

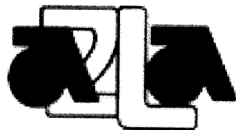
Specific Absorption Rate (SAR) Test Report

for
Cardionet, Inc.



on the
CardioNet ECG Monitor With Arrhythmia Detection
Model Number: CardioNet Monitor
FCC ID: QBI-1005

Test Report: 30388163
Date of Report: March 25, 2003

Job #: 3038816
Date of Test: March 10 to 11, 2003



A2LA Certificate Number: 1755-01

Tested by: 	Suresh Kondapalli
Reviewed by: 	David Chernomordik, Ph.D., EMC Technical Manager

Review Date: 3/31/03

All services undertaken are subject to the following general policy: Reports are submitted for exclusive use of the client to whom they are addressed. Their significance is subject to the adequacy and representative character of the samples and to the comprehensiveness of the tests, examinations or surveys made. This report shall not be reproduced except in full, without written consent of Intertek Testing Services, NA Inc.



Intertek Testing Services NA, Inc.

1365 Adams Court, Menlo Park, CA 94025

Telephone 650-463-2900 Fax 650-463-2910 Home Page www.etlsemko.com



TABLE OF CONTENTS

STATEMENT OF COMPLIANCE	3
1.0 JOB DESCRIPTION	4
1.1 Client Information.....	4
1.2 Equipment under test (EUT).....	4
1.3 Test Plan Reference	5
1.4 System Test Configuration	6
1.4.1 System Block Diagram & Support Equipment.....	6
1.4.2 Test Position	7
1.4.3 Test Condition	8
1.5 Modifications required for compliance.....	8
1.6 Additions, deviations and exclusions from standards.....	8
2.0 SAR EVALUATION	9
2.1 SAR Limits	9
2.2 Configuration Photographs	10
2.3 System Verification	16
2.4 Evaluation Procedures	16
2.5 Test Results.....	17
3.0 TEST EQUIPMENT.....	20
3.1 Equipment List.....	20
3.2 Muscle Tissue Simulating Liquid	22
3.3 E-Field Probe Calibration	23
3.4 Measurement Uncertainty	24
3.5 Measurement Tractability	25
4.0 WARNING LABEL INFORMATION - USA.....	26
5.0 REFERENCES.....	27
6.0 DOCUMENT HISTORY	28
APPENDIX A - SAR Evaluation Data.....	29
APPENDIX B - E-Field Probe Calibration Data	45
APPENDIX C – Phantom Certificate	58

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

STATEMENT OF COMPLIANCE

The Cardionet Monitor, FCC ID: QBI-1005 was evaluated for SAR in accordance with the requirements for RF Exposure compliance testing defined in FCC OET Bulletin 65, Supplement C (Edition 01-01). Testing was performed at the Intertek Testing Services facility in Menlo Park, California.

For the evaluation, the dosimetric assessment system DASY3 was used. The phantom employed was the "SAM Twin Phantom". The total uncertainty for the evaluation of the spatial peak SAR values averaged over a cube of 1g tissue mass had been assessed for this system to be $\pm 27.8\%$.

The device was tested at their maximum output power declared by the Cardionet Inc.

In summary, the maximum spatial peak SAR value for the sample device averaged over 1g was found to be:

Phantom	Position	SAR _{1g} , mW/g
Flat Section	1.5 cm from phantom	0.89

In conclusion, the tested sample device was found to be in compliance with the requirements defined in OET Bulletin 65, Supplement C (Edition 01-01) for body-worn configuration.

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

1.0 JOB DESCRIPTION**1.1 Client Information**

The CardioNet Monitor has been tested at the request of:

Company: Cardionet, Inc.
510 Market Street
San Diego , CA 92101
USA

Name of contact: Mr. Jason Dorsey
Telephone: 619/243-7550
Fax: 619/243-7700

1.2 Equipment under test (EUT)**Product Descriptions:**

Equipment	CardioNet ECG Monitor With Arrhythmia Detection		
Trade Name	CardioNet	P/N.	CardioNet Monitor
FCC ID	QBI-1005	S/N No.	11006
Category	Portable device	RF Exposure	Uncontrolled Environment
Frequency Band	824 - 849 MHz 1850 – 1910 MHz	System	CDMA

EUT Antenna Description			
Type	Monopole	Configuration	Fixed
Dimensions	13.5mm (L), 7mm (D)	Gain	from 0 dBi to -2 dBi
Location	Right side Top		

Use of Product: The Cardionet Monitor constantly monitors patient's heartbeat and sends information to the CardioNet Patient Service Center. When a patient is away from home it uses wireless communication throughout Cellular/ PCS CDMA modem manufactured by Sierra Wireless (SB555 model, FCC ID: N7NSB555).

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

Manufacturer: Cardionet, Inc.

Production is planned: ☒ Yes, ☐ No

EUT receive date: February 18, 2003

EUT received condition: Good working condition prototype. As declared by the Applicant it is identical to the production units.

Test start date: March 10, 2003

Test end date: March 11, 2003

1.3 Test Plan Reference

FCC Rule: Part 2.1093, FCC's OET Bulletin 65, Supplement C (Edition 01-01)

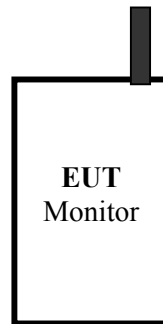
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

1.4 System Test Configuration

1.4.1 System Block Diagram & Support Equipment

The diagram, shown below details test configuration of the equipment under test.



The CardioNet Monitoring System consists of two devices - the CardioNet Sensor and the CardioNet Monitor.

The CardioNet Monitor communicates with the CardioNet Sensor at 902 - 928 MHz (ISM band) and also communicates with CardioNet Patient Service Center in cellular band (824 – 848 MHz) and/or PCS band (1850 – 1910 MHz).

The CardioNet Monitor transmitter operating at ISM band is very low power transmitter (1 mW), The Cellular/ PCS bands transmitter operates at 224 mW power in CDMA mode.

During the SAR test of CDMA transmitter, the ISM transmitter was also active. No Support Equipment was used. The test sample was operated in a test mode that allows control of the transmitter without the need to place actual phone calls. For the purposes of this test the device is commanded to test mode and manually set to the proper channel, transmitter power level and transmit mode of operation. The device was then placed in the SAR Measurement System with a fully charged battery.

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

1.4.2 Test Position

Two test configurations were used to show compliance with the FCC RF Exposure requirements. In all configurations, the CardioNet Monitor was configured for testing in a typical fashion (as a customer would normally use it). Due to the application and usage of the product, SAR measurements with the human head region are not necessary. Table 1 below describes the setup and condition:

Table 1, Equipment Setup	
Configuration	Description
A	<ul style="list-style-type: none"> Back Side 15 mm from phantom. Antenna in horizontal position, distance from antenna to Phantom = 31.5 mm. Simulating close proximity of human body.
B	<ul style="list-style-type: none"> Front Side 15 mm from phantom. Antenna in horizontal position, distance from antenna to Phantom = 28.5 mm Simulating close proximity of human body.

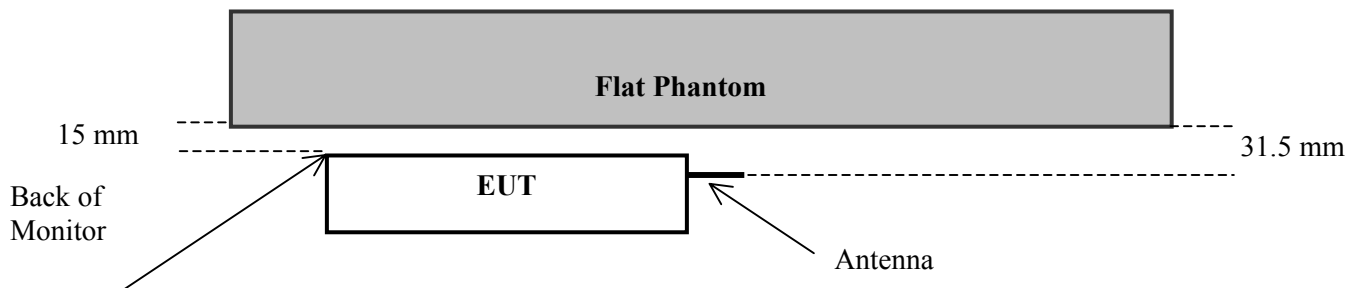


Figure 1: Configuration A Back Side 15mm from Phantom

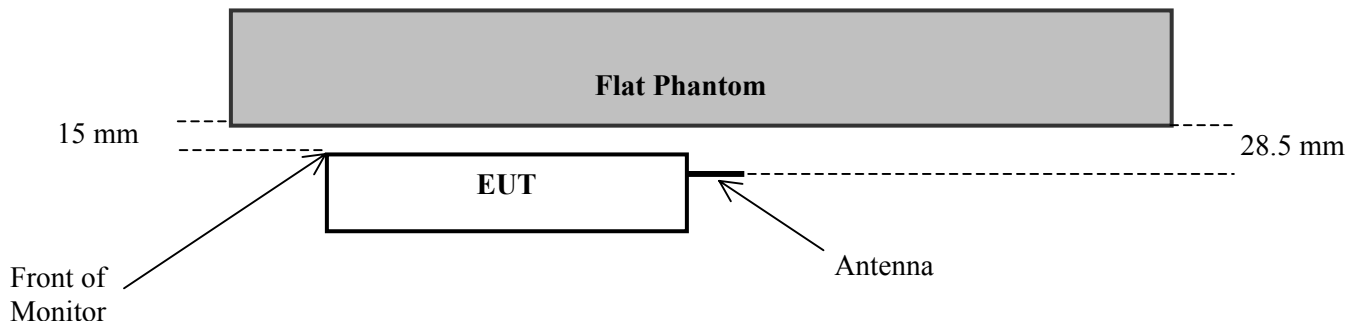


Figure 1: Configuration B Front Side 15mm from Phantom

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

1.4.3 Test Condition

During tests, the worst case data (max. RF coupling) was determined with following conditions:

EUT Antenna	Fixed length	Orientation	N/A
Usage	Stand-alone	Distance between antenna axis at the joint and the phantom surface:	31.5 mm – Back Side 15 mm from phantom. 28.5.5 mm –Front Side 15 mm from phantom
Simulating human Body/hand	body	EUT Battery	Unit powered from battery.
Power output	Frequency MHz		Power Output, dBm
	824.7		23.5
	836.5		23.5
	848.3		23.4
	1851.25		23.5
	1880.0		23.4
	1908.75		23.6

The spatial peak SAR values were accessed for lowest, middle and highest operating channels defined by the manufacturer.

Antenna port power measurement was performed, with the power meter, before and after the SAR tests to ensure that the CardioNet Monitor operated at the highest power level.

1.5 Modifications required for compliance

Intertek Testing Services implemented no modifications.

1.6 Additions, deviations and exclusions from standards

No additions, deviations or exclusions have been made from standard.

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

2.0 SAR EVALUATION

2.1 SAR Limits

The following FCC limits for SAR apply to devices operate in General Population/Uncontrolled Exposure environment:

EXPOSURE (General Population/Uncontrolled Exposure environment)	SAR (W/kg)
Average over the whole body	0.08
Spatial Peak (1g)	1.60
Spatial Peak for hands, wrists, feet and ankles (10g)	4.00

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

2.2 Configuration Photographs

SAR measurement Test Setup

Back Side 15 mm from Phantom



Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

2.2 Configuration Photographs (Continued)

SAR measurement Test Setup

Back Side 15 mm from Phantom



Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

2.2 Configuration Photographs (Continued)

SAR measurement Test Setup

Front Side 15 mm from Phantom



Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

2.2 Configuration Photographs (Continued)

SAR measurement Test Setup

Front Side 15 mm from Phantom



Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

2.2 Configuration Photographs (Continued)

SAR measurement Test Setup

EUT Photo



Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

2.2 Configuration Photographs (Continued)

SAR measurement Test Setup

EUT Photo



Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

2.3 System Verification

Prior to the assessment, the system was verified to the $\pm 10\%$ of the specifications by using the system validation kit. The validation was performed at 900 MHz. & 1800 MHz.

Validation kit	Targeted SAR _{1g} (mW/g)	Measured SAR _{1g} (mW/g)	Plot Number
D900V2	2.65	2.74	13
D1800V2	9.45	10.1	14

2.4 Evaluation Procedures

The SAR evaluation was performed with the following procedures:

- a. SAR was measured at a fixed location above the reference point and used as a reference value for the assessing the power drop.
- b. The SAR distribution at the exposed side of the flat Phantom was measured at a distance of 30 mm from the inner surface of the shell. The area covered the entire dimension of the head and the horizontal grid spacing was 20 mm x 20 mm. Based on this data, the area of the maximum absorption was determined by spline interpolation.
- c. Around this point, a volume of 32 mm x 32 mm x 34 mm was assessed by measuring 5 x 5 x 7 points. On the basis of this data set, the spatial peak SAR value was evaluated with the following procedure:
 - i) The data at the surface were extrapolated, since the center of the dipoles is 2.7 mm away from the tip of the probe and the distance between the surface and the lowest measurement point is 1.6 mm. The extrapolation was based on a least square algorithm. A polynomial of the fourth order was calculated through the points in Z-axes. This polynomial was then used to evaluate the points between the surface and the probe tip.
 - ii) The maximum interpolated value was searched with a straightforward algorithm. Around this maximum, the SAR values averaged over the spatial volumes (1g or 10g) were computed using the 3-D spline interpolation algorithm. The 3-D spline is composed of three one-dimensional splines with the "Not a knot" condition (in x, y and z directions). The volume was integrated with the trapezoidal algorithm. 1000 points (10 x 10 x 10) were interpolated to calculate the average.
 - iii) All neighboring volumes were evaluated until no neighboring volume with a higher average value was found.
- d. Re-measurements of the SAR value at the same location as in step a. above. If the value changed by more than 5 %, the evaluation was repeated.

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

2.5 Test Results

The results on the following page(s) were obtained when the device was tested in the condition described in this report. Detail measurement data and plots, which reveal information about the location of the maximum SAR with respect to the device, are reported in Appendix A.

