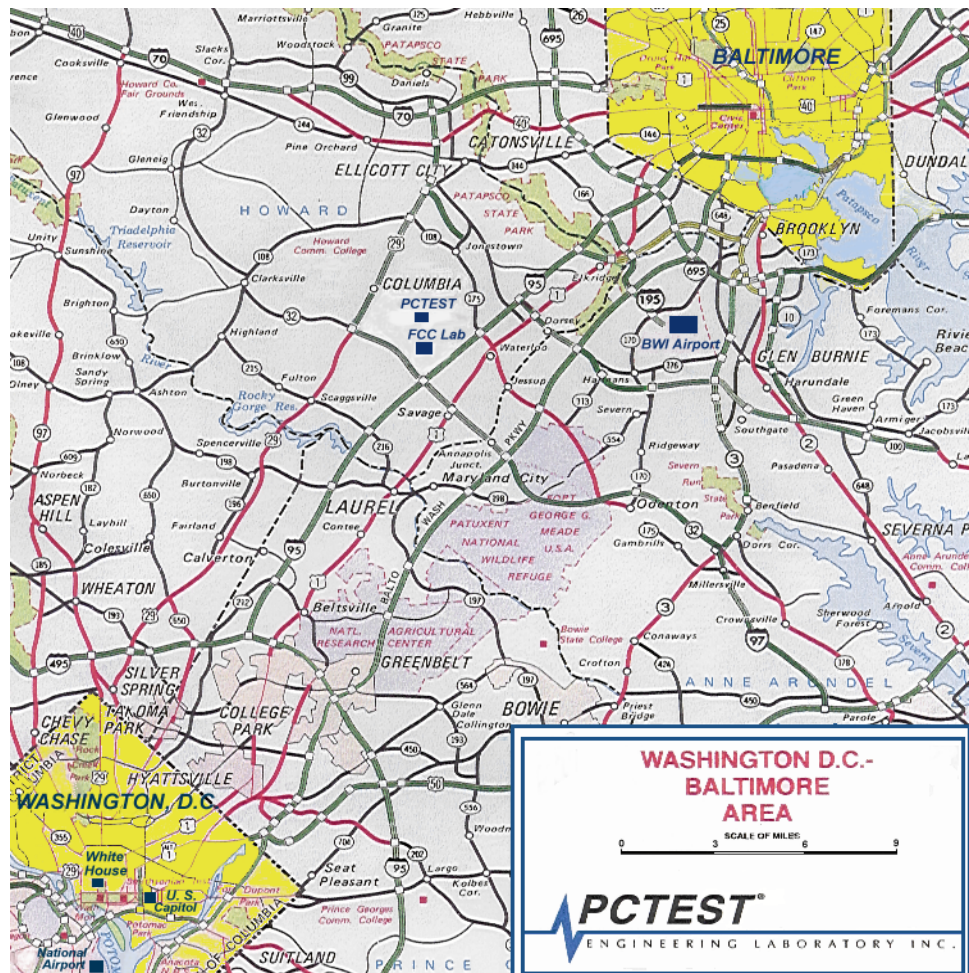


Figure 1. Map of the Greater Baltimore and Metropolitan Washington, D.C. area.

PCTEST™ PT. 15 REPORT	 Complete Wireless Lab™ www.pctestlab.com Measurement Report Reviewed By: Quality Manager			
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004.	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXX	Page 3 of 14

2.1 Product Information

2.2 Equipment Description

The Equipment Under Test (EUT) is a Wireless Pressure Sensor designed to be placed within a human vascular blood vessel for the purpose of providing a non-invasive capability for determining blood pressure. The system is only used in hospitals and similar medical facilities in a post operative period of approximately 2 weeks following surgery and, in unusual cases infrequently thereafter. Thus, proliferation of these devices over time is extremely limited.

