## ENGINEERING TEST REPORT



### CM-16, 2.4 GHz Cordless Telephone Model No.: CM-16

### **Tested For**

Aastra Telecom Limited 155 Snow Blvd. Concord, ON Canada L4K 4N9

### In Accordance With

SAR (Specific Absorption Rate) Requirements
using guidelines established in IEEE C95.1-1991,
FCC OET Bulletin 65 (Supplement C),
Industry Canada RSS-102(Issue 1) and
ACA Radiocommunications (Electromagnetic Radiation – Human Exposure)
Amendment Standard 2000 (No. 1)

UltraTech's File No.: AAST-007-SAR

This Test report is Issued under the Authority of Tri M. Luu, Professional Engineer, Vice President of Engineering UltraTech Group of Labs

Date: September 08, 2004

Report Prepared by: JaeWook Choi, SAR System Engineer

Issued Date: September 08, 2004



Tested by:

JaeWook Choi, SAR System Engineer

Test Dates: August 25, 2004

The results in this Test Report apply only to the sample(s) tested, which has been randomly selected.

## **UltraTech**

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### **TABLE OF CONTENTS**

EXHIB	1. INTRODUCTION	5
1.1.	SCOPE	5
1.2.	REFERENCES	
EXHIB	2. PERFORMANCE ASSESSMENT	6
2.1.	CLIENT AND MANUFACTURER INFORMATION	6
2.2.	DEVICE UNDER TEST (D.U.T.) DESCRIPTION	
2.3.	LIST OF D.U.T.'S ACCESSORIES:	
2.4.	SPECIAL CHANGES ON THE D.U.T.'S HARDWARE/SOFTWARE FOR TESTING PURPOSES	7
2.5.	ANCILLARY EQUIPMENT	
2.6.	GENERAL TEST CONFIGURATIONS	7
2.6	Equipment Configuration	7
2.6	Exercising Equipment	7
2.7.	SPECIFIC OPERATING CONDITIONS	7
2.8.	BLOCK DIAGRAM OF TEST SETUP	8
EXHIB	3. SUMMARY OF TEST RESULTS	9
3.1.	LOCATION OF TESTS	9
3.2.	APPLICABILITY & SUMMARY OF SAR RESULTS	
EXHIB		
4.1.	TEST SETUP	
4.2.	PHOTOGRAPH OF D.U.T. AND ALL ACCESORIES	
4.3.	PHOTOGRAPHS OF D.U.T. POSITION	
4.3	Head Left Configuration	16
	1.1. Cheek Position	
	1.2. Tilt Position	
	Head Right Configuration	
	2.1. Cheek Position	
4.3	2.2. Tilt Position	
	Body Configuration	
4.4.	MAXIMUM PEAK SPATIAL-AVERAGE SAR	
4.4		
4.4		
4.5.	SAR MEASUREMENT DATA	
4.5		
	1.1. Left Head	
4	1.2. Right Head	
4.5	Body Configuration	
	2.1. Back-side of DUT parallel to the phantom and in contact	
4.5	Power Measurement	
2	3.1. Power Drift	24
<b>EXHIB</b>	5. SAR SYSTEM CONFIGURATION & TEST METHODOLOGY	25
5.1.	MEASUREMENT SYSTEM SPECIFICATIONS	25
5.2.	Test Procedures	2.5

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CM-16, 2.4 GHz Cordeless Phone M/N: CM-16

FCC ID: SDVCM16

5.3. Phantom	25
5.4. SIMULATED TISSUE	26
5.4.1. Preparation	27
5.5. MEASUREMENT OF ELECTRICAL CHARACTERISTICS OF SIMULATED TISSUE	
5.5.1. Slotted Coaxial Waveguide	
5.5.1.1. Equipment set-up	
5.5.1.2. Measurement procedure	
5.5.2. HP Dielectric Strength Probe System (open-ended coaxial transmission-line probe/s	
5.5.2.1. Equipment set-up	
5.5.2.2. Measurement procedure	
5.6. System Calibration	31
5.6.1. Probe linearity	
5.6.2. Free Space Calibration	
5.6.2.1. Method	
5.6.2.2. Measurement procedure	
5.6.2.3. Definition of Amplifier Settings	
5.6.3. Calculable Waveguide Calibration	
5.6.3.1. Measurement Procedure	
5.6.4. Thermal Transfer Calibration	
5.6.4.1. Measurement procedure	
5.6.4.2. Determination of Conversion Factor (γ) in the simulated tissue	
5.6.5. Data Acquisition Methodology	
5.6.5.1. E-Field Measurement	
5.6.5.2. Sensitivity( $\zeta$ ) of probe in the simulated tissue	
5.6.5.3. SAR Measurement	
5.6.5.4. Data Extrapolation and boundary effect	
5.6.6. Determining the Heat Capacity of Simulated Tissue	45
5.6.6.1. Instruments and Materials	45
5.6.6.2. Method	45
5.6.6.3. Rationale	46
5.7. SAR MEASUREMENT SYSTEM VERIFICATION	47
5.7.1. Standard Source	47
5.7.2. Standard Source Input Power Measurement	47
5.7.3. System Validation Procedure	48
5.8. Power Measurement	
5.9. Positioning of D.U.T.	50
5.10. SAR MEASUREMENT UNCERTAINTY	
5.10.1. Measurement Uncertainty	
5.10.1.1. Measurement Uncertainty evaluation for handset SAR test	
5.10.1.2. Measurement Uncertainty for System Performance Check	
EXHIBIT 6. SAR PRESCANS	55
6.1.1. Body Configuration	55
6.1.1.1. Test configurations used	
6.1.1.2. Equipment permutation investigated for each orientation	
6.1.1.3. Comments on non-tested configurations	
6.2. RECOMMENDED CAUTION STATEMENTS TO BE INCLUDED IN USERS MANUAL	
EXHIBIT 7. SAR MEASUREMENT	
7.1. HEAD CONFIGURATION	
7.1.1. Left Head	
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CM-16, 2.4 GHz Cordeless Phone M/N: CM-16