Cardionet, Inc., Model No: CardioNet Monitor
 FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

Measurement Results

Trade Name:	CardioNet	Model No.:	CardioNet Monitor
Serial No.:	Not Labeled	Test Engineer:	Suresh Kondapalli

TEST CONDITIONS					
Ambient Temperature	23.5 °C	Relative Humidity	55 %		
Test Signal Source	Test Mode	Signal Modulation	CDMA		
Output Power Before SAR Test	See Page 8	Output Power After SAR Test	Within -0.32 to 0.12 dB		
Test Duration	20 Min. each test	Number of Battery Change	New battery for each scan		
Position - 15 mm from Phantom					
Frequency MHz	Operating Mode	Crest Factor	Measured SAR _{1g} (mW/g)	Position 15 mm From Phantom	Plot Number
824	CDMA	1	0.634	Front Side	1
824	CDMA	1	0.746	Back Side	2
835	CDMA	1	0.890	Front Side	3
835	CDMA	1	1.070	Back Side	4
849	CDMA	1	0.547	Front Side	5
849	CDMA	1	0.651	Back Side	6
1850	CDMA	1	0.132	Front Side	9
1850	CDMA	1	0.124	Back Side	10
1880	CDMA	1	0.116	Front Side	11
1880	CDMA	1	0.094	Back Side	12
1910	CDMA	1	0.102	Front Side	13
1910	CDMA	1	0.088	Back Side	14

D900V2					
Frequency MHz	Operating Mode	Crest Factor	Measured SAR _{1g} (mW/g)	Measured SAR _{10g} (mW/g)	Plot Number
900	CW	1	2.74	1.75	8

D1800V2					
Frequency MHz	Operating Mode	Crest Factor	Measured SAR _{1g} (mW/g)	Measured SAR _{10g} (mW/g)	Plot Number
1800	CW	1	10.9	6.19	15

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

Z-Scan					
Frequency MHz	Operating Mode	Crest Factor	Measured SAR _{1g} (mW/g)	Measured SAR _{10g} (mW/g)	Plot Number
835	CDMA	1	1.07	0.804	7

Note: a) Worst case data were reported
b) Duty cycle factor included in the measured SAR data
c) Uncertainty of the system is not included

Cardionet, Inc., Model No: CardioNet Monitor
 FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

3.0 TEST EQUIPMENT

3.1 Equipment List

The Specific Absorption Rate (SAR) tests were performed with the SPEAG model DASY 3 automated near-field scanning system, which is a package, optimized for dosimetric evaluation of mobile radios [3].

The following major equipment/components were used for the SAR evaluations:

SAR Measurement System			
EQUIPMENT	SPECIFICATIONS	S/N #	Last Cal. Data
Robot	Stäubi RX60L	597412-01	N/A
	Repeatability: ± 0.025 mm Accuracy: 0.806×10^{-3} degree Number of Axes: 6		
E-Field Probe	ER3DV4R	1576	2/27/02 *
	Dynamic Range: $5 \mu\text{W/g}$ to $>100 \text{ mW/g}$ Tip diameter: 6.8 mm Probe Linearity: $\pm 0.2 \text{ dB}$ (30 MHz to 3 GHz) Axial isotropy: $\pm 0.2 \text{ dB}$ Spherical isotropy: $\pm 0.4 \text{ dB}$ Length: 34.5 cm Distance between the probe tip and the dipole center: 2.7 mm Calibration: 835/900 MHz and 1800/1900 MHz for head & body simulating liquid		
Data Acquisition	DAE3	317	N/A
	Measurement Range: $1 \mu\text{V}$ to $>200 \text{ mV}$ Input offset Voltage: $< 1 \mu\text{V}$ (with auto zero) Input Resistance: 200 M		
Phantom	SAM Twin V4.0	TP-1243	QD000P40CA
	Complies with IEEE P1528-200x, draft 6.5 (see certificate in App. C) Type SAM Twin, Homogenous Shell Material: Fiberglass Thickness: $2 \pm 0.2 \text{ mm}$ Capacity: 20 liter Size of the flat section: approx. 320 x 230 mm		
Device holder	Non-conductive holder supplied with DASY3, dielectric constant less than 5.0	N/A	N/A
Simulated Tissue	Mixture	N/A	3/10/03
	Please see section 6.2 for details		
Power Meter	HP 8900D w/ 84811A sensor	3607U00673	08/27/02
	Frequency Range: 100kHz to 18 GHz Power Range: $300 \mu\text{W}$ to 3W		

* See Note on the next page.

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

Note:

At the time of the test, the calibration interval of the probe was longer than one year. To verify that its calibration remained within acceptable limits, the test results with the Probe Serial #1576 were compared with test results with the Probe Serial # 1577 on April 25, 2003, having last calibration date of February 5, 2003.

Another device was tested on April 22, 2003 at ITS using Probe #1576 and at another test Laboratory (CCS, Compliance Certification Services) using Probe #1577 on April 25, 2003. The comparison of the results indicates that SAR readings are within the overall measurement uncertainty requirements of IEEE Standard 1528-200X. The comparison of the results obtained with two probes is below:

SAR mW/g Measured with Probe # 1576 (ITS)	SAR mW/g Measured with Probe # 1577 (CCS)	Variance
0.326	0.280	+16.4%
0.183	0.197	-7.1%
0.245	0.239	+2.5%

Additionally, the system validation, performed with the Dipoles at the time of Cardionet Monitor testing was within 10% of the target values (in particular: +8.6% for 900 MHz, and +0.5% for 1800 MHz).

Based on this information, we can conclude that the conversion factors for Probe # 1576 are still within the tolerance ($\pm 8.9\%$) and the test results for Cardionet Monitor are valid.

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

3.2 Muscle Tissue Simulating Liquid

Muscle Ingredients Frequency (835 MHz)	
Sugar	41.76%
HEC	1.21%
Water	56.0%
Bactericide	0.27%
Salt	0.76%

The dielectric parameters were verified prior to assessment using the HP 85070A dielectric probe kit and the HP 8753C network Analyzer. The dielectric parameters were:

Frequency (MHz)	ϵ_r^*	σ^*(mho/m)	ρ^{**}(kg/m³)
835	53.2	0.98	1000

* Worst case uncertainty of the HP 85070A dielectric probe kit

** Worst-case assumption

Muscle Ingredients Frequency (1800 MHz)	
Water	52.90 %
Salt	0.181%
DGBE Diethylene Glycol	44.92%
ton X-100 (Polyethylene Glycol Mono) Ether	0.1 %

The dielectric parameters were verified prior to assessment using the HP 85070A dielectric probe kit and the HP 8753C network Analyzer. The dielectric parameters were:

Frequency (MHz)	ϵ_r^*	σ^*(mho/m)	ρ^{**}(kg/m³)
1800	55.8	1.49	1000

* Worst case uncertainty of the HP 85070A dielectric probe kit

** Worst-case assumption

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

3.3 E-Field Probe Calibration

The manufacturer in the TEM cell ifi 110 calibrated probes. To ensure consistency, a strict protocol was followed. The conversion factor (ConF) between this calibration and the measurement in the tissue simulation solution was performed by comparison with temperature measurement and computer simulations. Probe calibration factors are included in Appendix B.

Cardionet, Inc., Model No: CardioNet Monitor
 FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

3.4 Measurement Uncertainty

The Table below includes the uncertainty budget suggested by the IEEE Std 1528-200X and determined by SPEAG for the DASY3 measurement System.

The extended uncertainty (K=2) was assessed to be 27.8 %

Uncertainty Component	Tolerance (± %)	Probability Distribution	Divisor	c_i	Standard Uncertainty, (± %)	v_i^2 or v_{eff}
Measurement System						
Probe Calibration	4.7	Normal	1	1	4.7	Inf.
Axial Isotropy	4.7	Rectangular	$\sqrt{3}$	$(1-c_p)^{1/2}$	1.9	Inf.
Spherical Isotropy	9.6	Rectangular	$\sqrt{3}$	$\sqrt{c_p}$	3.9	Inf.
Boundary Effect	5.5	Rectangular	$\sqrt{3}$	1	3.2	Inf.
Linearity	4.7	Rectangular	$\sqrt{3}$	1	2.7	Inf.
System Detection Limits	1.0	Rectangular	$\sqrt{3}$	1	0.6	Inf.
Readout Electronics	1.0	Normal	1	1	1.0	Inf.
Response Time	0.8	Rectangular	$\sqrt{3}$	1	0.5	Inf.
Integration Time	1.4	Rectangular	$\sqrt{3}$	1	0.8	Inf.
RF Ambient Conditions	3.0	Rectangular	$\sqrt{3}$	1	1.7	Inf.
Probe Positioner Mechanical Tolerance	0.4	Rectangular	$\sqrt{3}$	1	0.2	Inf.
Probe Positioning with respect to Phantom Shell	2.9	Rectangular	$\sqrt{3}$	1	1.7	Inf.
Extrapolation, interpolation and Integration Algorithms for Max. SAR Evaluation	3.9	Rectangular	$\sqrt{3}$	1	2.3	Inf.
Test sample Related						
Test Sample Positioning	6.0	Normal	0.89	1	6.7	12
Device Holder Uncertainty	5.0	Normal	0.84	1	5.9	8
Output Power Variation - SAR drift measurement	7.0	Rectangular	$\sqrt{3}$	1	4.0	Inf.
Phantom and Tissue Parameters						
Phantom Uncertainty (shape and thickness tolerances)	4.0	Rectangular	$\sqrt{3}$	1	2.3	Inf.
Liquid Conductivity Target tolerance	5.0	Rectangular	$\sqrt{3}$	0.6	1.7	Inf.
Liquid Conductivity - measurement uncertainty	10.0	Rectangular	$\sqrt{3}$	0.6	3.5	Inf.
Liquid Permittivity Target tolerance	5.0	Rectangular	$\sqrt{3}$	0.6	1.7	Inf.
Liquid Permittivity - measurement uncertainty	5.0	Rectangular	$\sqrt{3}$	0.6	1.7	Inf.
Combined Standard Uncertainty					13.9	
Expanded Uncertainty (95% CONFIDENCE INTERVAL)					27.8	

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

Notes.

1. The Divisor is a function of the probability distribution and degrees of freedom (v_i and v_{eff}). See NIST Technical Note TN1297, NIS 81 and NIS 3003.
2. c_i is the sensitivity coefficient that should be applied to convert the variability of the uncertainty component into a variability of SAR.

3.5 Measurement Tractability

All measurements described in this report are traceable to National Institute of Standards and Technology (NIST) standards or appropriate national standards.

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

4.0 WARNING LABEL INFORMATION - USA

See users manual.

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

5.0 REFERENCES

- [1] ANSI, *ANSI/IEEE C95.1-1991: IEEE Standard for Safety Levels with Respect to Human Exposure to Radio Frequency Electromagnetic Fields, 3kHz to 300 GHz*, The Institute of electrical and Electronics Engineers, Inc., New York, NY CardioNet Monitor7, 1992
- [2] Federal Communications Commission, “Evaluating Compliance with FCC Guidelines for Human Exposure to Radiofrequency Electromagnetic Fields”, OET Bulletin 65, FCC, Washington, D.C. 20554, 1997
- [3] Thomas Schmid, Oliver Egger, and Niels Kuster, “Automated E-field scanning system for dosimetric assessments”, *IEEE Transaction on Microwave Theory and Techniques*, vol. 44, pp. 105-113, Jan. 1996.
- [4] Niels Kuster, Ralph Kastle, and Thomas Schmid, “Dosimetric evaluation of mobile communications equipment with know precision”, *IEICE Transactions on Communications*, vol. E80-B, no. 5, pp.645-652, May 1997.
- [5] NIS81, NAMAS, “The treatment of uncertainty in EMC measurement”, Tech. Rep., NAMAS Executive, National Physical Laboratory, Teddinton, Middlesex, England, 1994.
- [6] Barry N. Taylor and Chris E. Kuyatt, “Guidelines for evaluating and expressing the uncertainty of NIST measurement results”, Tech. Rep., National Institute of Standards and Technology, 1994.

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

6.0 DOCUMENT HISTORY

Revision/ Job Number	Writer Initials	Date	Change
1.0 /3038816	SS	March 25, 2003	Original document

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

APPENDIX A - SAR Evaluation Data

Please note that the graphical visualization of the phone position onto the SAR distribution gives only limited information on the current distribution of the device, since the curvature of the head results in graphical distortion. Full information can only be obtained either by H-field scans in free space or SAR evaluation with a flat phantom.

Power drift is the measurement of power drift of the device over one complete SAR scan.