2.3 EMI Suppression Device(s)

EMI suppression device(s) added and/or modified during testing:

- none

PCTEST™ PT. 15 REPORT	 Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXXX	Page 4 of 14

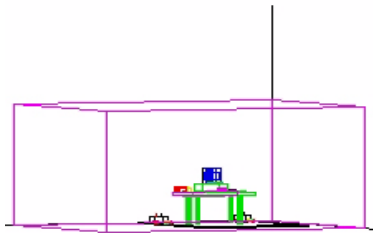


Figure 2. Shielded Enclosure Line-Conducted Test Facility

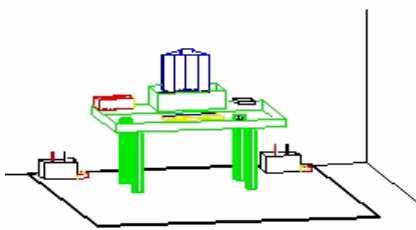


Figure 3. Line Conducted Emission Test Set-Up

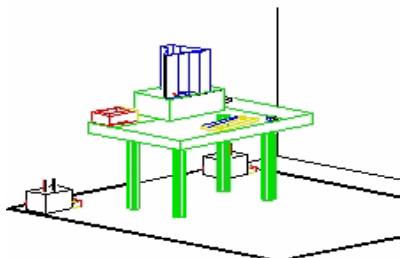


Figure 4. Wooden Table & Bonded LISNs

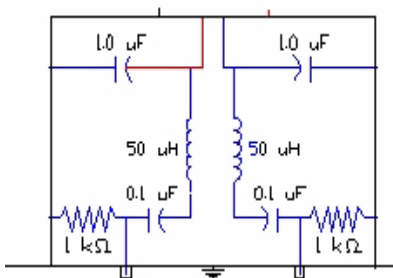


Figure 5. LISN Schematic

3.1 Description of Tests

3.2 Conducted Emissions

The line-conducted facility is located inside a 16'x20'x10' shielded enclosure. It is manufactured by Ray Proof Series 81 (see Figure 2). The shielding effectiveness of the shielded room is in accordance with MIL-Std-285 or NSA 65-6. A 1m. x 1.5m. wooden table 80cm. high is placed 40cm. away from the vertical wall and 1.5m away from the side wall of the shielded room (see Figure 3). Solar Electronics and EMC0 Model 3725/2 (10kHz-30MHz) 50Ω/50μH Line-Impedance Stabilization Networks (LISNs) are bonded to the shielded room (see Figure 4). The EUT is powered from the Solar LISN and the support equipment is powered from the EMC0 LISN. Power to the LISNs are filtered by a high-current high-insertion loss Ray Proof power line filters (100dB 14kHz-10GHz). The purpose of the filter is to attenuate ambient signal interference and this filter is also bonded to the shielded enclosure. All electrical cables are shielded by braided tinned copper zipper tubing with inner diameter of 1/2". If the EUT is a DC-powered device, power will be derived from the source power supply it normally will be powered from and this supply lines will be connected to the Solar LISN. LISN schematic diagram is shown in Figure 5. All interconnecting cables more than 1 meter were shortened by non-inductive bundling (serpentine fashion) to a 1-meter length. Sufficient time for the EUT, support equipment, and test equipment was allowed in order for them to warm up to their normal operating condition. The RF output of the LISN was connected to the spectrum analyzer to determine the frequency producing the maximum EME from the EUT. The spectrum was scanned from 450kHz to 30MHz with 20 msec. sweep time. The frequency producing the maximum level was reexamined using EMI/ Field Intensity Meter and Quasi-Peak adapter. The detector function was set to CISPR quasi-peak mode. The bandwidth of the receiver was set to 10 kHz. The EUT, support equipment, and interconnecting cables were arranged and manipulated to maximize each EME

PCTEST™ PT. 15 REPORT		Measurement Report		Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXXX	Page 5 of 14