FCC ID: SDVCM16

File #: AAST-007-SAR September 08, 2004

7.1.1.2. Left Tilt Position; #05	61
7.1.2. Right Head	
7.1.2.1. Right Cheek Position; #08	
7.1.2.2. Right Tilt Position; #11	
7.2. BODY CONFIGURATION	
7.2.1. Back-side of DUT parallel to the phantom and in contact 7.2.1.1. D.U.T. perpendicular to the phantom and in contact, Generic Head	
7.2.1.1. D.U.T. perpendicular to the phantom and in contact, Generic Head 7.2.1.2. D.U.T. perpendicular to the phantom and in contact, Generic Head	
7.2.1.3. D.U.T. perpendicular to the phantom and in contact, Generic Head	
EXHIBIT 8. TISSUE DIELECTRIC PARAMETER CALIBRATION	N82
8.1. SIMULATED TISSUE AT 2450 MHz	82
8.1.1. 2450 MHz Brain Tissue	
8.1.2. 2450 MHz Muscle Tissue	
EXHIBIT 9. SAR SYSTEM CALIBRATION	83
9.1. GENERAL INFORMATION OF THE PROBE	83
9.2. PROBE LINEARITY AND DYNAMIC RANGE	
9.2.1. Diode Compression Potential	
9.2.2. Channel 1	
9.2.3. Channel 2	
9.2.4. Channel 3	
9.3. PROBE FREE SPACE CALIBRATION	
9.3.1. Calibration Setup at 2450 MHz	
9.3.2. Amplifier Settings	
9.3.3. Isotropic response	
9.4. CALCULABLE WAVEGUIDE CALIBRATION	
9.4.1. Calibration Setup	
9.4.1.1. Setup for 2400 MHz	
9.4.2. Simulated Tissue	
9.4.2.1. Brain Tissue at 2400 MHz	
9.4.2.2. Muscle Tissue at 2400 MHz	
9.4.3. Conversion Factor	
<ul><li>9.4.3.1. Calculable waveguide calibration at 2400 MHz for simulated brain</li><li>9.4.3.2. Calculable waveguide calibration at 2400 MHz for simulated mus</li></ul>	
<u> </u>	
EXHIBIT 10. SAR SYSTEM VERIFICATION	93
10.1. VERIFICATION SETUP	93
10.1.1. Test setup at 2400 MHz using the dipole reference	
10.2. SIMULATED TISSUE	
10.2.1. Simulated brain tissue at 2400 MHz	
10.3. VERIFICATION RESULT	
10.3.1. Reference SAR values for simulated brain tissue at 2400 MHz	
10.3.2. Verification result at 2400 MHz	
EXHIBIT 11. SAR CALCULATION SUMMARY	
11.1. Terminology	
11.1.1. Sensor factor( $\eta_{pd}$ and $\eta_{E2}$ ) in the air ( $Z_0 = 377[\Omega]$ )	
11.1.2. Diode Compensation (Linearity correction)	
11.1.3. Amplifier settings( $AS_i$ ) and probe output	

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### **SPECIFIC ABSORPTION RATE (SAR)**

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FCC ID: SDVCM16

IEEE C95.1-1991, FCC OET Bulletin 65 (Supplement C), Industry Canada RSS-102(Issue 1) and ACA Radiocommunications (Electromagnetic Radiation – Human Exposure) Amendment Standard 2000 (No. 1)

CM-16, 2.4 GHz Cordeless Phone M/N: CM-1	/I-16
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1111		100
11.1.4.	Conversion factor ( $\gamma$ ) in the simulated tissue	102
11.1.5.	Conversion factor (γ) Calculation #1 – Thermal Trnasfer Calibration (Below 1 GHz)	102
11.1.6.	Conversion factor (γ) Calculation #2 – Calculable Waveguide Calibration (Above 1 GHz)	103
11.1.7.	Sensitivity ( $\zeta$ ) in the simulated tissue	103
1118	SAR calculation	103

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### **EXHIBIT 1. INTRODUCTION**

### 1.1. SCOPE

Reference:	SAR (Specific Absorption Rate) Requirements		
	IEEE C95.1-1991,		
	FCC OET Bulletin 65 (Supplement C Edition 01-01)		
	Industry Canada RSS-102 (Issue 1).		
	ACA Radiocommunications (Electromagnetic Radiation – Human Exposure), Amendment		
	Standard 2000 (No. 1)		
Title	Safety Levels with respect to human exposure to Radio Frequency Electromagnetic Fields		
	Guideline for Evaluating the Environmental Effects of Radio Frequency Radiation		
Purpose of Test:	To verify compliance with Federal regulated SAR requirements in Canada, Chatswood NSW 2067		
	and the US.		
Method of Measurements:	IEEE C95.1-1991, FCC OET Bulletin 65 (Supplement C Edition 01-01) and Industry Canada		
	RSS-102 (Issue 1)		
<b>Device Category</b>	Portable		
<b>Exposure Category</b>	General Population/Uncontrolled		

### 1.2. REFERENCES

The methods and procedures used for the measurements contained in this report are details in the following reference standards:

Publications	Year	Title
IEEE Std. 1528/D1.2		Draft Recommended practice for determining the Peak Spatial-Average Specific
	2003	Absorption rate (SAR) in the Human Body Due to Wireless Communications Devices:
		Experimental Techniques.
Industry Canada RSS102	1999	"Evaluation Procedure for Mobile and Portable Radio Transmitters with respect to
	1999	Health Canada's Safety Code 6 for Exposure of Humans to Radio Frequency Fields"
ACA	2000	ACA Radiocommunications (Electromagnetic Radiation – Human Exposure)
2000		Amendment Standard 2000 (No. 1)
NCRP Report No.86	1986	"Biological Effects and Exposure Criteria for radio Frequency Electromagnetic Fields"
FCC OET Bulletin 65	lletin 65 "Evaluating Compliance with FCC Guidelines for Human Exposure to radio	
1997		Fields"
ANSI/IEEE C95.3	1992	"Recommended Practice for the Measurement of Potentially Hazardous
	1992	Electromagnetic Fields - RF and Microwave"
ANSI/IEEE C95.1 "Safe		"Safety Levels with Respect to Human Exposure to Radio Frequency Electromagnetic
	1992	Fields, 3kHz to 300GHz"
AS/NZS 2722.1	2722.1 Interim Chatswood NSW 2067n/New Zealand Standard. "Radiofrequency 2	
	1998	1:Maximum exposure levels – 3kHz to 300GHz "

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### **EXHIBIT 2. PERFORMANCE ASSESSMENT**

### 2.1. CLIENT AND MANUFACTURER INFORMATION

APPLICANT:	
Name:	Aastra Telecom Limited
Address:	155 Snow Blvd.
	Concord, ON
	Canada L4K 4N9
Contact Person:	Mr. Wah Kung
	Phone #: +1 905 760 4200
	FAX #: +1 905 760 4235
	Email Address: wkung@aastra.com

MANUFACTURER:		
Name:	Victory Telecom (Huizhou) Ltd.	
Address:	No. 11, Lon Feng Du Tian Industrial District	
	Huizhou Guandong	
	China	
Contact Person:	Mr. Lai Jian Hui	
	Phone #: (0752) 2371871	
	FAX #: (0752) 2372118	
	Email Address: vchz@pub.huizhou.gd.cn	

### 2.2. DEVICE UNDER TEST (D.U.T.) DESCRIPTION

The following is the information provided by the applicant.

Trade Name	Aastra Telecom Limited
Type/Model Number	CM-16
Frequency of Operation	2400 ~ 2483 MHz
Rated RF Output Power	23 dBm
Modulation Employed	GFSK, TDD
Antenna	Tyco's surface mount antenna (M/N: 1513150-1, Gain: > 4.5 dBi)
Power Supply Rechargeable Ni-MH battery (3.6 V)	
Primary User Functions of D.U.T.	Voice communication through air

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### 2.3. LIST OF D.U.T.'S ACCESSORIES:

Rechargeable Ni-MH battery (3.6 V), Belt clip

## 2.4. SPECIAL CHANGES ON THE D.U.T.'S HARDWARE/SOFTWARE FOR TESTING PURPOSES

N/A

### 2.5. ANCILLARY EQUIPMENT

N/A

### 2.6. GENERAL TEST CONFIGURATIONS

### 2.6.1. Equipment Configuration

Power and signal distribution, grounding, interconnecting cabling and physical placement of equipment of a test system shall simulate the typical application and usage in so far as is practicable, and shall be in accordance with the relevant product specifications of the manufacturer.