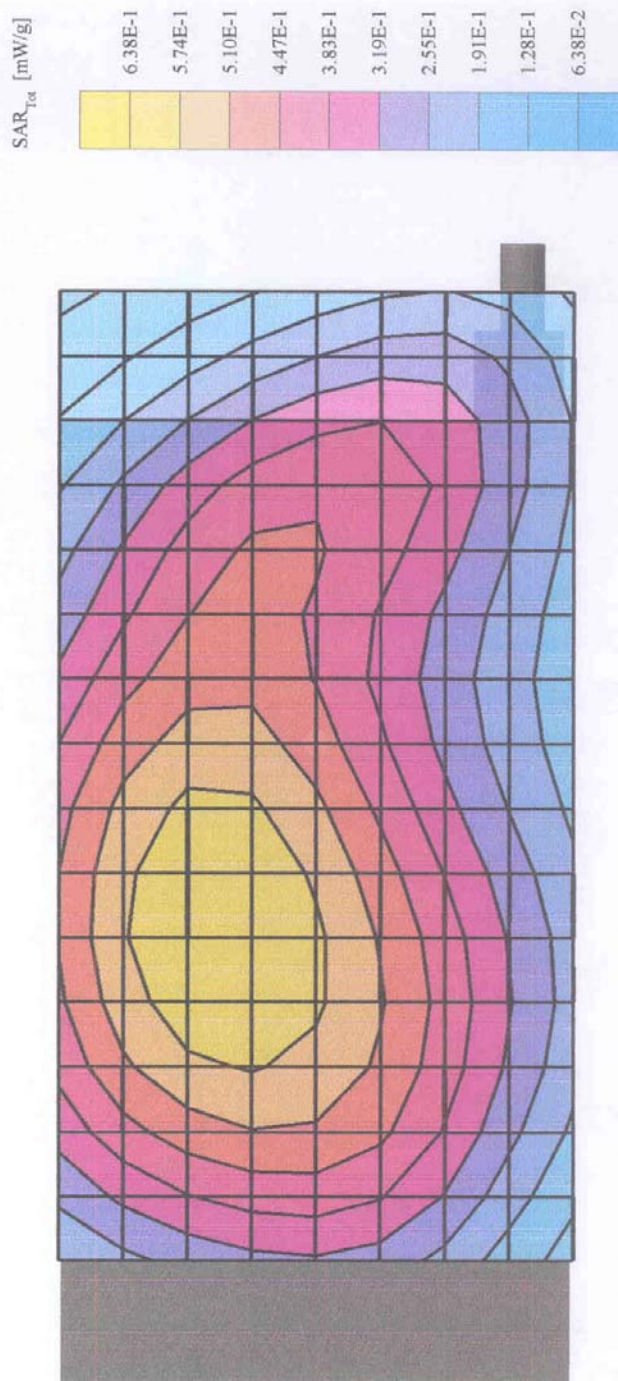
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

03/10/03

CardioNet Monitor CDMA

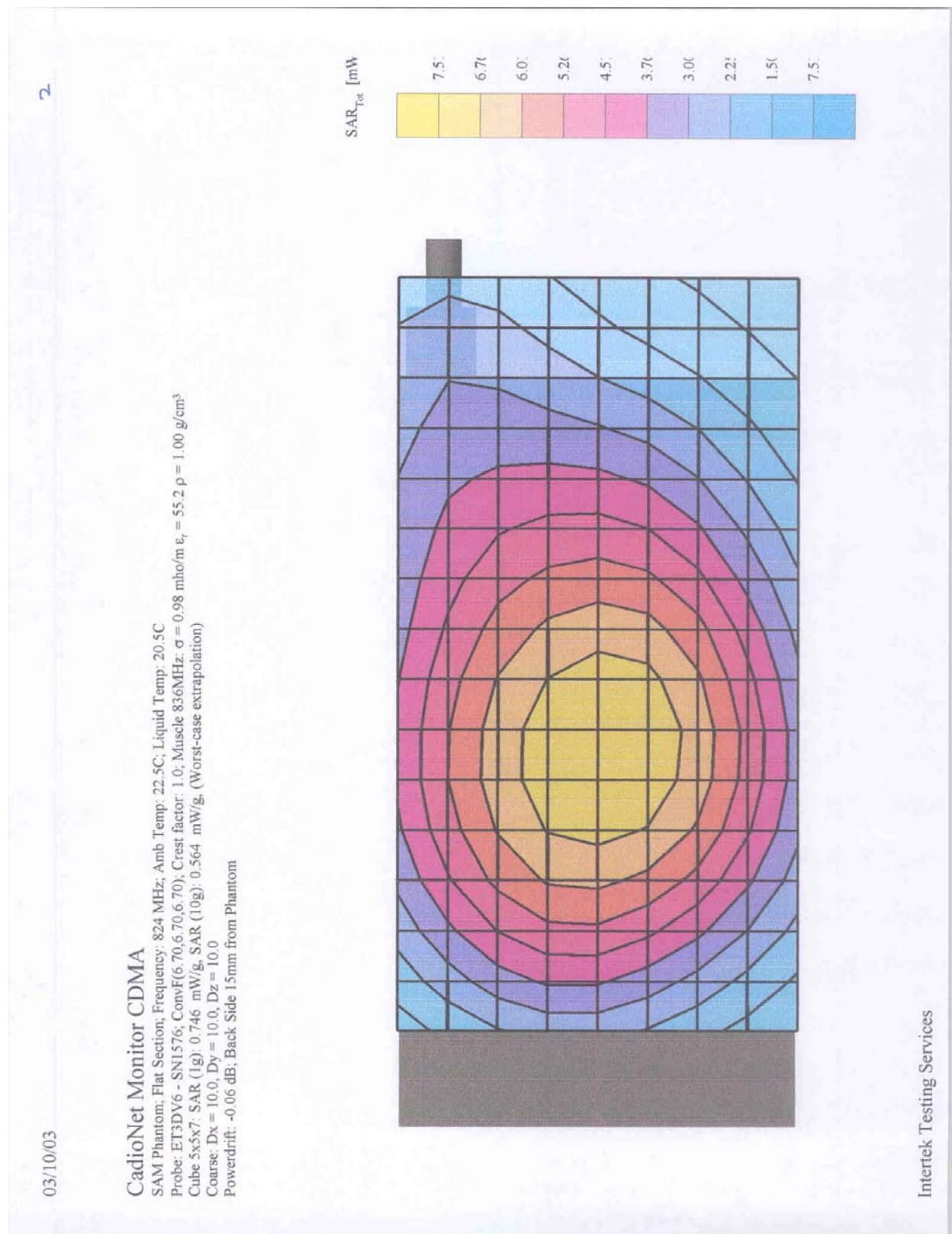
SAM Phantom: Flat Section; Frequency: 824 MHz; Amb Temp: 22.5°C; Liquid Temp: 20.5°C
Probe: ET3DV6 - SN1576; ConvF(6.70,6.70,6.70); Crest factor: 1.0; Muscle 836MHz: $\sigma = 0.98$ mho/m $\epsilon_r = 55.2$ $\rho = 1.00$ g/cm³
Cube 5x5x7: SAR (1g): 0.634 mW/g, SAR (10g): 0.477 mW/g (Worst-case extrapolation)
Coarse: Dx = 10.0, Dy = 10.0, Dz = 10.0
Powerdrift: -0.01 dB; Front Side 15mm from Phantom



Intertek Testing Services

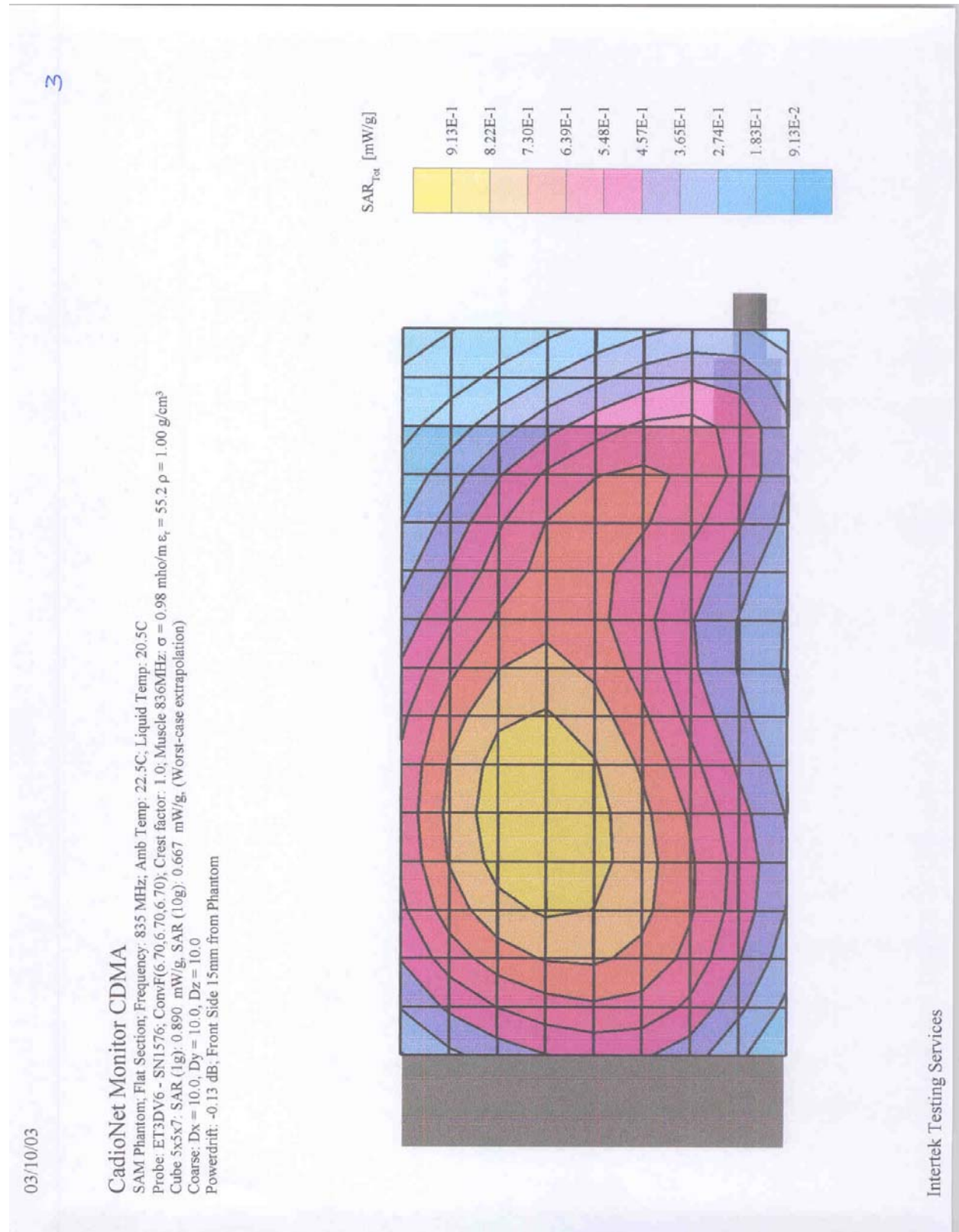
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



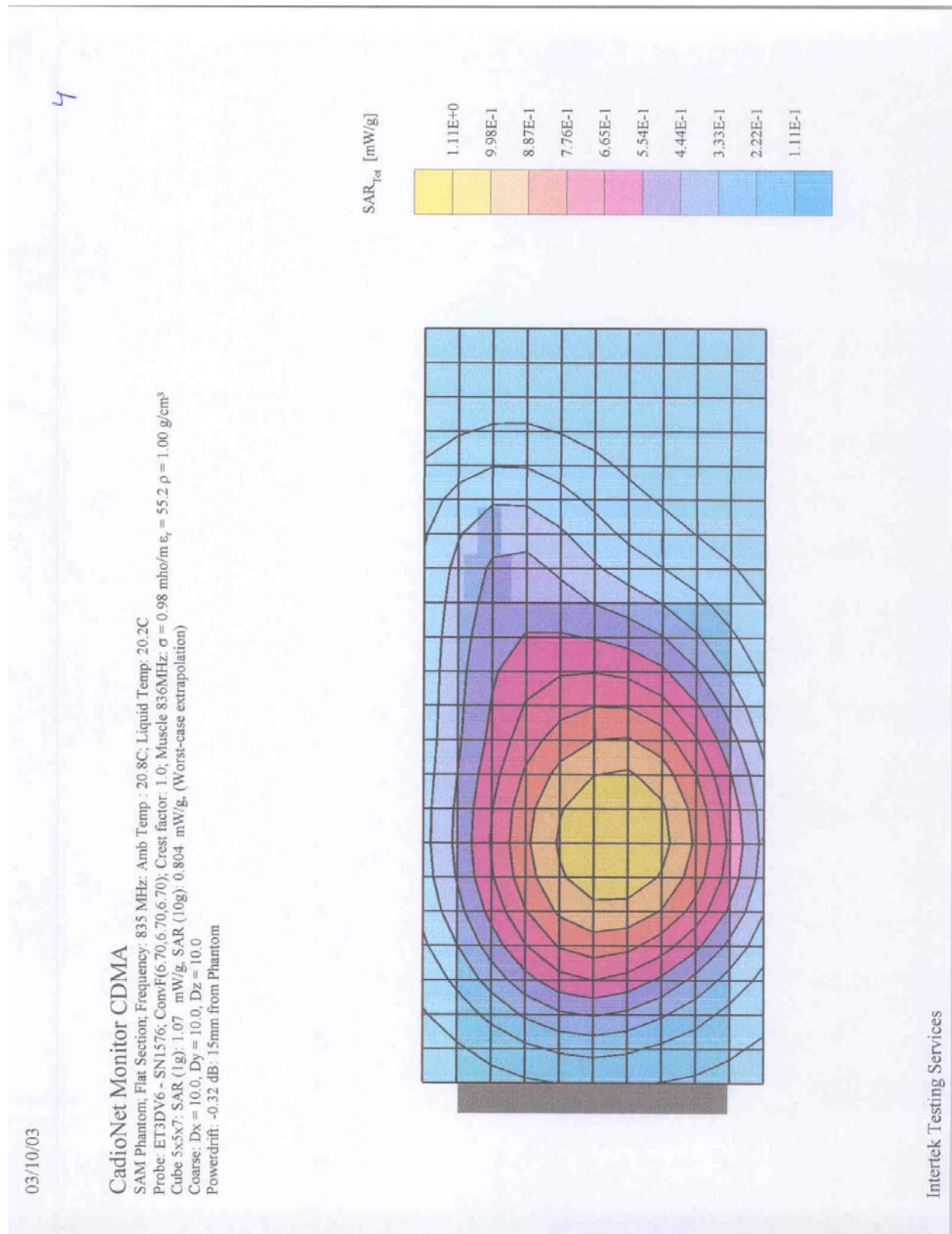
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