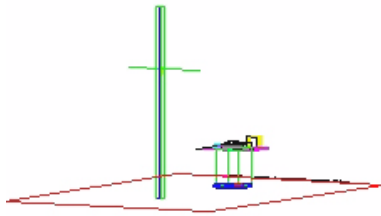


Figure 6. 3-Meter Test Site

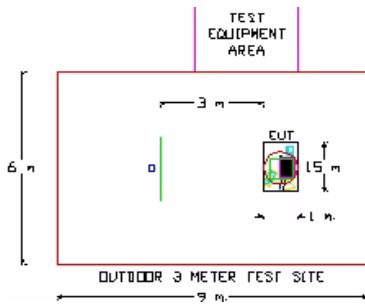


Figure 7. Dimensions of Outdoor Test Site

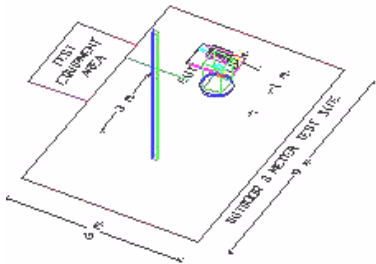


Figure 8. Turntable and System Setup

emission. Each emission was maximized by: switching power lines; varying the mode of operation or resolution; clock or data exchange speed; scrolling H pattern to the EUT and/or support equipment, and powering the monitor from the floor mounted outlet box and the computer aux AC outlet, if applicable; whichever determined the worst-case emission. Photographs of the worst-case emission can be seen in Appendix D. Each EME reported was calibrated using the HP8640B signal generator.

3.1 Description of Tests (continued)

3.3 Radiated Emissions

Preliminary measurements were made indoors at 1 meter using broadband antennas, broadband amplifier, and spectrum analyzer to determine the frequency producing the maximum EME. Appropriate precaution was taken to ensure that all EME from the EUT were maximized and investigated. The system configuration, clock speed, mode of operation or video resolution, turntable azimuth with respect to the antenna were noted for each frequency found. The spectrum was scanned from 30 to 200 MHz using biconical antenna and from 200 to 1000 MHz using log-spiral antenna. From 1 to 4.2 GHz, linearly polarized double ridge horn antennas were used.

Final measurements were made outdoors at 3-meter test range using Roberts™ Dipole antennas or horn antenna (see Figure 6). The test equipment was placed on a wooden and plastic bench situated on a 1.5 x 2 meter area adjacent to the measurement area (see Figure 7). Sufficient time for the EUT, support equipment, and test equipment was allowed in order for them to warm up to their normal operating condition. Each frequency found during pre-scan measurements was re-examined and investigated using EMI/Field Intensity Meter and Quasi-Peak Adapter. The detector function was set to CISPR quasi-peak mode and the bandwidth of the receiver was set to 100kHz or 1 MHz depending on the frequency or type of signal.

PCTEST™ PT. 15 REPORT			Measurement Report		Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXX	Page 6 of 14	

The half-wave dipole antenna was tuned to the frequency found during preliminary radiated measurements. The EUT, support equipment and interconnecting cables were re-configured to the set-up producing the maximum emission for the frequency and were placed on top of a 0.8-meter high non-metallic 1 x 1.5 meter table (see Figure 8). The EUT, support equipment, and interconnecting cables were re-arranged and manipulated to maximize each EME emission. The turntable containing the system was rotated; the antenna height was varied 1 to 4 meters and stopped at the azimuth or height producing the maximum emission. Each emission was maximized by: varying the mode of operation or resolution; clock or data exchange speed; scrolling H pattern to the EUT and/or support equipment, and powering the monitor from the floor mounted outlet box and the computer aux AC outlet, if applicable; and changing the polarity of the antenna, whichever determined the worst-case emission. Photographs of the worst-case emission can be seen in Appendix C. Each EME reported was calibrated using the HP8640B signal generator. The Theoretical Normalized Site Attenuation Curves for both horizontal and vertical polarization are shown in Figure 9.

4.1 Support Equipment Used

(See "Appendix G - Test Photographs" for actual system test setup.)