The configuration that tends to maximize the D.U.T.'s emission or minimize its immunity is not usually intuitively obvious and in most instances selection will involve some trial and error testing. For example, interface cables may be moved or equipment re-orientated during initial stages of testing and the effects on the results observed.

Only configurations within the range of positions likely to occur in normal use need to be considered.

The configuration selected shall be fully detailed and documented in the test report, together with the justification for selecting that particular configuration.

### 2.6.2. Exercising Equipment

The exercising equipment and other auxiliary equipment shall be sufficiently decoupled from the D.U.T. so that the performance of such equipment does not significantly influence the test results.

### 2.7. SPECIFIC OPERATING CONDITIONS

The D.U.T had been tested with three different kind of generic headsets from a local retail store plugged into the handset respectively as a consideration of the normal operating usage.

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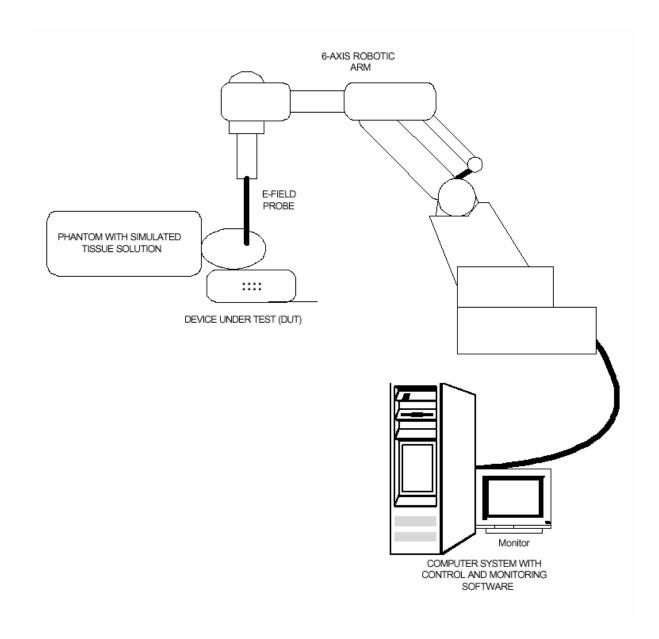
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### 2.8. BLOCK DIAGRAM OF TEST SETUP

The D.U.T. was configured as normal intended use. The following block diagram shows a representative equipment arrangement during tests:



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### **EXHIBIT 3. SUMMARY OF TEST RESULTS**

### 3.1. LOCATION OF TESTS

All of the measurements described in this report were performed at UltraTech Group of Labs located at:

3000 Bristol Circle, in the city of Oakville, Province of Ontario, Canada.

All measurements were performed in UltraTech's shielded chamber, 24' x 16' x 8'.

### 3.2. APPLICABILITY & SUMMARY OF SAR RESULTS

The maximum peak spatial - average SAR measured was found to be 0.33 W/Kg.

Exposure Category and SAR Limits	Test Requirements	Compliance (Yes/No)
General population/Uncontrolled exposure  0.08W/kg whole body average and spatial peak SAR of  1.6W/kg, averaged over 1gram of tissue  Hands, wrist, feet and ankles have a peak SAR not to exceed  4 W/kg, averaged over 10 grams of tissue.	Requirements using guidelines established in IEEE C95.1-1991  FCC OET Bulletin 65 (Supplement C Edition 01-01)  Industry Canada RSS-102 (Issue 1).  ACA Radiocommunications (Electromagnetic Radiation – Human Exposure) Amendment Standard 2000 (No. 1)	YES
Occupational/Controlled Exposure  0.4W/kg whole body average and spatial peak SAR of  8W/kg, averaged over 1gram of tissue Hands, wrist, feet and ankles have a peak SAR not to exceed 20 W/kg, averaged over 10 grams of tissue.	Requirements using guidelines established in IEEE C95.1-1991  FCC OET Bulletin 65 (Supplement C Edition 01-01),  Industry Canada RSS-102 (Issue 1)  ACA Radiocommunications (Electromagnetic Radiation – Human Exposure) Amendment Standard 2000 (No. 1)	N/A

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## **EXHIBIT 4. MEASUREMENTS, EXAMINATIONS & TEST DATA**

### 4.1. TEST SETUP

D.U.T. Information		Condition			
Product Name	CM-16, 2.4 GHz Cordless Telephone	Robot Type	6 Axis		
Model Number	CM-16	Scan Type	SAR - Area/Zoom/Att Vs Depth		
Serial Number	Pre Production	Measured Field	Е		
Frequency Band [MHz]	2400 ~ 2483	Phantom Type	2 <sub>mm</sub> base Flat Phantom		
Frequency Tested [MHz]	2401.056, 2440.800, 2482.272	Phantom Position	Waist		
Rated RF Output Power [dBm]	23	Room Temperature [°C]	$24.0 \pm 1$		
Antenna Type	Surface mount (M/N: 1513150-1, Gain: > 4.5 dBi)	Room Humidity [%]	40 ± 10		
Modulation	TDD	Tissue Temperature [°C]	$24.0 \pm 1$		
Worst Case Duty Cycle	12.5 % (1:8)				
<b>Duty Cycle Tested</b>	12.5 % (1:8)				
Source-based time-average	1				

Type of Tissue	Brain	Muscle
Test Frequency [MHz]	2450	2450
Measured Dielectric Constant	38.2 (-2.6 %)	50.2 (-4.8 %)
Target Dielectric Constant	39.2	52.7
Measured Conductivity [S/m]	1.78 (-1.0 %)	1.94 (-0.6 %)
Target Conductivity [S/m]	1.80	1.95
Penetration Depth (Plane Wave Excitation) [mm]	18.7	19.6
Probe Model Number	ET20	ET20
Probe Serial Number	03JUN-0028	03JUN-0028
Probe Orientation	Isotropic	Isotropic
Probe Offset [mm]	2.00	2.00
Probe Tip Diameter [mm]	4.00	4.00
Sensor Factor $(\eta_{pd})_{[mV/(mW/cm)]}^2$	10.8	10.8
Conversion Factor (γ)	4.539	4.068
Sensitivity (ζ) <sub>[W/Kg/mV]</sub>	1.369E-01	1.613E-01

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### 4.2. PHOTOGRAPH OF D.U.T. AND ALL ACCESORIES



< Front View >

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< Rear View >

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< Belt clip>

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< Generic Headset #1 >



< Generic Headset #2 >

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< Generic Headset #3 >

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### 4.3. PHOTOGRAPHS OF D.U.T. POSITION