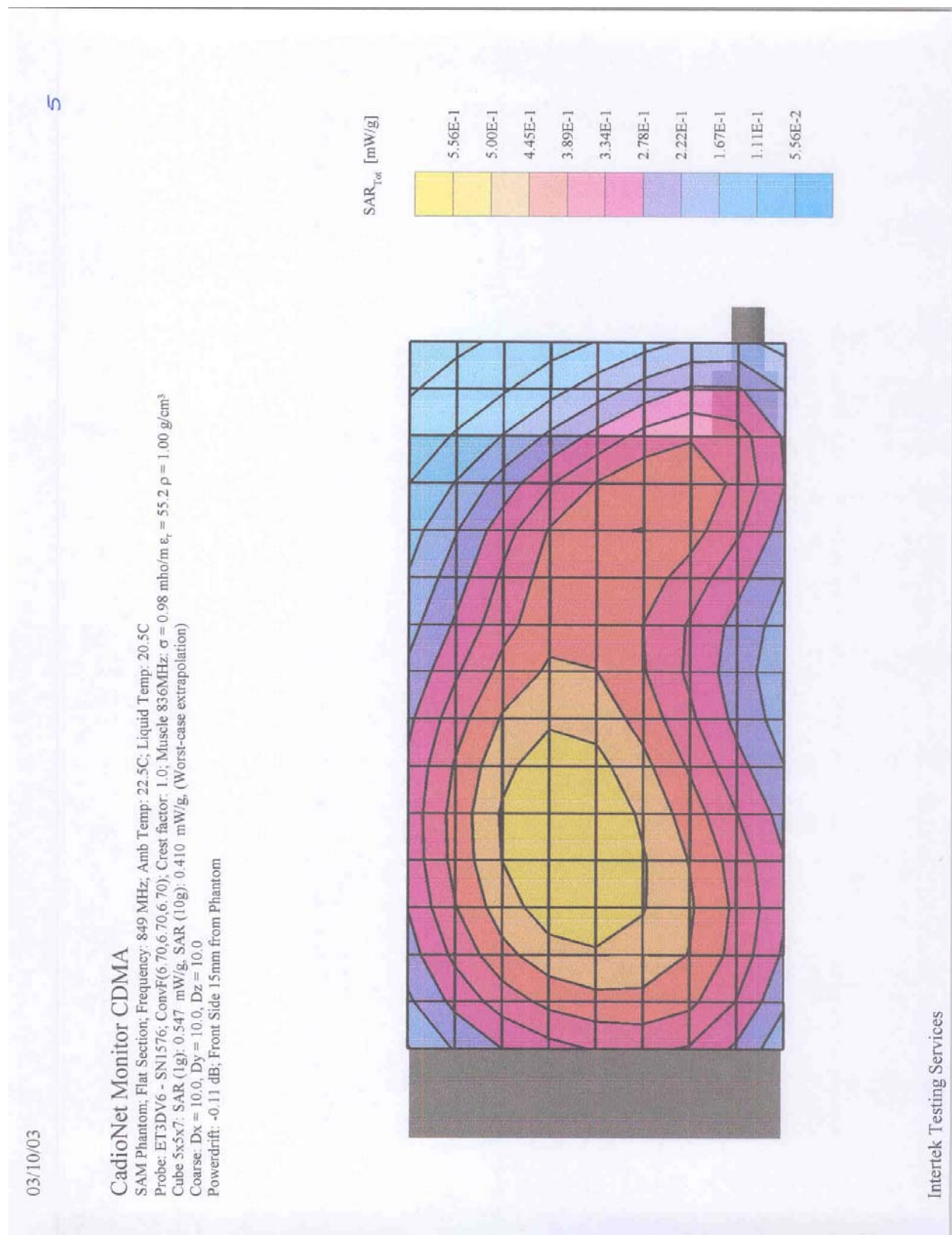
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



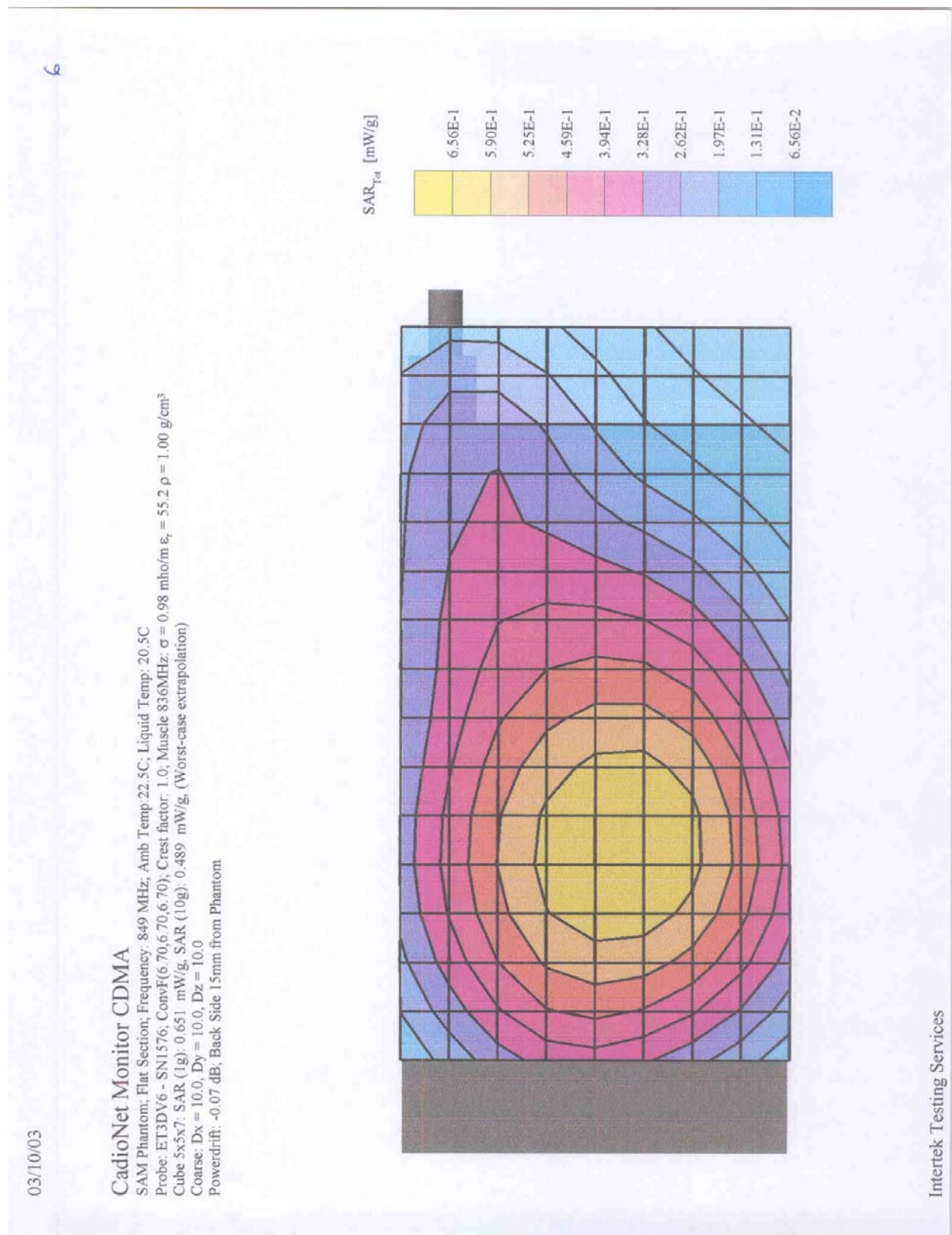
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



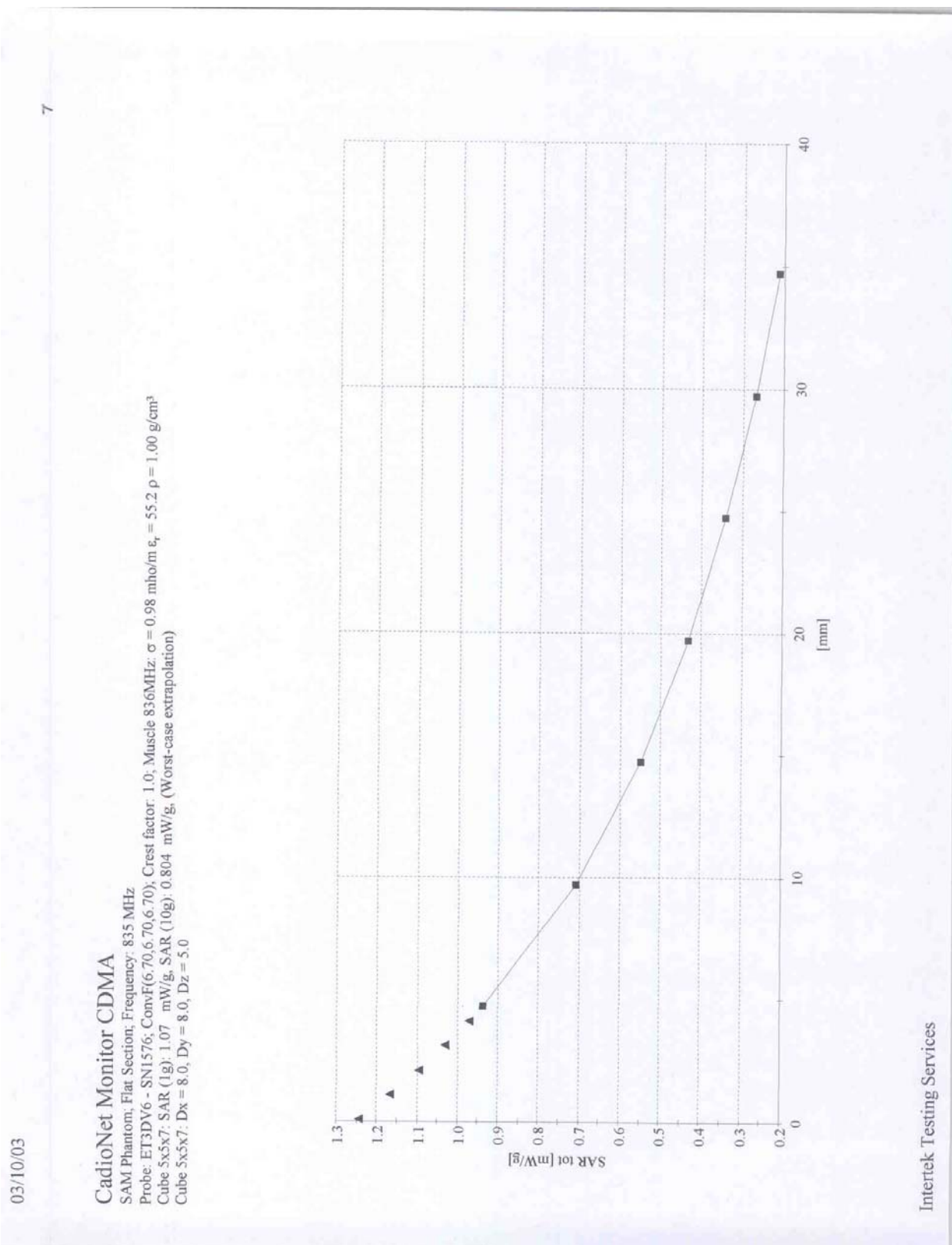
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



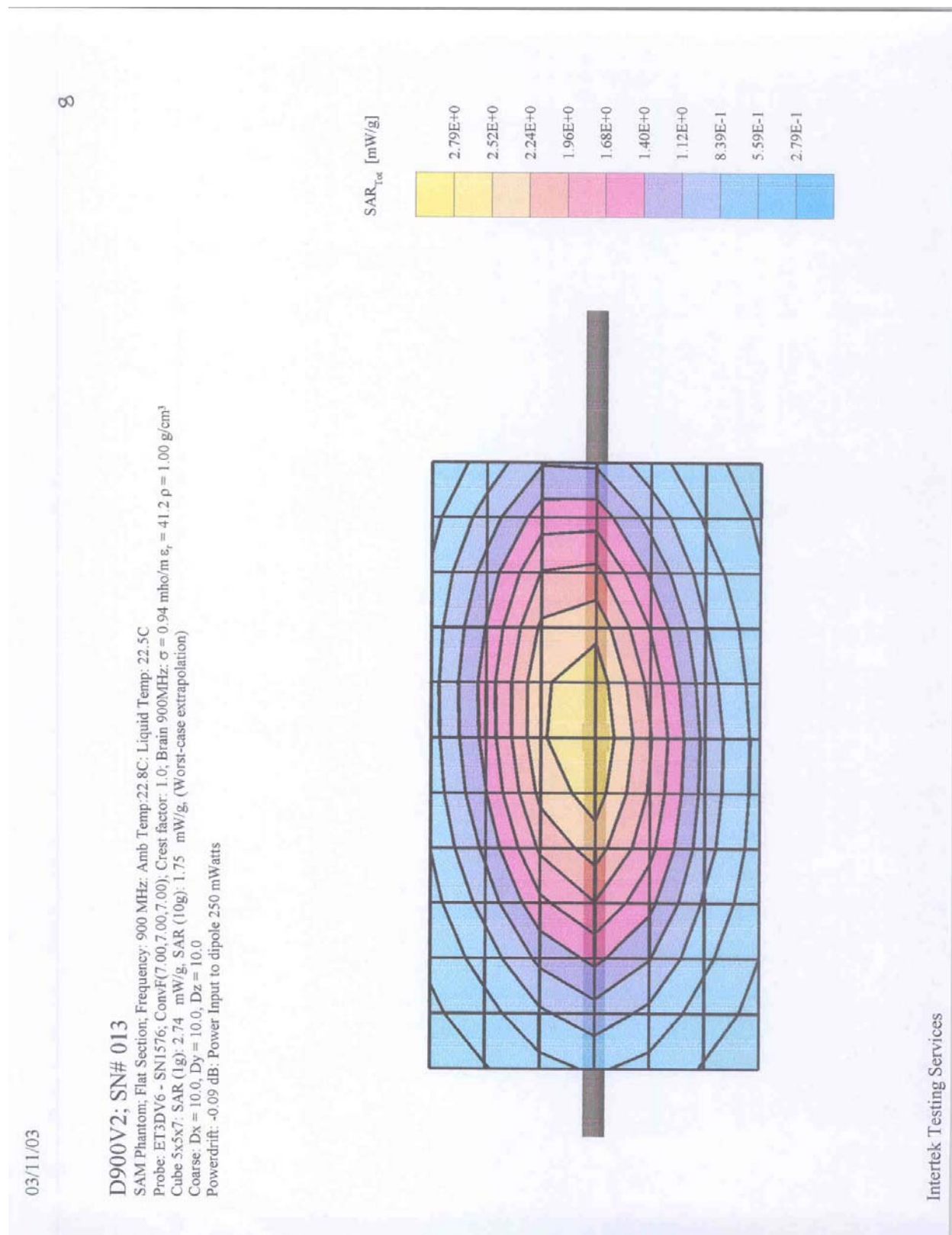
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