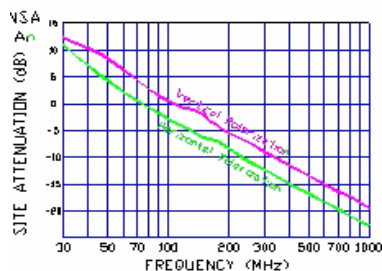


Figure 9. Normalized Site Attenuation Curves (H&V)

PCTEST™ PT. 15 REPORT	 Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXX	Page 7 of 14

5.1 LINE-CONDUCTED TEST DATA

5.2 Conducted Emissions

- **

Device is not powered via AC mains

(See Plots - ATTACHMENT D)(N/A)

NOTES:

1. All modes of operation were investigated and the worst-case emissions are reported.
2. The limit for a Class B device is Specified in EN55022 Standard from 150kHz to 30MHz.
3. Line A = Phase; Line B = Neutral
4. Deviations to the Specifications: None.

* All readings are calibrated by HP8640B signal generator with accuracy traceable to the National Institute of Standards and Technology (formerly NBS).

PCTEST™ PT. 15 REPORT	 Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXX	Page 8 of 14

** Measurements using CISPR quasi-peak mode.

PCTEST™ PT. 15 REPORT	 Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXXX	Page 9 of 14

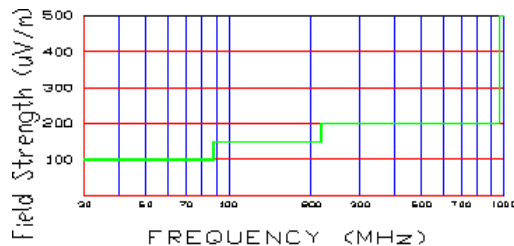
6.1 RADIATED TEST DATA

6.2 Radiated and Spurious Emissions

FREQ. (MHz)	Level* (dBm)	AFCL** (dB)	POL (H/V)	Height (m)	Azimuth (° angle)	F/S (μV/m)	Margin* ** (dB)
36.0	- 92	0.24	H	3.2	180	5.8	- 24.8

Table 1. Radiated Measurements at 3-meters.

**All other emissions were more than 30 dB below the limit.*



NOTES:

1. All modes of operation were investigated and the worst-case emissions are reported.
2. The radiated limits are shown on Figure A-1. Above 1 GHz the limit is 500μV/m.

Figure 10. Limits at 3 meters

- * All readings are calibrated by HP8640B signal generator with accuracy traceable to the National Institute of Standards and Technology (formerly NBS).
- ** AFCL = Antenna Factor (Roberts dipole) and Cable Loss (30 ft. RG58C/U).
- *** Measurements using CISPR quasi-peak mode. Above 1GHz, peak detector function mode is used using a resolution bandwidth of 1MHz and a video bandwidth of 1MHz. The peak level complies with the average limit. Peak mode is used with linearly polarized horn antenna and low-loss microwave cable.

Measurement Procedure

In order to determine the radiated field strength of the wireless AAA Pressure sensor transmitter it was necessary to use a substitution technique.

A signal test point on the associated receiver section was coupled to an oscilloscope. The separation distance from the membrane type transmitter (EUT) to its associated receiver antenna was initially 3.16 cm. The EUT was transmitting at max power and the receiver test point level was measured with an oscilloscope and recorded. The same level at the Rx section test point was then matched by a signal generator feeding a dipole antenna (substituted transmitter) 3 meters away and the level of the signal generator was recorded. This process was repeated with the membrane transmitter moved as far as possible away from

PCTEST™ PT. 15 REPORT	 Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXX	Page 10 of 14