### 4.3.1. Head Left Configuration

### 4.3.1.1. Cheek Position



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### 4.3.1.2. Tilt Position



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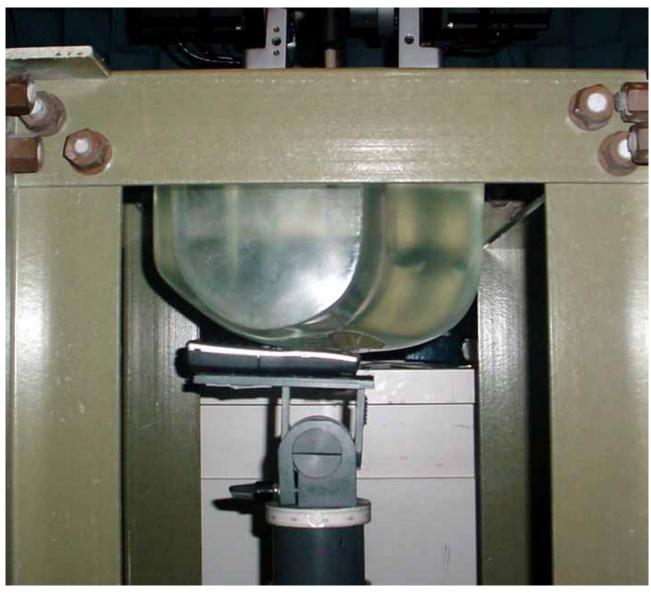
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### 4.3.2. Head Right Configuration

### 4.3.2.1. Cheek Position



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### 4.3.2.2. Tilt Position



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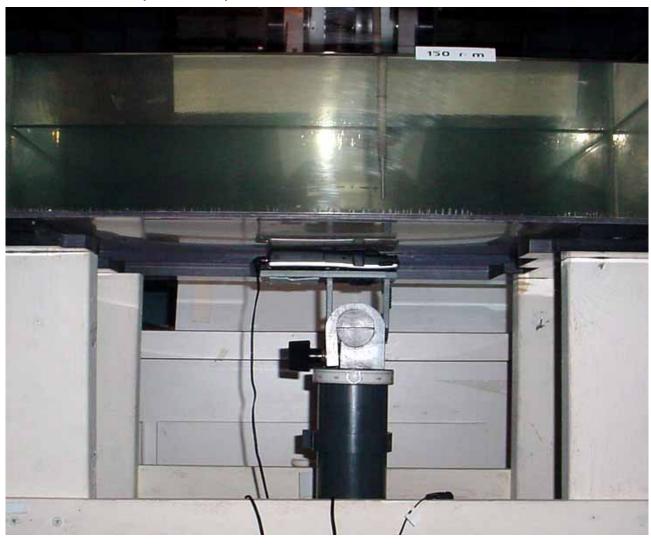
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### 4.3.3. Body Configuration

### 4.3.3.1. Back- side of D.U.T. parallel to the phantom and in contact



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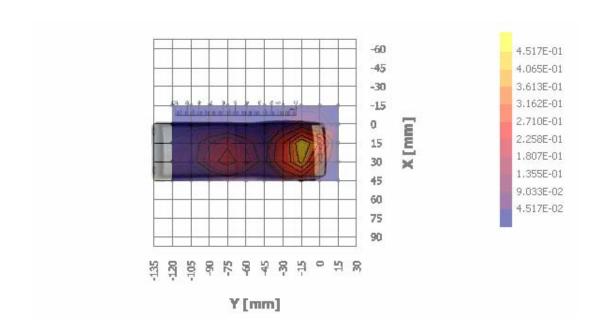
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#### **MAXIMUM PEAK SPATIAL-AVERAGE SAR** 4.4.

#### 4.4.1. Maximum Peak Spatial-average SAR Data

#	Configuration	Device Test Positions	Antenna Position	Freq.	Channel	MAX. SAR [W/Kg]
*	General Population/Uncontrolled Exposure Category Limit					1.6
20	Headset #3 Back-side of DUT parallel to the phantom and in contact	Body	Integrated	2440.800	Middle	0.33

#### 4.4.2. **Maximum Peak Spatial-Average SAR Location**



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File #: AAST-007-SAR September 08, 2004

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### 4.5. SAR MEASUREMENT DATA

### 4.5.1. Head Configuration\*

### 4.5.1.1. Left Head

#	Configuration	Antenna Position	Frequency [MHz]	Channel	SAR <sub>local</sub> Before [W/Kg]	SAR <sub>local</sub> After [W/Kg]	MAX SAR <sub>1g</sub> [W/Kg]
*	* General population/Uncontrolled Exposure Category Limit						1.6
01	Left Head		2401.056	Lowest			-
02	Cheek Position	Integrated	2440.800	Middle	0.01	0.01	0.04
03			2482.272	Highest			-
04	Left Head		2401.056	Lowest			-
05	Tilt Position	Integrated	2440.800	Middle	0.01	0.01	0.03
06			2482.272	Highest			-

### 4.5.1.2. Right Head

#	Configuration	Antenna Position	Frequency [MHz]	Channel	SAR <sub>local</sub> Before [W/Kg]	SAR <sub>local</sub> After [W/Kg]	$\begin{array}{c} MAX \\ SAR_{1g} \\ {}_{[W/Kg]} \end{array}$
*	* General population/Uncontrolled Exposure Category Limit						1.6
07	Right Head		2401.056	Lowest			-
08	Cheek Position	Integrated	2440.800	Middle	0.02	0.02	0.05
09			2482.272	Highest			-
10	Right Head		2401.056	Lowest			-
11	Tilt Position	Integrated	2440.800	Middle	0.02	0.01	0.04
12			2482.272	Highest			-

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<sup>\*</sup> If the SAR measured at the middle channel for each test configuration is at least 3.0 dB lower than the SAR limit, testing at the high and low channel is optional for such test configuration(s).

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#### **Body Configuration**\* 4.5.2.

#### 4.5.2.1. Back-side of DUT parallel to the phantom and in contact

#	Configuration	Antenna Position	Frequency [MHz]	Channel	SAR <sub>local</sub> Before [W/Kg]	SAR <sub>local</sub> After [W/Kg]	$\begin{array}{c} MAX \\ SAR_{1g} \\ {\tiny [W/Kg]} \end{array}$
*	* General population/Uncontrolled Exposure Category Limit						1.6
13	Headset #1		2401.056	Lowest			-
14	Back-side of DUT parallel to the phantom and in contact	Integrated	2440.800	Middle	0.12	0.10	0.29
15			2482.272	Highest			-
16	Headset #2		2401.056	Lowest			-
17	Back-side of DUT parallel to the phantom and in	Integrated	2440.800	Middle	0.16	0.15	0.31
18	contact		2482.272	Highest			-
19	Headset #3		2401.056	Lowest			-
20	Back-side of DUT parallel to the phantom and in	Integrated	2440.800	Middle	0.15	0.15	0.33
21	contact		2482.272	Highest			-

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<sup>\*</sup> If the SAR measured at the middle channel for each test configuration is at least 3.0 dB lower than the SAR limit, testing at the high and low channels is optional for such test configuration(s).