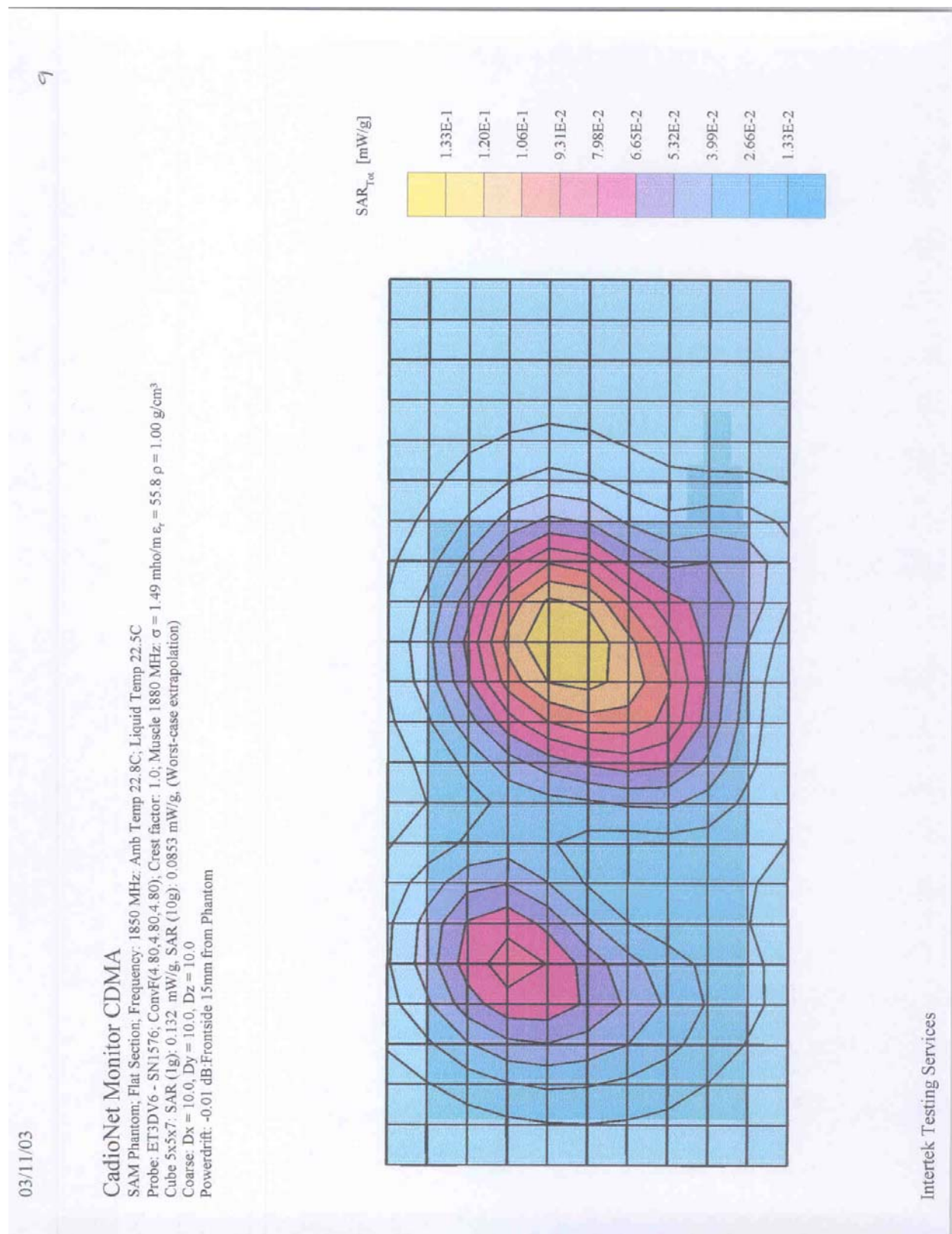
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



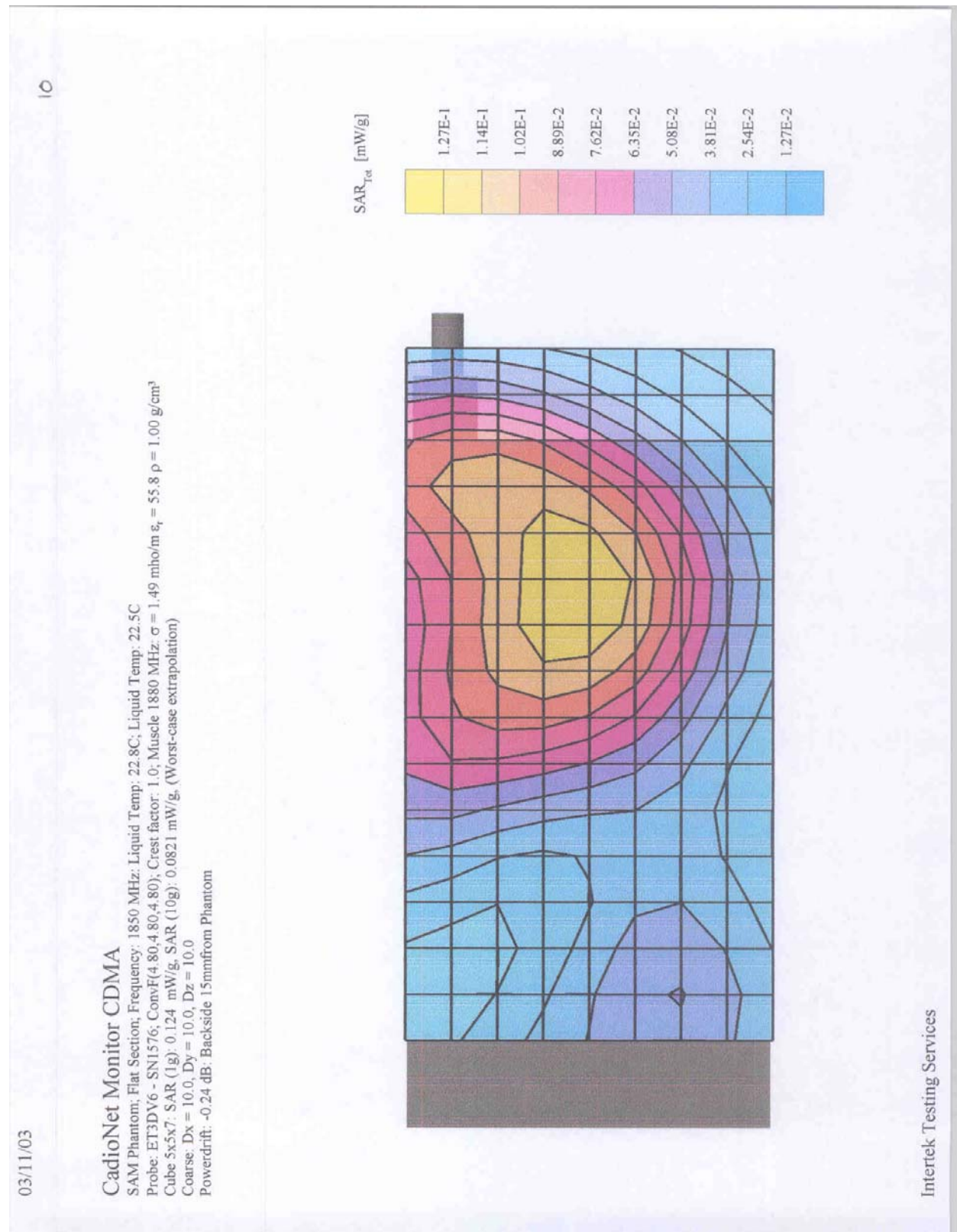
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



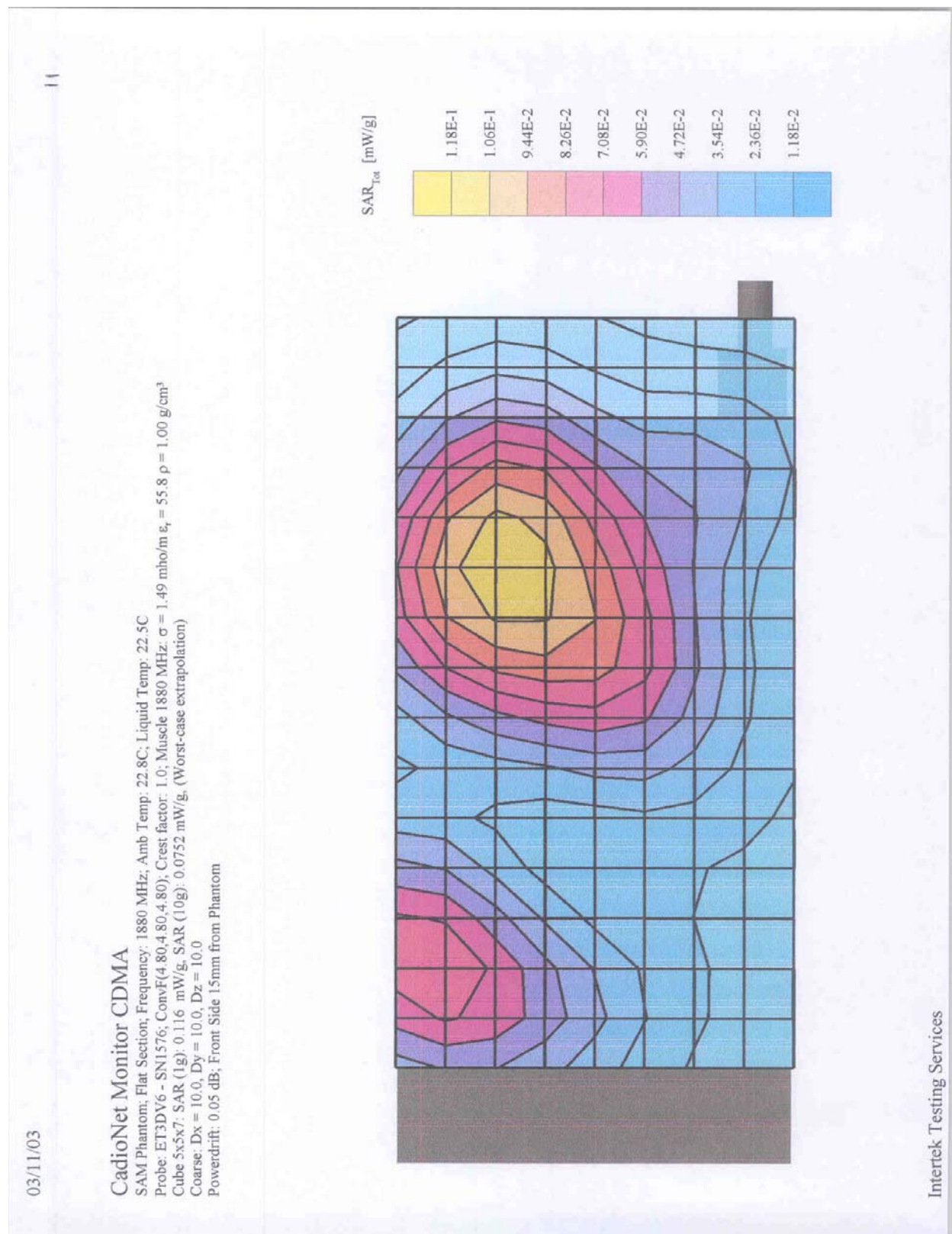
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



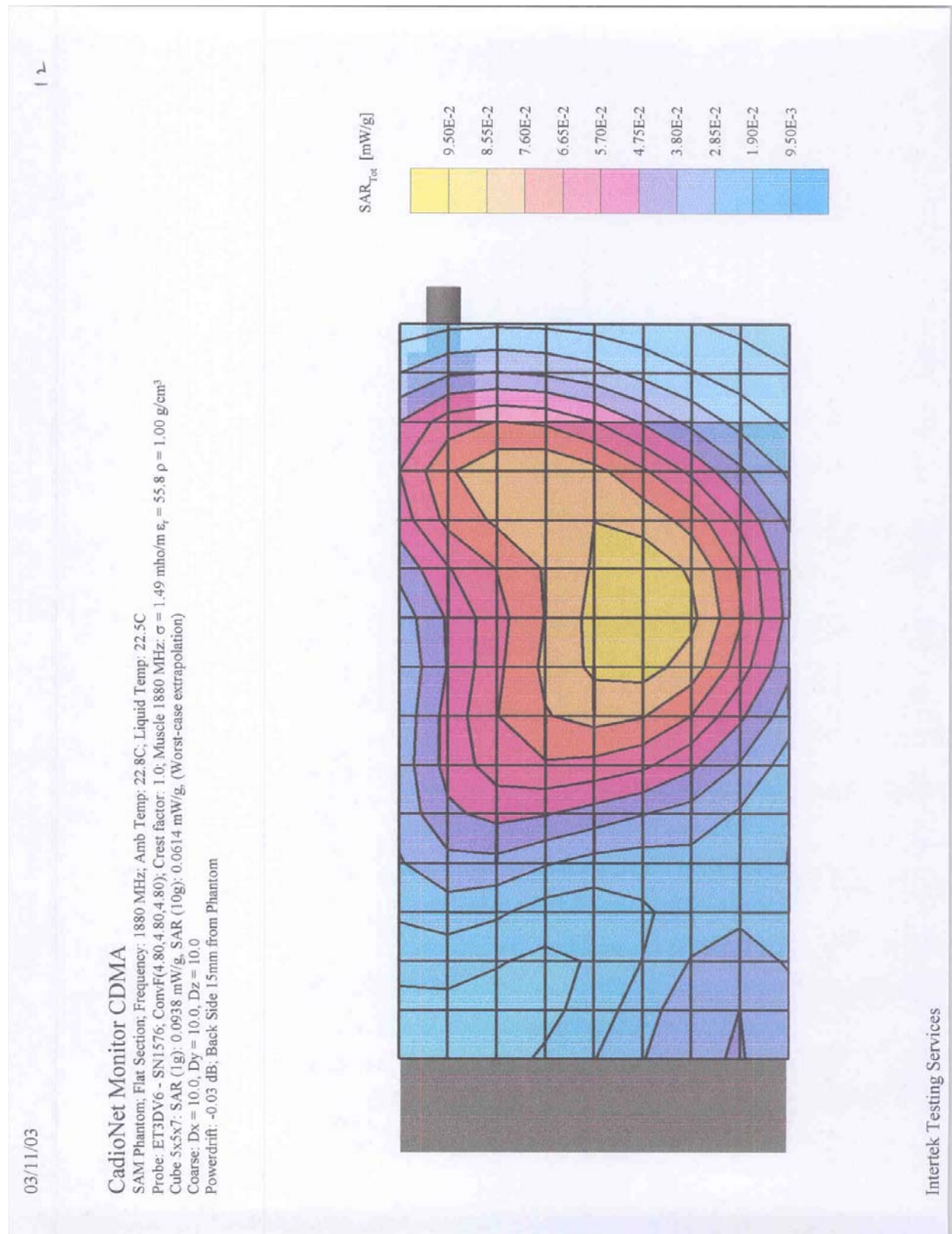
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