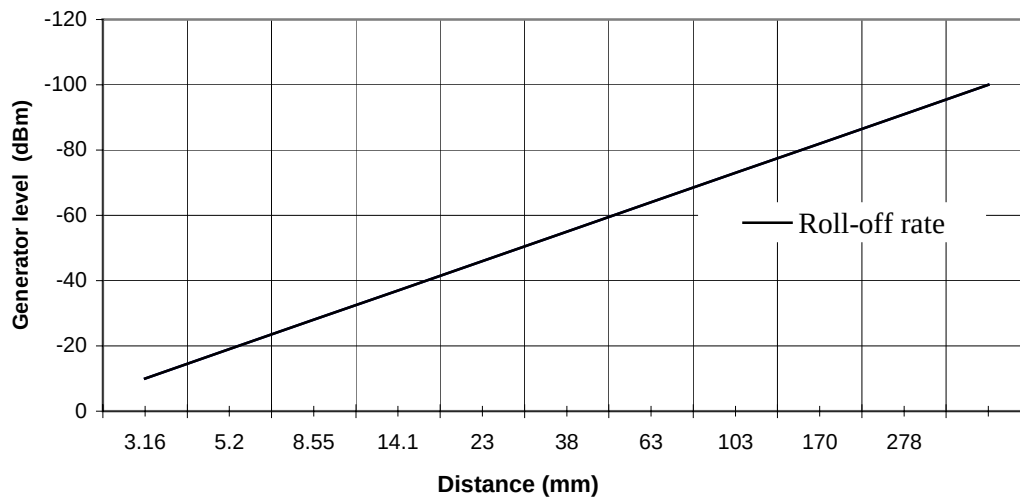
the receiver antenna. This data was then used to extrapolate the data to a 3 meter distance. Levels reported are peak detector levels.

PCTEST™ PT. 15 REPORT	 Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXXX	Page 11 of 14

7.1 Data used to extrapolate to 3 meter (300mm)

d (mm)	To match TP1 Sig GEN (dBm)	Frequency	Data Type
3.16	- 10	36 MHz	Measured
5.2	- 19	36 MHz	Measured
8.6	- 28	36 MHz	Measured
14.1	- 37	36 MHz	Trendline
23	- 46	36 MHz	Trendline
38	- 55	36 MHz	Trendline
63	- 64	36 MHz	Trendline
103	- 73	36 MHz	Trendline
170	- 82	36 MHz	Trendline
278	- 91	36 MHz	Trendline
300.00	-92	36 MHz	Trendline

Wireless Blood Pressure Transmitter Attenuation w/distance



PCTEST™ PT. 15 REPORT	Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXX	Page 12 of 14

8.1 Sample Calculations

$$\begin{aligned} \text{dB}\mu\text{V} &= 20 \log_{10} (\mu\text{V}/\text{m}) \\ \text{dB}\mu\text{V} &= \text{dBm} + 107 \end{aligned}$$

8.2 Example 1:

@ 20.3 MHz

$$\begin{aligned} \text{Class B limit} &= 250 \mu\text{V} = 47.96 \text{ dB}\mu\text{V} \\ \text{Reading} &= -67.8 \text{ dBm (calibrated level)} \\ \text{Convert to dB}\mu\text{V} &= -67.8 + 107 = 39.2 \text{ dB}\mu\text{V} \\ 10^{(39.2/20)} &= 91.2 \mu\text{V} \\ \text{Margin} &= 39.2 - 47.96 = -8.76 \\ &= \mathbf{8.8 \text{ dB below limit}} \end{aligned}$$

8.3 Example 2:

@ 66.7 MHz

$$\begin{aligned} \text{Class B limit} &= 100 \mu\text{V}/\text{m} = 47.96 \text{ dB}\mu\text{V}/\text{m} \\ \text{Reading} &= -76.0 \text{ dBm (calibrated level)} \\ \text{Convert to dB}\mu\text{V}/\text{m} &= -76.0 + 107 = 31.0 \text{ dB}\mu\text{V}/\text{m} \\ \text{Antenna Factor + Cable Loss} &= 5.8 \text{ dB} \\ \text{Total} &= 36.8 \text{ dB}\mu\text{V}/\text{m} \\ \text{Margin} &= 36.8 - 40.0 = -3.2 \\ &= \mathbf{3.2 \text{ dB below limit}} \end{aligned}$$

PCTEST™ PT. 15 REPORT	 Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXX	Page 13 of 14

9.1 Accuracy of Measurement

9.2 Measurement Uncertainty Calculations:

The measurement uncertainties stated were calculated in accordance with the requirements of NIST Technical Note 1297 and NIS 81 (1994).