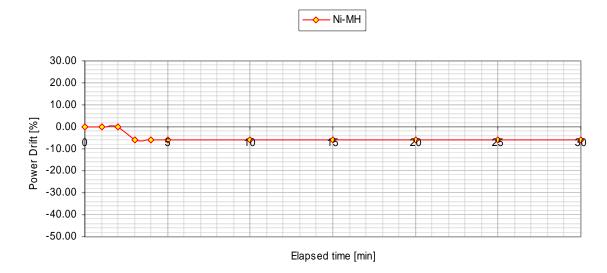
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### 4.5.3. Power Measurement

### 4.5.3.1. Power Drift

The local SAR was measured at the arbitrary location in the vicinity of the antenna fed point in the simulated tissue at 2440.800 MHz (11 Mbps) during the period of 30 minute.

The power (SAR) drift after 30 minutes of the continuous exposure at the maximum power level (at the maximum duty cycle\*) was found to be less than -5.88 [%].



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<sup>\*</sup> Refer to 2.7. Specific Operating Conditions

# EXHIBIT 5. SAR SYSTEM CONFIGURATION & TEST METHODOLOGY

### 5.1. MEASUREMENT SYSTEM SPECIFICATIONS

Positioning Equipment	Probe
Type : 3D Near Field Scanner	Sensor : E-Field
Location Repeatability: 0.1 [mm]	Spatial Resolution : 1 [mm <sup>3</sup> ]
Speed 180 [°/sec]	Isotropic Response : ±0.25 [dB]
AC motors	Dynamic Range: 0.01 to 100 [W/Kg]
Computer	Phantom
Type: Pentium III 500MHz	Tissue : Simulated Tissue with electrical characteristics similar to those
Memory: 256 MB RAM	of the human at normal body temperature.
Operating System : Windows 2000 Pro	Left/Right Head: IEEE P1528 Compliant SAM manufactured by Aprel Body/Frontal Head: IEEE Flat Phantom 2 [mm] Base
Monitor : 19" SVGA	

### 5.2. TEST PROCEDURES

In the SAR measurement, the positioning of the probes must be performed with sufficient accuracy to obtain repeatable measurements in the presence of rapid spatial attenuation phenomena. The accurate positioning of the E-field probe is accomplished by using a high precision robot. The robot can be taught to position the probe sensor following a specific pattern of points. In a first sweep, the sensor is positioned as close as possible to the interface, with the sensor enclosure touching the inside of the phantom shell. The SAR is measured on a grid of points, which covers the curved surface of the phantom in an area larger than the size of the D.U.T. After the initial scan, a high-resolution volume gird is used to locate the absolute maximum measured energy point and to calculate the peak spatial-average SAR. At this location, attenuation versus depth scan will be accomplished by the measurement system in order to verify the peak spatial-average SAR measured.

### 5.3. PHANTOM

For Head mounted devices placed next to the ear, the phantom used in the evaluation of the RF exposure of the user of the wireless device is a IEEE P1528 compliant SAM phantom, shaped like a human head and filled with a mixture simulating the dielectric characteristics of the brain. A left sided head and a right sided head are evaluated to determine the worst case orientation for SAR. For body mounted and frontal held push-to-talk devices, a flat phantom of dimensions 70x42x20cm with a base plate thickness of 2mm is used.

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File #: AAST-007-SAR September 08, 2004

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#### 5.4. SIMULATED TISSUE

Simulated Tissue: Suggested in a paper by George Hartsgrove and colleagues in University of Ottawa Ref.: Bioelectromagnetics 8:29-36 (1987)

Ingredient	Quantity
Water	40.4 %
Sugar	56.0 %
Salt	2.5 %
HEC	1.0 %
Bactericide	0.1 %

Table 5.4. Example of composition of simulated tissue

This simulated tissue is mainly composed of water, sugar and salt. At higher frequencies, in order to achieve the proper conductivity, the solution does not contain salt. Also, at these frequencies, D.I. water and alcohol is preferred.

Target Frequency	Не	ad	Во	ody
(MHz)	$\epsilon_{\rm r}$	σ (S/m)	$\epsilon_{\rm r}$	σ (S/m)
150	52.3	0.76	61.9	0.80
300	45.3	0.87	58.2	0.92
450	43.5	0.87	56.7	0.94
835	41.5	0.90	55.2	0.97
900	41.5	0.97	55.0	1.05
915	41.5	0.98	55.0	1.06
1450	40.5	1.20	54.0	1.30
1610	40.3	1.29	53.8	1.40
1800 – 2000	40.0	1.40	53.3	1.52
2450	39.2	1.80	52.7	1.95
3000	38.5	2.40	52.0	2.73
5800	35.3	5.27	48.2	6.00

 $(\varepsilon_r = \text{relative permittivity}, \sigma = \text{conductivity and } \rho = 1000 \text{ Kg/m}^{3*})$ 

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<sup>\*</sup> The actual mass density of the equivalent tissue vary based on the composition of the tissue from 990 Kg/m<sup>3</sup> to 1,300 Kg/m<sup>3</sup>.

File #: AAST-007-SAR September 08, 2004

### 5.4.1. Preparation

The weight requirements is determined and measured carefully for all the components. A clean container is used where the ingredients will be mixed. A stirring paddle mounted to a drill press is used to stir the mixture. First the heat is applied to the DI water to approximately 40 °C to help the ingredients dissolve well and then the salt and the bactericide are added. It is stirred until all the ingredients are completely dissolved. It is continuously stirred slowly while adding the sugar. Rotation of stirring paddle at a high RPM is avoided to prevent air bubbles in the mixture. Later on, the HEC is added to maintain the solution homogeneous. Mixing time is approximately 2 hours.

### 5.5. MEASUREMENT OF ELECTRICAL CHARACTERISTICS OF SIMULATED TISSUE

- 1) Slotted Coaxial Waveguide
- 2) HP Dielectric Strength Probe System

### 5.5.1. Slotted Coaxial Waveguide

### 5.5.1.1. Equipment set-up

The test equipment consists of a slotted coaxial transmission line with a probe connected to a vector network analyzer, as shown in Figure 5.5.1.1 The log-magnitude and phase of  $S_{21}$  should be displayed simultaneously. Source power should be set to a level high enough to provide good signal-to-noise ratio. Periodically (annually or whenever the measuring scale along the line length is changed) a measurement is made on a reference liquid to validate the system. Since the measured quantities are magnitude and phase changes versus distance, the accuracy of the scale is very critical.

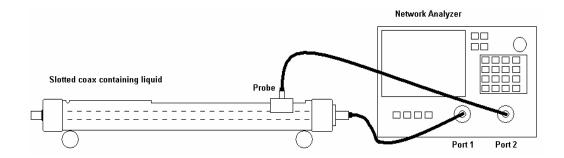


Figure 5.5.1.11 Slotted line set-up

The network analyzer injects a signal into one end of the slotted coaxial transmission line. The probe inserted through the slot into the tissue-equivalent material detects the RF amplitude and phase for each measurement position along the length of the line. A full two-port calibration of the network analyzer should be carried out prior to connecting the sample holder, and the following precautions should be observed:

a) Fill the slotted line carefully to avoid trapping air bubbles. This operation should be performed while the slotted line is horizontal.