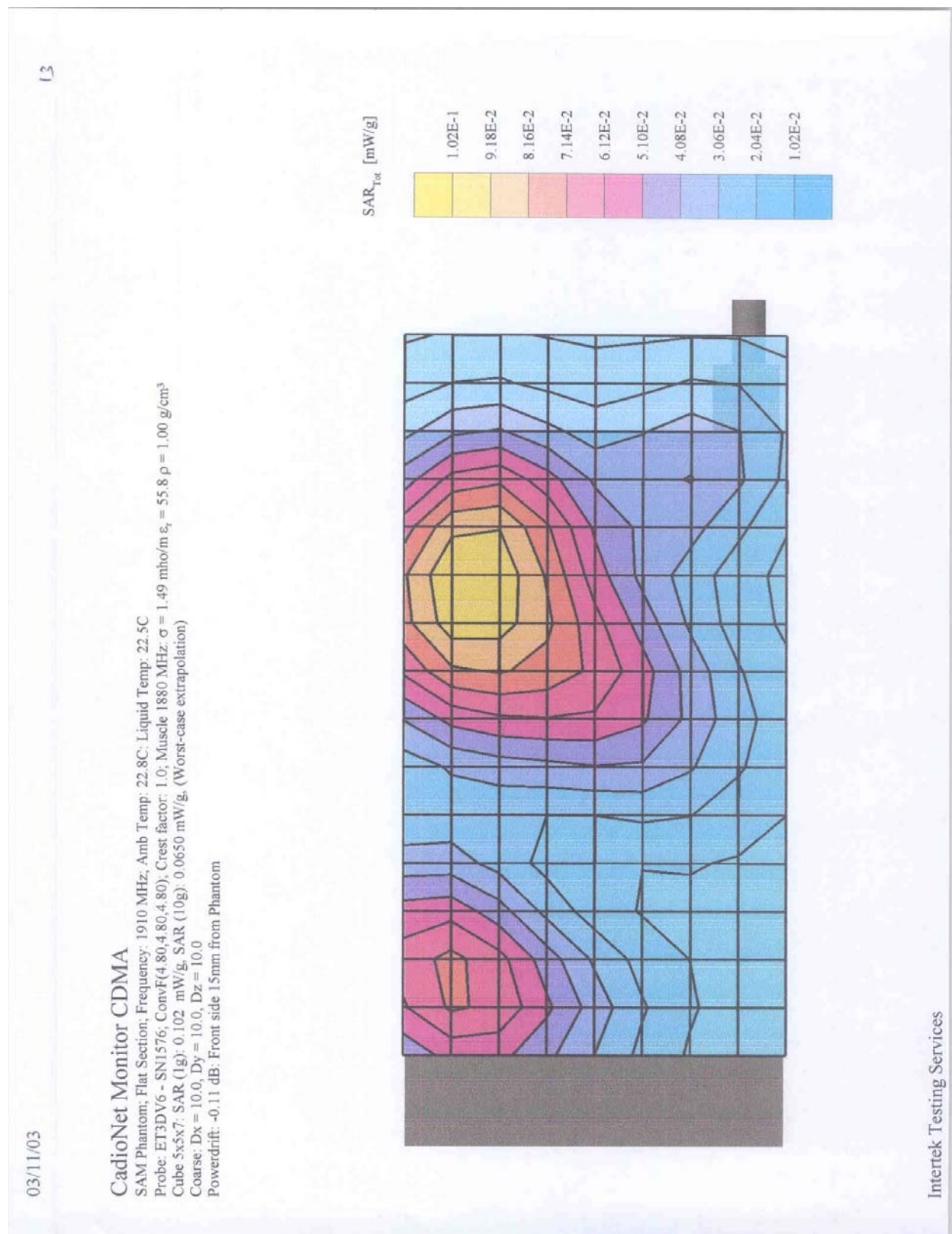
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



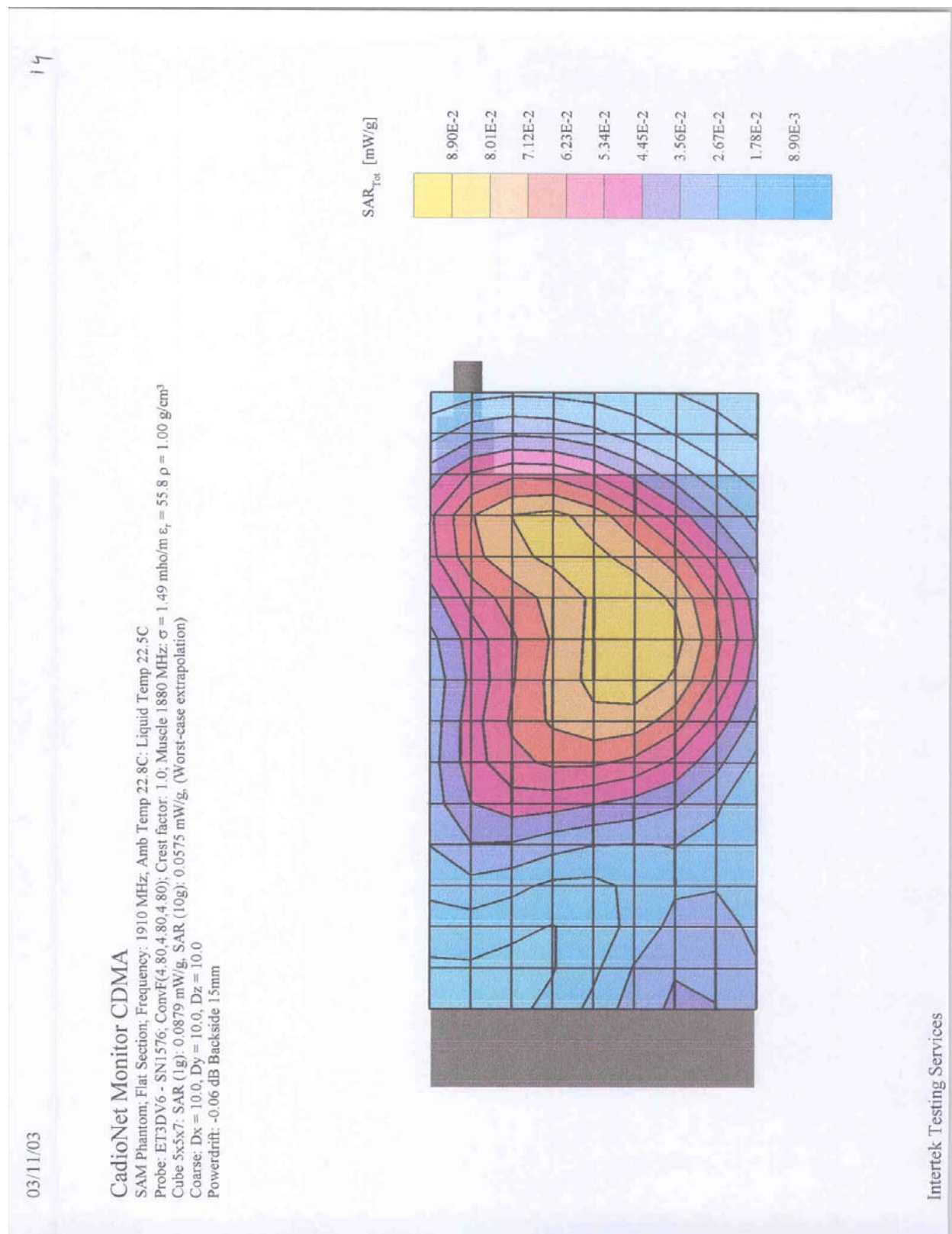
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



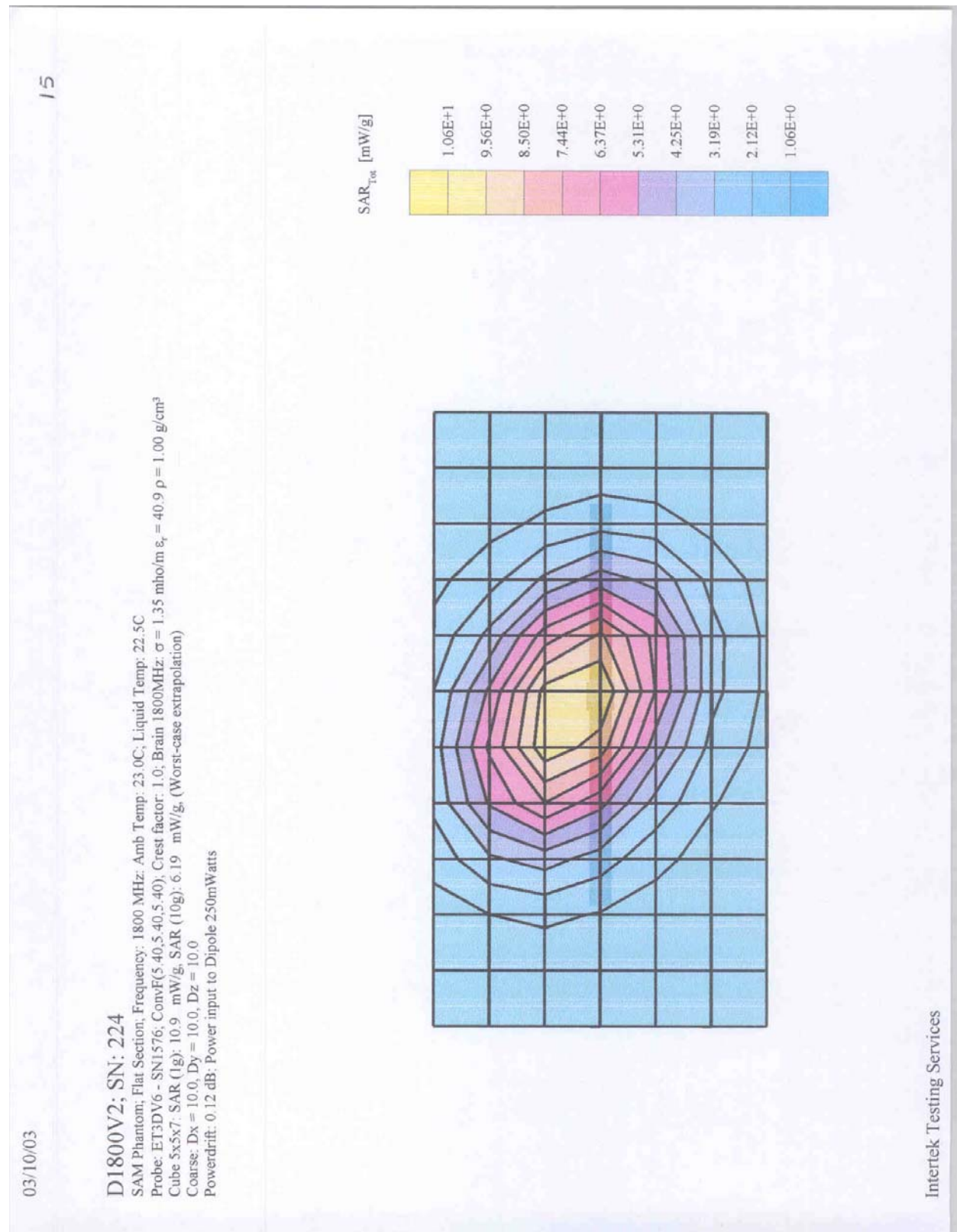
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003



Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

APPENDIX B - E-Field Probe Calibration Data

See attached.

Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

**Schmid & Partner
Engineering AG**

Zeughausstrasse 43, 8004 Zurich, Switzerland, Phone +41 1 245 97 00, Fax +41 1 245 97 79

Calibration Certificate

Dosimetric E-Field Probe

Type:

ET3DV6

Serial Number:

1576

Place of Calibration:

Zurich

Date of Calibration:

February 27, 2002

Calibration Interval:

12 months

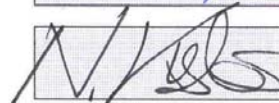
Schmid & Partner Engineering AG hereby certifies, that this device has been calibrated on the date indicated above. The calibration was performed in accordance with specifications and procedures of Schmid & Partner Engineering AG.

Wherever applicable, the standards used in the calibration process are traceable to international standards. In all other cases the standards of the Laboratory for EMF and Microwave Electronics at the Swiss Federal Institute of Technology (ETH) in Zurich, Switzerland have been applied.