Contribution (Line Conducted)	Probability Distribution	Uncertainty (± dB)	
		9kHz-150MHz	150-30MHz
Receiver specification	Rectangular	1.5	1.5
LISN coupling specification	Rectangular	1.5	1.5
Cable and input attenuator calibration	Normal (k=2)	0.3	0.5
Mismatch: Receiver VRC $\Gamma_1 = 0.03$ LISN VRC $\Gamma_R = 0.8$ (9kHz) 0.2 (30MHz) Uncertainty limits $20\log(1 \pm \Gamma_1 \Gamma_R)$	U-Shaped	0.2	0.35
System repeatability	Std. deviation	0.2	0.05
Repeatability of EUT		-	-
Combined standard uncertainty	Normal	1.26	1.30
Expanded uncertainty	Normal (k=2)	2.5	2.6

Calculations for 150kHz to 30MHz:

$$u_c(y) = \sqrt{\sum_{i=1}^m u_i^2(y)} = \pm \sqrt{\frac{1.5^2 + 1.5^2}{3} + \left(\frac{0.5}{2}\right)^2 + 0.35} = \pm 1.298 \text{ dB}$$

$$U = 2U_c(y) = \pm 2.6 \text{ dB}$$

Contribution (Radiated Emissions)	Probability Distribution	Uncertainties (± dB)	
		3 m	10 m
Ambient Signals		-	-
Antenna factor calibration	Normal (k=2)	± 1.0	± 1.0
Cable loss calibration	Normal (k=2)	± 0.5	± 0.5
Receiver specification	Rectangular	± 1.5	± 1.5
Antenna directivity	Rectangular	+ 0.5 / - 0	+ 0.5
Antenna factor variation with height	Rectangular	± 2.0	± 0.5
Antenna phase centre variation	Rectangular	0.0	± 0.2
Antenna factor frequency interpolation	Rectangular	± 0.25	± 0.25
Measurement distance variation	Rectangular	± 0.6	± 0.4
Site imperfections	Rectangular	± 2.0	± 2.0
Mismatch: Receiver VRC $\Gamma_1 = 0.2$ Antenna VRC $\Gamma_R = 0.67$ (Bi) 0.3 (Lp) Uncertainty limits $20\log(1 \pm \Gamma_1 \Gamma_R)$	U-Shaped	+ 1.1 - 1.25	± 0.5
System repeatability	Std. Deviation	± 0.5	± 0.5
Repeatability of EUT		-	-
Combined standard uncertainty	Normal	+ 2.19 / - 2.21	+ 1.74 / - 1.72
Expanded uncertainty U	Normal (k=2)	+ 4.38 / - 4.42	+ 3.48 / - 3.44

Calculations for 3m biconical antenna. Coverage factor of k=2 will ensure that the level of confidence will be approximately 95%, therefore:

PCTEST™ PT. 15 REPORT	 Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXX	Page 14 of 14