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- b) The probe should be inserted into the slot at the end nearest to the input connector of the slotted line, ensuring that the tissue-equivalent liquid is flush with the inside surface of the line, and aligned with a well-defined position on the distance scale of the slotted line.
- c) The probe should be inserted perpendicular to the slotted—line longitudinal axis until a stable and adequate amplitude response is achieved. Do not insert the probe too deeply into the coaxial line, because it can overly perturb the field distribution.

### 5.5.1.2. Measurement procedure.

- a) Configure and calibrate the network analyzer.
- b) Measure 10 to 20 log-magnitude and phase data points along the slotted line corresponding to about a 30 dB change in magnitude.
- c) Plot  $S_{21}$  log-magnitude and phase vs. measurement distance.
- d) Determine if the graphed points closely follow a straight-line approximation, based on the correlation coefficient or a similar statistical measure. The data should produce a good linear curve fit (expected correlation coefficient r² > 0.99 for lossy materials). If not, re-measure the liquid by increasing the sample points to extend the magnitude change from 30 to 40 dB. Note: for low loss materials, ensure that the slotted line is long enough to avoid reflections from the load-terminated end.
- e) Calculate the conductivity and relative permittivity of the tissue-equivalent material using Equations (5.5.1.21) derived from

$$\overline{\alpha} = \frac{m_m \ln(10)}{20} \qquad \text{Np/cm}$$

$$\overline{\beta} = \frac{m_p \pi}{180} \qquad \text{rad/cm}$$

$$\varepsilon'_r = \frac{(\overline{\beta})^2 - (\overline{\alpha})^2}{\omega^2 \mu_0 \varepsilon_0}$$

$$\sigma = \frac{2\overline{\alpha} \overline{\beta}}{\omega \varepsilon_0} \left(\frac{100 \text{ cm}}{\text{m}}\right) \qquad \text{S/m}$$
(Eq 5.5.1.21)

where,  $m_m$  and  $m_p$  are the slopes of the least-squares linear fits of the log-magnitude and phase plots, respectively, and  $\overline{\alpha}$  and  $\overline{\beta}$  are the average attenuation and propagation coefficients along the line.

### 5.5.2. HP Dielectric Strength Probe System (open-ended coaxial transmission-line probe/sensor)

### 5.5.2.1. Equipment set-up

The equipment consists of a probe connected to one port of a vector network analyzer. The probe is an open-ended coaxial line, as shown in Figure 5.5.2.11. Cylindrical coordinates  $(\rho, \phi, z)$  are used where  $\rho$  is the radial distance from the axis,  $\phi$  is the angular displacement around the axis, z is the displacement along the axis, z is the inner conductor radius, and z is the outer conductor inner radius.

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FCC ID: SDVCM16

The sample holder is a non-metallic container that is large compared with the size of the probe immersed in it. A probe with an outer diameter b of 2 to 4 mm is suitable for the measurement of tissue-equivalent materials in the 300 MHz to 3 GHz frequency range. This probe size is commensurate with sample volumes of 50 cc or higher. Larger probes of up to 7 mm outer diameter b may be used with larger sample volumes. A flange is typically included to better represent the infinite ground-plane assumption used in admittance calculations.

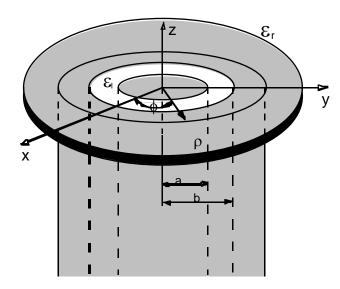


Figure 5.5.2.1An open-ended coaxial probe with inner and outer radii a and b, respectively

The accuracy of the short-circuit measurement should be verified for each calibration at a number of frequencies. A short circuit can be achieved by gently pressing a piece of aluminum foil against the open end. For best electrical contact, the probe end should be flat and free of oxidation. Larger the sensors generally have better foil short-circuit repeatability. It is possible to obtain good contact with some commercial 4.6 mm probes using the metal-disk short-circuit supplied with the kit. For best repeatability, it may be necessary to press the disk by hand.

The network analyzer is configured to measure the magnitude and phase of the admittance. A one-port reflection calibration is performed at the plane of the probe by placing materials for which the reflection coefficient can be calculated in contact with the probe. Three standards are needed for the calibration, typically a short circuit, air, and deionized water at a well-defined temperature (other reference liquids such as methanol or ethanol may be used for calibration). The calibration is a key part of the measurement procedure, and it is therefore important to ensure that it has been performed correctly. It can be checked by re-measuring the short circuit to ensure that a reflection coefficient of  $\Gamma = -1.0$  (linear units) is obtained consistently.

### 5.5.2.2. Measurement procedure

a) Configure and calibrate the network analyzer and probe system.

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- b) Place the sample in a non-metallic container and immerse the probe. A fixture or clamp is recommended to stabilize the probe, mounted such that the probe face is at an angle with respect to the liquid surface to minimize trapped air bubbles beneath the flange.
- c) Measure the complex admittance with respect to the probe aperture.
- d) Compute the complex relative permittivity  $\varepsilon_r = \varepsilon_r' j \sigma / \omega \varepsilon_0$ .

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### 5.6. SYSTEM CALIBRATION

The SAR measurement system has two main components:

- a) the probe, which is connected to the inputs of
- b) the instrumentation amplifier whose outputs are connected through the optical transmission line to
- c) the computer.

The system is calibrated as one unit not as individual components. If any components is modified or replaced, the system must be re-calibrated.

The system calibration is performed by three steps:

- 1) determination of the diode compression potential for each channel by introducing the probe into the well-defined RF field and and record the probe output while increasing the RF field intensity,
- 2) determination of the sensitivity of the probe in the air by introducing it into the well-defined RF field, and
- 3) correlation of the measured E-field in the dielectric medium to the temperature rise in a dielectric medium.

### 5.6.1. Probe linearity

Detector diodes at the dipole feed-point are used to rectify the sensor voltage output. The rectified signal is transmitted through resistive (RF-transparent) lines to the sensor amplifier. At low field strength levels the output voltage is proportional to the square of the amplitude of the incident field; at higher signal levels, the output voltage is not linearly proportional to  $|E|^2$ , but becomes proportional to E. The compensation for diode compression is carried out for the each detector diode using the follwing algorithm before any further evaluation.

$$U_{L,i} = U_{O,i} + \frac{U_{O,i}^{2}}{DCP_{i}}$$

Where,

 $U_{O.i}$  Probe raw output for channel i

U<sub>Li</sub> Linearized probe output for channel i

DCP<sub>i</sub> Diode specific compression potential for channel i

### 5.6.2. Free Space Calibration

- RF Signal Generator frequency range to at least 6 GHz,
- RF Amplifier if needed to generate the required power density in the test cell,
- Test Cell TEM (Crawford) cell, waveguide, or other device capable of maintaining a uniform field,
- RF Power Meter capable of measuring at least 5 Watts (current calibration is mandatory!) if possible traceable to the National Institute of Standards and Technology (NIST).
- E-Field Probe (under calibration)
- Probe Support Fixture
- Instrumentation Amplifier

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- Transmission Line
- Computer Program with the Automated Calibration System Program

### 5.6.2.1. Method

Due to impedance variations in the diodes and the transmission line, and slight differences in gain among the channels of the instrumentation amplifier, a normalization method had been designed. The calibration method actually used is to determine the factors necessary adjust each channel of the system so it's indicated output can then be equated to the well-defined RF field. These factors are referred to as "Amplifier Settings".