Calibrated by:



Approved by:



**Schmid & Partner
Engineering AG**

Zeughausstrasse 43, 8004 Zurich, Switzerland, Telephone +41 1 245 97 00, Fax +41 1 245 97 79

Probe ET3DV6

SN:1576

Manufactured:	April 6, 2001
Last calibration:	April 20, 2001
Recalibrated:	February 27, 2002

Calibrated for System DASY3

ET3DV6 SN:1576

February 27, 2002

DASY3 - Parameters of Probe: ET3DV6 SN:1576

Sensitivity in Free Space

NormX	1.77 $\mu\text{V}/(\text{V}/\text{m})^2$
NormY	1.81 $\mu\text{V}/(\text{V}/\text{m})^2$
NormZ	1.76 $\mu\text{V}/(\text{V}/\text{m})^2$

Diode Compression

DCP X	98	mV
DCP Y	98	mV
DCP Z	98	mV

Sensitivity in Tissue Simulating Liquid

Head	900 MHz	$\epsilon_r = 41.5 \pm 5\%$	$\sigma = 0.97 \pm 5\%$ mho/m
Head	835 MHz	$\epsilon_r = 41.5 \pm 5\%$	$\sigma = 0.90 \pm 5\%$ mho/m
ConvF X	7.0 $\pm 9.5\%$ (k=2)	Boundary effect:	
ConvF Y	7.0 $\pm 9.5\%$ (k=2)	Alpha	0.30
ConvF Z	7.0 $\pm 9.5\%$ (k=2)	Depth	2.51
Head	1800 MHz	$\epsilon_r = 40.0 \pm 5\%$	$\sigma = 1.40 \pm 5\%$ mho/m
Head	1900 MHz	$\epsilon_r = 40.0 \pm 5\%$	$\sigma = 1.40 \pm 5\%$ mho/m
ConvF X	5.4 $\pm 9.5\%$ (k=2)	Boundary effect:	
ConvF Y	5.4 $\pm 9.5\%$ (k=2)	Alpha	0.45
ConvF Z	5.4 $\pm 9.5\%$ (k=2)	Depth	2.30

Boundary Effect

Head	900 MHz	Typical SAR gradient: 5 % per mm		
Probe Tip to Boundary		1 mm	2 mm	
SAR _{be} [%]	Without Correction Algorithm	7.6	4.3	
SAR _{be} [%]	With Correction Algorithm	0.3	0.5	
Head	1800 MHz	Typical SAR gradient: 10 % per mm		
Probe Tip to Boundary		1 mm	2 mm	
SAR _{be} [%]	Without Correction Algorithm	9.7	6.6	
SAR _{be} [%]	With Correction Algorithm	0.2	0.3	

Sensor Offset

Probe Tip to Sensor Center	2.7	mm
Optical Surface Detection	1.9 ± 0.2	mm

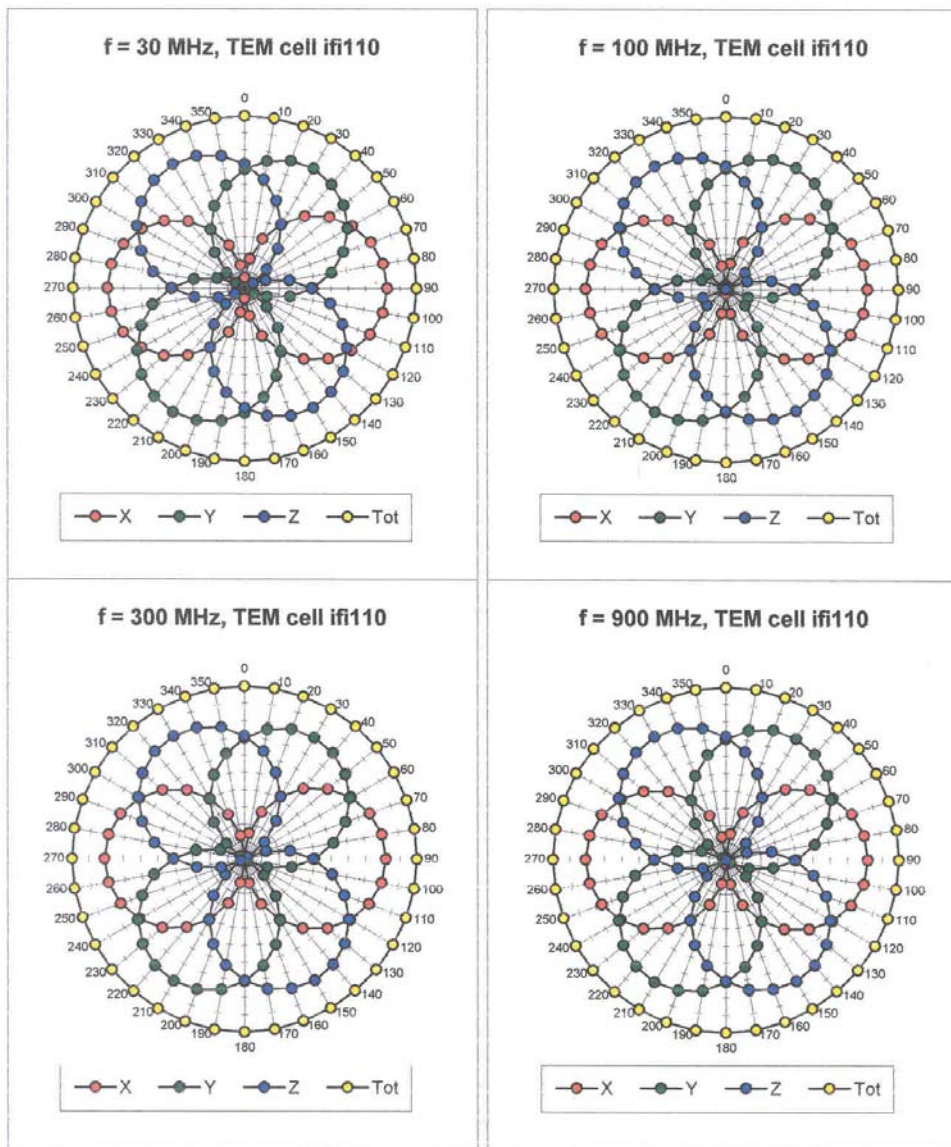
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

ET3DV6 SN:1576

February 27, 2002

Receiving Pattern (ϕ), $\theta = 0^\circ$

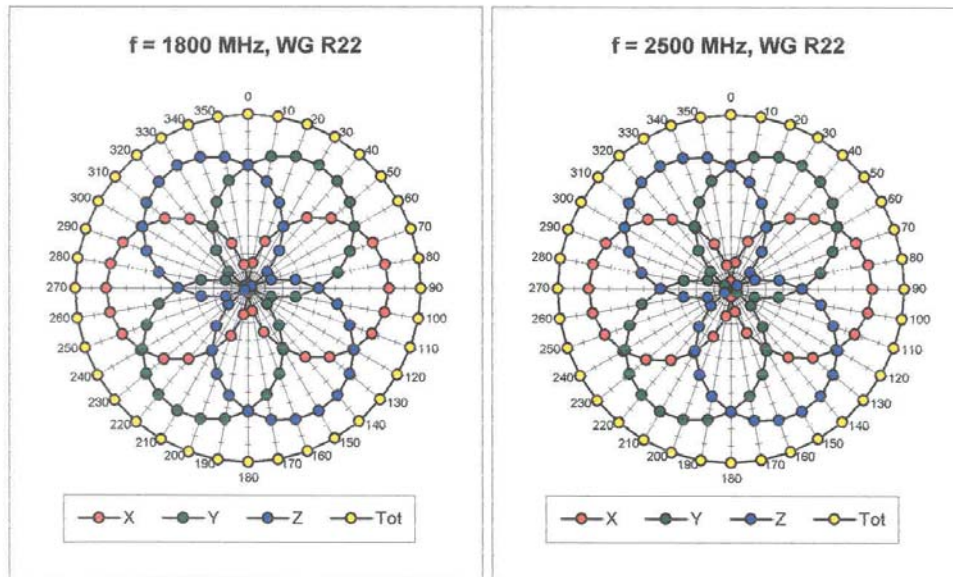


Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

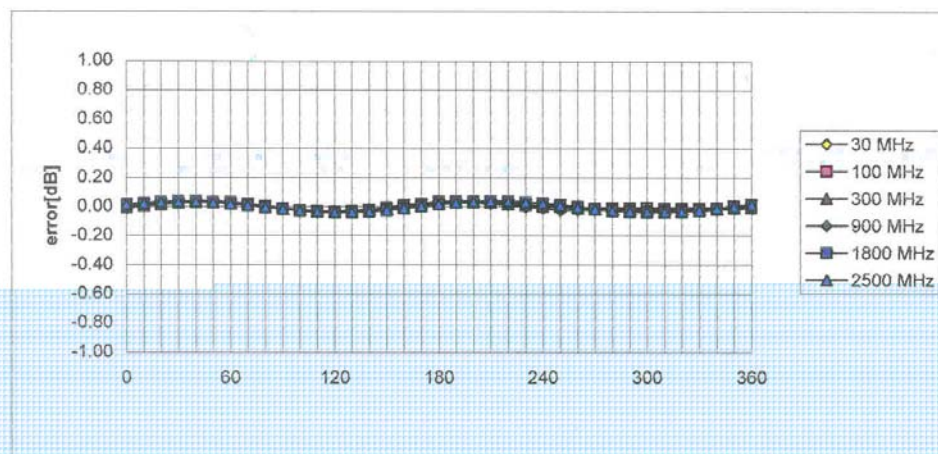
Date of Test: March 10 to 11, 2003

ET3DV6 SN:1576

February 27, 2002



Isotropy Error (ϕ), $\theta = 0^\circ$



Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

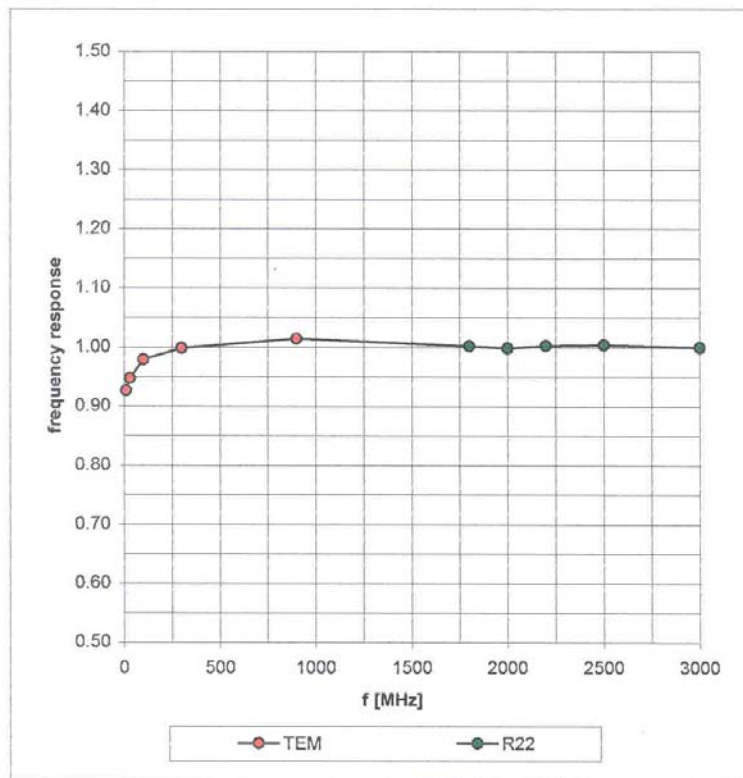
Date of Test: March 10 to 11, 2003

ET3DV6 SN:1576

February 27, 2002

Frequency Response of E-Field

(TEM-Cell:ifi110, Waveguide R22)



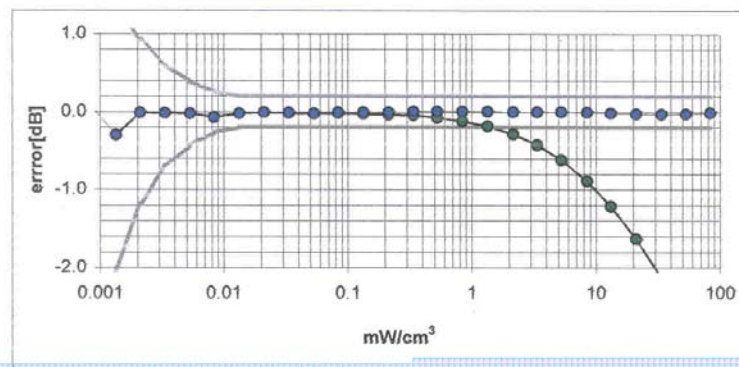
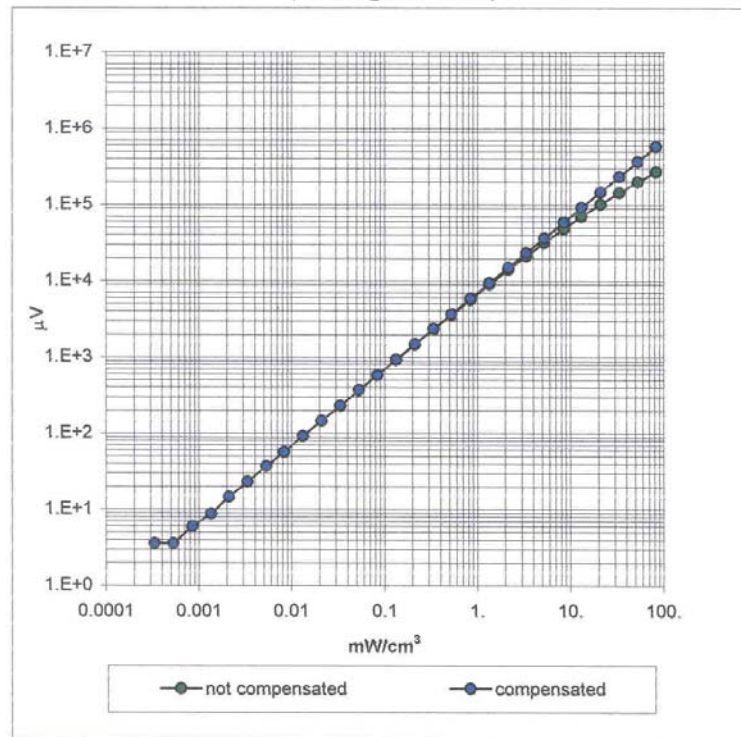
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

ET3DV6 SN:1576

February 27, 2002

Dynamic Range $f(\text{SAR}_{\text{brain}})$ (Waveguide R22)