$$U=2u_c(y) = 2 \times \pm 2.19 = \pm 4.38\text{dB}$$

10.1 Test Equipment

10.2 Type	Model	Cal. Due Date	S/N
Microwave Spectrum Analyzer	HP 8566B (100Hz-22GHz) 12/05/04	3638A08713	
Microwave Spectrum Analyzer	HP 8566B (100Hz-22GHz) 04/17/04	2542A11898	
Spectrum Analyzer/Tracking Gen.	HP 8591A (9kHz-1.8GHz)	06/02/04	3144A02458
Spectrum Analyzer	HP 8591A (9kHz-1.8GHz)	10/15/04	3108A02053
Spectrum Analyzer	HP 8594A (9kHz-2.9GHz)	11/02/04	3051A00187
Signal Generator*	HP 8640B (500Hz-1GHz)	06/02/04	2232A19558
Signal Generator*	HP 8640B (500Hz-1GHz)	06/02/04	1851A09816
Signal Generator*	Rohde & Schwarz (0.1-1000MHz)	09/11/04	894215/012
Ailtech/Eaton Receiver	NM 37/57A-SL (30-1000MHz)	04/12/04	0792-03271
Ailtech/Eaton Receiver	NM 37/57A (30-1000MHz)	03/11/05	0805-03334
Ailtech/Eaton Receiver	NM 17/27A (0.1-32MHz)	09/17/04	0608-03241
Quasi-Peak Analyzer	HP 85650A	08/09/04	2043A00301
Ailtech/Eaton Analyzer	CCA-7 CISPR/ANSI QP Analyzer	03/11/05	0194-04082
RG58 Coax Test Cable	No. 167		n/a
Harmonic/Flicker Test System	HP 6841A (IEC 555-2/3)		3531A00115
Broadband Amplifier (2)	HP 8447D		1145A00470, 1937A03348
Broadband Amplifier	HP 8447F		2443A03784
Transient Limiter	HP 11947A (9kHz-200MHz)		2820A00300
Horn Antenna	EMCO Model 3115 (1-18GHz)		9704-5182
Horn Antenna	EMCO Model 3115 (1-18GHz)		9205-3874
Horn Antenna	EMCO Model 3116 (18-40GHz)		9203-2178
Biconical Antenna (4)	Eaton 94455/Eaton 94455-1/Singer 94455-1/Compliance Design 1295, 1332, 0355		
Log-Spiral Antenna (3)	Ailtech/Eaton 93490-1		0608, 1103, 1104
Roberts Dipoles	Compliance Design (1 set) A100		5118
Ailtech Dipoles	DM-105A (1 set)		33448-111
EMCO LISN (2)	3816/2		1077, 1079
EMCO LISN	3725/2		2009
Microwave Pre-amplifier 40dB Gain	HP 83017A (0.5-26.5GHz)		3123A00181
Microwave Cables	MicroCox (1.0-26.5GHz)		
Ailtech/Eaton Receiver	NM37/57A-SL		0792-03271
Spectrum Analyzer	HP 8591A		3034A01395
Modulation Analyzer	HP 8901A		2432A03467
NTSC Pattern Generator	Levler 408		0377433
Noise Figure Meter	HP 8970B		3106A02189
Noise Figure Meter	Ailtech 7510		TE31700
Noise Generator	Ailtech 7010		1473
Microwave Survey Meter	Holaday Model 1501 (2.450GHz)		80931
Digital Thermometer	Extech Instruments 421305		426966
Attenuator	HP 8495A (0-70dB) DC-4GHz		
Bi-Directional Coax Coupler	Narda 3020A (50-1000MHz)		
Shielded Screen Room	RF Lindegren Model 26-2/2-0		6710 (PCT270)
Shielded Semi-Anechoic Chamber	Roy Prof Model S81		R2437 (PCT278)
Environmental Chamber	Associated Systems Model 1025 (Temperature/Humidity)		PCT285

* Calibration traceable to the National Institute of Standards and Technology (NIST).

PCTEST™ PT. 15 REPORT	Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXXXX	Page 15 of 14

11.1 Test Software Used

```

10  CLS:COLOR 7,0
20  FOR I = 1 TO 80
30  PRINT H;
40  NEXT I
50  FOR K= 1 TO 25
60  LPRINT H;
70  NEXT K
80  OPEN COM1:1200,N,8,1,CS0,DS0" FOR OUTPUT AS #1
90  PRINT#1,ATDT,0123456789"
100 CLOSE:GOTO 20

```

NOTE:

This is a sample of a basic program used during testing of various devices. However, during testing, a different software program may be used; whichever determines the worst-case condition. In addition the program used, if any, also depends on the number and type of devices being tested.

PCTEST™ PT. 15 REPORT	 Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXXX	Page 16 of 14

12.1 Conclusion

The data collected shows that the **CARDIOMEMS wireless blood pressure transmitter, FCC ID: XXXCS-A-000051**, is in compliance with §§ 15.207 and 15.209 of the FCC Rules.

No modifications were made to the device.

PHOTOGRAPHS OF EUT



PCTEST™ PT. 15 REPORT	Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXX	Page 17 of 14

PCTEST™ PT. 15 REPORT	 Measurement Report			Reviewed By: Quality Manager
Test Report S/N: 15.240205064.CD M	Test Dates: February 6, 2004	EUT Type: CARDIOMEMS Blood Pressure Implant	FCC ID: XXXXXXXX	Page 18 of 14