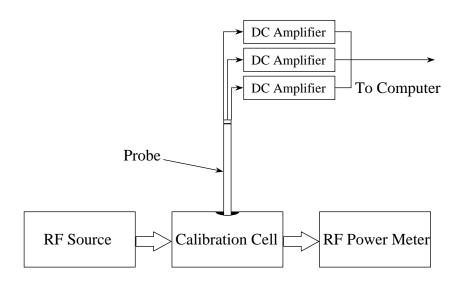


Figure 5.6.2.1 Free Space Calibration Setup for Amplifier Setting

### 5.6.2.2. Measurement procedure

Free Space Calibration of E-field probes can be performed using a TEM cell manufactured by IFI (Instrumentation for Industry, Farmingdale, NY 11735) with operating frequency at or below 1 GHz. Above 1 GHz, waveguides are used to calibrate the probes in free space.

• Connect the equipments as shown in Figure 5.6.2.1;

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- Adjust the RF generator output so that the power density at the calibration point inside the TEM cell is well-defined. (For the IFI model CC-110 cell, the uniform power density of 1.0 [mW/cm<sup>2</sup>] requires the power level of 271.0 [mW]);
- Mount the probe of the system to calibrate in the support fixture. Insert the probe through the aperture of the TEM cell. The probe handle should be at the geometric center of the aperture, i.e. midway between the septum and the upper surface, and orthogonal to the side of the cell. The sensing portion of the probe should be located at a point halfway across the depth of the cell (volumetric center).
- Once the prescribed position is obtained, it must be maintained during the rest of the measurement. The only movement of the probe allowed is rotation on its axis to position the dipole in the plane of the E-field and, for channel 3 only, parallel to the vertical uniform field (max./min. output).
- Verify that the RF power level remains constant throughout the measurement. While the probe is being rotated through 360 degrees, software indicators will show the maximum measured on each channel.

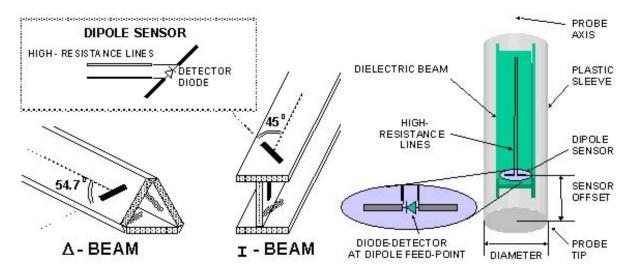


Figure 5.6.2.2 E-field probe construction

### 5.6.2.3. Definition of Amplifier Settings

The initial sequence of probe calibrations steps performed with SAR determinations produces the factors used in scaling probe output voltage to RF power density. For historical reasons all probes factors are compared to a factor 10.8 [mV] per [mW/cm²] that was typical of a prototype probe, but is in fact an arbitrary number used as an intermediately constant. The factor of 10.8 [mV/(mW/cm²)] is known as the sensor factor to the uniform power density  $(\eta_{pd})$ , but does not change. Also we can derive 10.8/3,770 [mV/(V/m)²] of the sensor factor to the  $|E|^2$   $(\eta_{E2})$ , providing 377 [ $\Omega$ ] as free space impedance.

$$\eta_{Pd} = 10.8[mV/(mW/cm^2)] \equiv \eta_{E2} = \frac{10.8}{3770}[mV/(V/m)^2]$$

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$$Pd[mW/cm^{2}] = \frac{PO_{tot}}{\eta_{Pd}}, |E|^{2}[(V/m)^{2}] = \frac{PO_{tot}}{\eta_{E2}} \text{ and } SAR = \frac{\sigma \times \frac{PO_{tot}}{\eta_{E2}}}{\rho}$$

To calibrate a probe, each channel is assigned an amplifier setting. This factor is obtained from the sum of the probe output voltage measured during probe calibration. This probe output voltage is corrected for any DC offset of the instrumentation amplifier, usually a very small amount.

During calibration, the sensitivity for the E-field tangential to the dipole axis caused by the geometry of the probe construction is carefully considered to obtain the correct amplifier setting for each channel. Thus, the amplifier settings for each channel are as follows:

$$R_{ISO,i,1} = \frac{\sum_{\theta}^{360^{\circ}} U_{L,1}(\theta)}{\sum_{\theta}^{360^{\circ}} U_{L,i}(\theta)}$$

$$AS_{i} = \eta_{Pd} [mV/(mW/cm^{2})] \times Pd[mW/cm^{2}] \times \cos^{2}(\varphi - \phi_{i}) \times \frac{R_{ISO,i,1}}{U_{L,1,peak}}$$

Where,

R<sub>ISO,i,1</sub> The isotropic correction factor normalized to channel 1

AS<sub>i</sub> Amplifier Setting for channel i

 $\eta_{pd}$  Sensor Factor to the uniform power density, an arbitrary value 10.8 [mV/(mW/cm<sup>2</sup>)]

 $\eta_{\rm E2}$  Sensor Factor to the uniform  $|E|^2$ , an arbitrary value 10.8/3,770 [mV/(V/m)<sup>2</sup>]

 $U_{L,1,peak}$  The peak linearized probe output recorded for channel 1 by a rotation about the probe axis with the probe in a test cell

 $\varphi$  Smaller angle between the probe axis and the direction of the E-field (i.e. 90° when the probe axis is parallel to the plane of the septum inside TEM cell)

 $\phi_i$  Smaller angle between the probe axis and the dipole sensor axis of the channel i ( $\phi_1 = \phi_2 = 45^{\circ}$ ,  $\phi_3 = 90^{\circ}$  for I-beam probe, and  $\phi_1 = \phi_2 = \phi_3 = 54.7^{\circ}$  for triangular-beam probe)

Pd Well-defined power density [mW/cm<sup>2</sup>] at the calibration point in a test cell

### **5.6.3.** Calculable Waveguide Calibration

### 5.6.3.1. Measurement Procedure

This method utilizes a setup in which the field can be calculated analytically based on measurements of other physical parameters, e.g., input power. While this method corresponds to the standard field method for probe calibration in air, there is no standard method defined for field generation in lossy liquids.

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When this method is used for calibrating probes in lossy liquids, the following points must be considered in the uncertainty assessment:

- 1. The net RF power dissipated in the waveguide must be measured accurately. This requirement implies precise measurements of two out of the following three quantities: incident power, reflected power, reflection coefficient at the waveguide input port.
- 2. The accuracy of the calculated field strength will depend on the assessment of the dielectric parameters of the liquid.
- 3. Because of the short wavelength in liquids with high permittivity, higher order modes may be excitable even in small waveguides. Consequently, stringent electrical and mechanical tolerances are required in the construction of calibration fixtures. The actual field distribution in the calibration setup must be carefully checked for conformity with the theoretical field distribution.