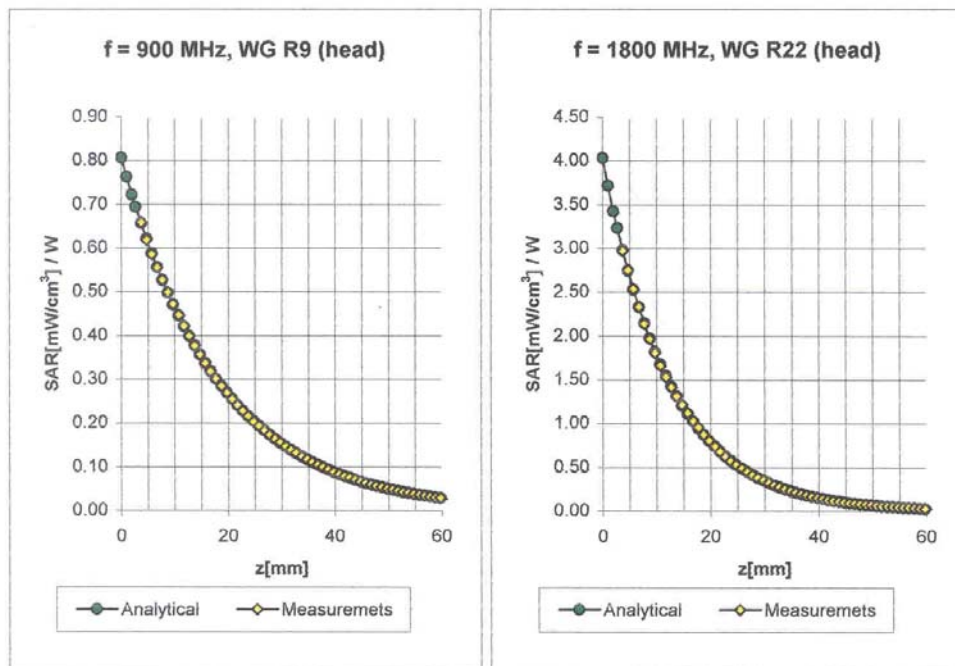
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

ET3DV6 SN:1576

February 27, 2002

Conversion Factor Assessment



Head	900 MHz	$\epsilon_r = 41.5 \pm 5\%$	$\sigma = 0.97 \pm 5\%$ mho/m
Head	835 MHz	$\epsilon_r = 41.5 \pm 5\%$	$\sigma = 0.90 \pm 5\%$ mho/m
	ConvF X	$7.0 \pm 9.5\%$ (k=2)	Boundary effect:
	ConvF Y	$7.0 \pm 9.5\%$ (k=2)	Alpha 0.30
	ConvF Z	$7.0 \pm 9.5\%$ (k=2)	Depth 2.51
Head	1800 MHz	$\epsilon_r = 40.0 \pm 5\%$	$\sigma = 1.40 \pm 5\%$ mho/m
Head	1900 MHz	$\epsilon_r = 40.0 \pm 5\%$	$\sigma = 1.40 \pm 5\%$ mho/m
	ConvF X	$5.4 \pm 9.5\%$ (k=2)	Boundary effect:
	ConvF Y	$5.4 \pm 9.5\%$ (k=2)	Alpha 0.45
	ConvF Z	$5.4 \pm 9.5\%$ (k=2)	Depth 2.30

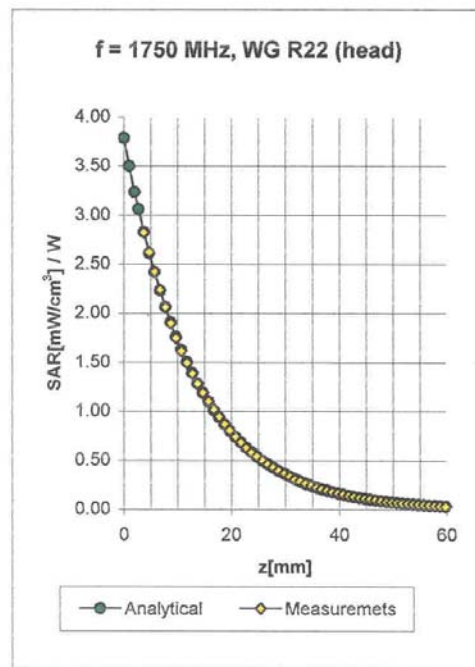
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

ET3DV6 SN:1576

February 27, 2002

Conversion Factor Assessment



Head	1750 MHz	$\epsilon_r = 40.0 \pm 5\%$	$\sigma = 1.40 \pm 5\% \text{ mho/m}$
ConvF X	5.4 $\pm 8.9\%$ (k=2)	Boundary effect:	
ConvF Y	5.4 $\pm 8.9\%$ (k=2)	Alpha	0.45
ConvF Z	5.4 $\pm 8.9\%$ (k=2)	Depth	2.27

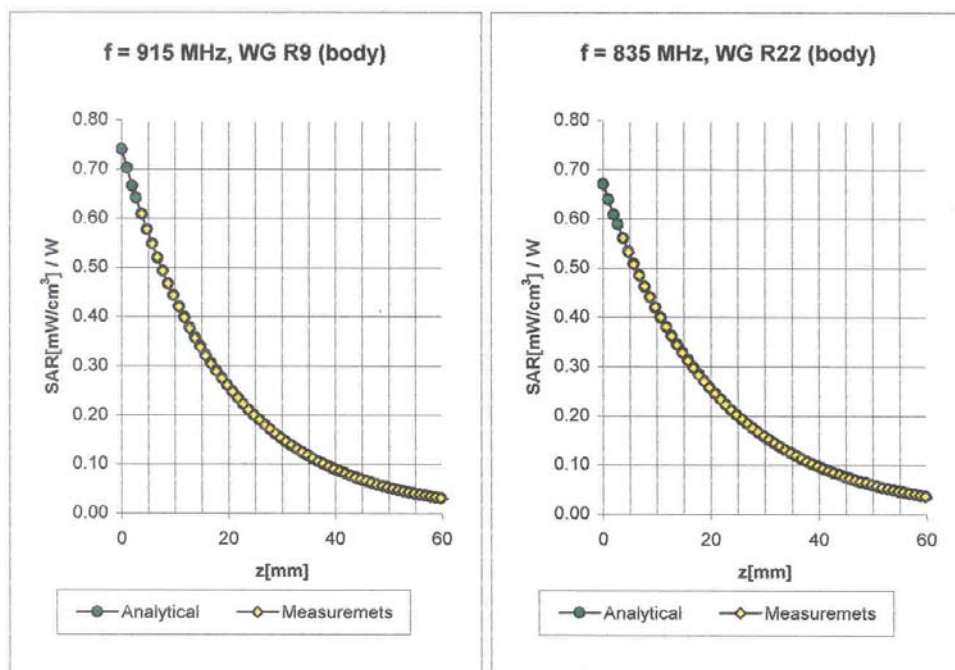
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

ET3DV6 SN:1576

February 27, 2002

Conversion Factor Assessment



Body	915 MHz	$\epsilon_r = 55.0 \pm 5\%$	$\sigma = 1.06 \pm 5\% \text{ mho/m}$
ConvF X	6.7 $\pm 8.9\%$ (k=2)	Boundary effect:	
ConvF Y	6.7 $\pm 8.9\%$ (k=2)	Alpha	0.45
ConvF Z	6.7 $\pm 8.9\%$ (k=2)	Depth	2.01
Body	835 MHz	$\epsilon_r = 55.2 \pm 5\%$	$\sigma = 0.97 \pm 5\% \text{ mho/m}$
ConvF X	6.7 $\pm 8.9\%$ (k=2)	Boundary effect:	
ConvF Y	6.7 $\pm 8.9\%$ (k=2)	Alpha	0.34
ConvF Z	6.7 $\pm 8.9\%$ (k=2)	Depth	2.37

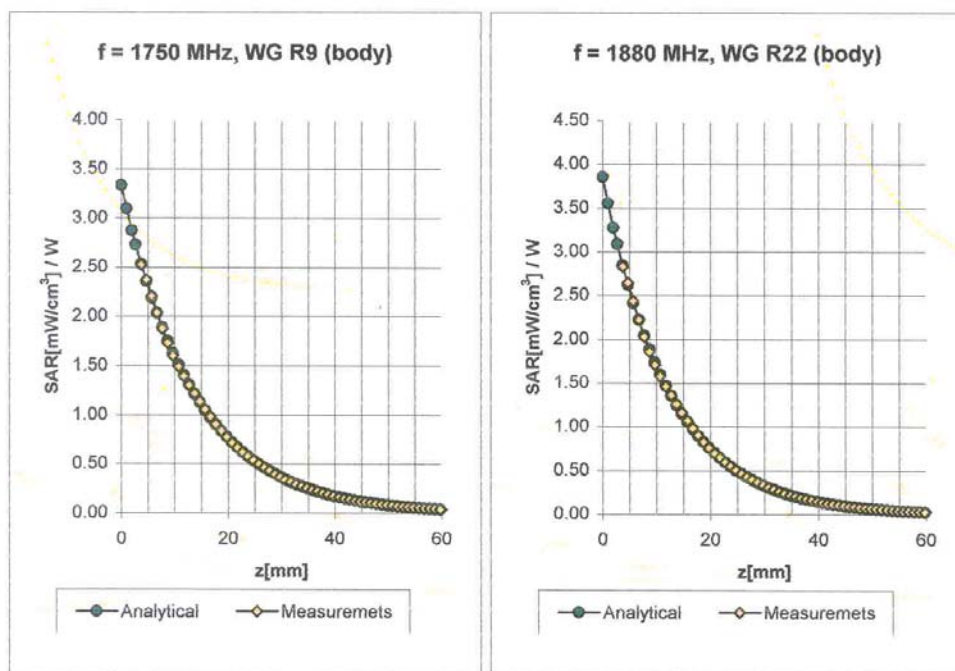
Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

ET3DV6 SN:1576

February 27, 2002

Conversion Factor Assessment



Body 1750 MHz $\epsilon_r = 53.3 \pm 5\%$ $\sigma = 1.52 \pm 5\% \text{ mho/m}$

ConvF X	5.1 $\pm 8.9\%$ (k=2)	Boundary effect:
ConvF Y	5.1 $\pm 8.9\%$ (k=2)	Alpha 0.51
ConvF Z	5.1 $\pm 8.9\%$ (k=2)	Depth 2.31

Body 1880 MHz $\epsilon_r = 53.3 \pm 5\%$ $\sigma = 1.52 \pm 5\% \text{ mho/m}$

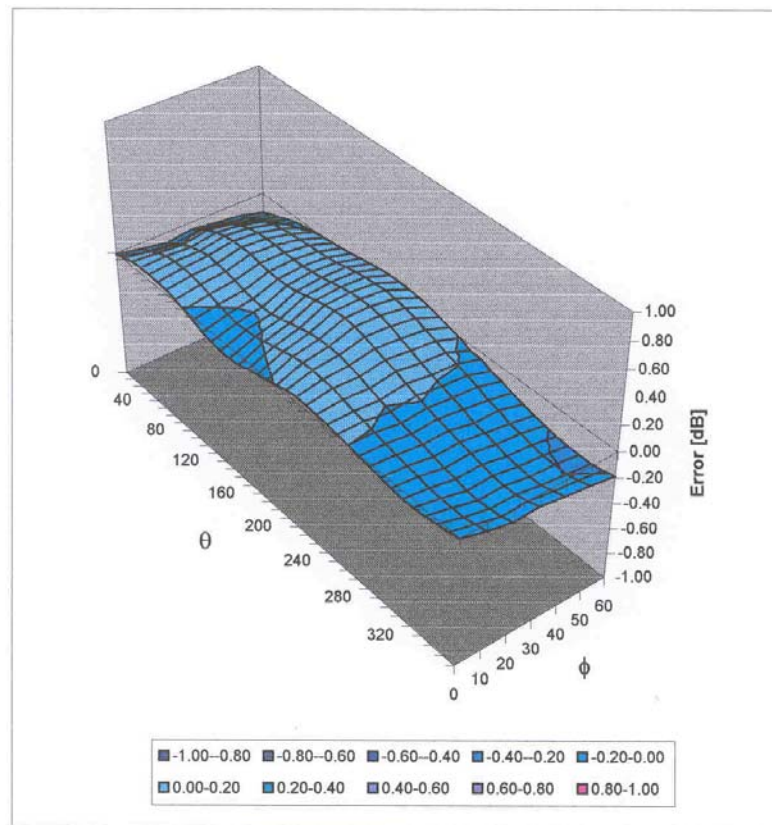
ConvF X	4.8 $\pm 8.9\%$ (k=2)	Boundary effect:
ConvF Y	4.8 $\pm 8.9\%$ (k=2)	Alpha 0.63
ConvF Z	4.8 $\pm 8.9\%$ (k=2)	Depth 2.10

ET3DV6 SN:1576

February 27, 2002

Deviation from Isotropy in HSL

Error (θ, ϕ), $f = 900$ MHz



Cardionet, Inc., Model No: CardioNet Monitor
FCC ID: QBI-1005

Date of Test: March 10 to 11, 2003

APPENDIX C – Phantom Certificate

**Schmid & Partner
Engineering AG**

Zeughausstrasse 43, 8004 Zurich, Switzerland, Phone +41 1 245 97 00, Fax +41 1 245 97 79

Certificate of conformity / First Article Inspection

Item	SAM Twin Phantom V4.0
Type No	QD 000 P40 BA
Series No	TP-1002 and higher
Manufacturer / Origin	Untersee Composites Hauptstr. 69 CH-8559 Fruthwilen Switzerland

Tests

The series production process used allows the limitation to test of first articles.
Complete tests were made on the pre-series Type No. QD 000 P40 AA, Serial No. TP-1001 and on the series first article Type No. QD 000 P40 BA, Serial No. TP-1006. Certain parameters have been retested using further series units (called samples).

Test	Requirement	Details	Units tested
Shape	Compliance with the geometry according to the CAD model.	IT'S CAD File (*)	First article, Samples
Material thickness	Compliant with the requirements according to the standards	2mm +/- 0.2mm in specific areas	First article, Samples
Material parameters	Dielectric parameters for required frequencies	200 MHz – 3 GHz Relative permittivity < 5 Loss tangent < 0.05.	Material sample TP 104-5
Material resistivity	The material has been tested to be compatible with the liquids defined in the standards	Liquid type HSL 1800 and others according to the standard.	Pre-series, First article

Standards

- [1] CENELEC EN 50361
- [2] IEEE P1528-200x draft 6.5
- [3] IEC PT 62209 draft 0.9
- (*) The IT'S CAD file is derived from [2] and is also within the tolerance requirements of the shapes of [1] and [3].

Conformity

Based on the sample tests above, we certify that this item is in compliance with the uncertainty requirements of SAR measurements specified in standard [1] and draft standards [2] and [3].

Date 18.11.2001

Signature / Stamp

Doc No 881 – QD 000 P40 BA – B

Page 1 (1)