Waveguides can be utilized to generate an analytically known field inside tissue-equivalent liquids, e.g., using the setup presented in Figure 5.6.3.1. In this setup, a portion of an upright-standing open waveguide is filled with a tissue-equivalent liquid. A dielectric slab at a distance greater than one free-space wavelength from the input coupler provides an impedance match between air and the liquid with a return loss typically greater than 10 dB. Proper fabrication symmetry and the high losses in the liquid ensure that the field distribution inside the tissue-equivalent liquid follows the  $TE_{10}$  mode pattern, although it is theoretically possible that higher order modes could be excited. The field distribution was carefully validated by means of a complete volume scan in the liquid, which showed a deviation from the theoretical  $TE_{10}$  pattern of less than 2%.

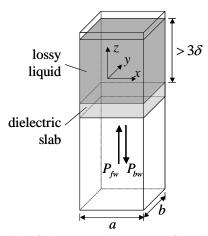


Figure 5.6.3.1 – Vertical rectangular waveguide calibration setup

Because of the low cutoff frequency, the field inside the liquid nearly propagates as a TEM wave. The depth of the medium (greater than three penetration depths) ensures that reflections at the upper surface of the liquid are negligible. The power absorbed in the liquid is determined by measuring the waveguide forward and reflected power. Equation (5.6.3.11) shows the relationship between the SAR at the cross-sectional center (x = y = 0) of the lossy waveguide and the longitudinal distance (z) from the dielectric separator

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$$SAR(z) = \frac{4(P_f - P_r)}{\rho \delta \, ab} e^{-2z/\delta} \,, \tag{Eq. 5.6.3.11}$$

where the density  $\rho$  is conventionally assumed to be 1000 kg/m³, ab is the cross-sectional area of the waveguide,  $P_f$  and  $P_r$  are the forward and reflected power inside the lossless section of the waveguide, respectively. The penetration depth  $\delta$ , which is the reciprocal of the waveguide-mode attenuation coefficient  $\alpha$ , is determined from a scan along the z-axis and compared with the theoretical value determined from Equation (5.6.3.12) using the measured dielectric properties of the lossy liquid.

$$\delta = \alpha^{-1} = \left\{ \text{Re} \left[ \sqrt{(\pi/a)^2 + j\omega\mu_0 (\sigma + j\omega\varepsilon_0 \varepsilon_r')} \right] \right\}^{-1}.$$
 (Eq. 5.6.3.12)

Table A.1 can be used for designing calibration waveguides with a return loss greater than 30 dB at the most important frequencies used for personal wireless communications. Values for the penetration depth for these specific fixtures and tissue-simulating mixtures are also listed in Table 5.6.3.11.

This calibration technique provides excellent accuracy, with standard uncertainty reported to be less than 3.6% depending on the frequency and medium. The calibration itself is reduced to power measurements traceable to a standard calibration procedure.

The waveguide method described here has the additional benefit of providing an alternative method to determine the electrical conductivity of the tissue-equivalent liquid. For a known relative permittivity, the conductivity is obtained using the equation

$$\sigma = \frac{2\alpha}{\omega\mu_0} \sqrt{\alpha^2 + \omega^2 \mu_0 \varepsilon_0 \varepsilon_r' - \left(\frac{\pi}{a}\right)^2} .$$
 (Eq. 5.6.3.13)

<b>Table 5.6.3.11 – Guidelines for designing calibration waveguides</b>	

	Head Tissue Simulant		Waveguide Dimension	Penetration Depth	Dielectric	Separator
Frequency (MHz)	$\mathcal{E}_r'$	σ(S/m)	a ( <b>mm</b> )	$\delta$ (mm)	$\mathcal{E}_r'$	Thickness (mm)
1800-2000	40	1.40	109.2	24.15	4.8	19.4
2450	39	1.80	109.2	18.59	5.7	12.6
5200	36	4.70	40.0	6.60	6.0	5.3
5800	35.3	5.30	40.0	6.00	6.0	4.4

NOTES — (1) Permittivity and thickness of the dielectric separator may vary from the values shown to accommodate commercially available materials. Changes of the order of  $\pm$  10% should not produce substantial performance degradation.

(2) By convention, the length of the cross-section short edge is one-half that of the long edge, i.e., b = a/2.

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CM-16, 2.4 GHz Cordeless Phone M/N: CM-16 FCC ID: SDVCM16

#### 5.6.4. **Thermal Transfer Calibration**

#### 5.6.4.1. Measurement procedure

An RF transparent thermistor-based temperature probe and a isotropic E-field probe are placed side-by-side in a planar phantom while both are exposed to RF energy from a half wave dipole antenna located below the phantom The E-field probe and amplifiers were previously calibrated.

First, the location of the maximum E-field close to the phantom's bottom is determined as a function of power into the dipole

Then, the E-field probe is moved sideways so that the temperature probe, while affixed to the E-field probe is placed at the previous location of the E-field probe.

Finally, temperature changes for a certain amount of time (generally 10 to 30 seconds) exposures at the same RF power levels used for the E-field are recorded. Care is taken to allow cooling down to the original temperature and temperature stabilization between tests.

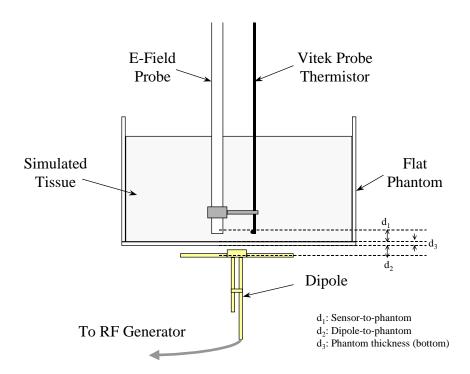


Figure 5.6.4.11 Flat Phantom, Thermistor and E-Field Probe

The following simple equation relates SAR to the initial temperature slope:

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$$SAR_{t} = \frac{c \cdot \Delta T}{\Delta t}$$
 (Eq. 5.6.4.11)

In (Eq. 5.6.4.11)  $\Delta t$  is the exposure time [sec], c is the specific heat capacity of the simulated tissue [J/Kg/ $^{\circ}$ C] and  $\Delta T$  is the temperature increase [ $^{\circ}$ C] due to the RF exposure. SAR is proportional to  $\Delta T/\Delta t$ , the initial rate of tissue heating, before thermal diffusion takes place.

From (Eq. 5.6.4.11) it is possible to quantify the electric field in the simulated tissue by equating the thermally-derived SAR to the E-field:

$$SAR = \frac{\left|E\right|^2 \cdot \sigma}{\rho}$$
 (Eq. 5.6.4.12)

where  $\sigma$  is the simulated tissue conductivity [S/m] and  $\rho$  its mass density [Kg/m³]; The actual mass density of the simulated tissue is required during the thermal transfer calibration, while mass density of 1,000 [Kg/m³] is conventionally chosen during the SAR measurements.

## 5.6.4.2. Determination of Conversion Factor ( $\gamma$ ) in the simulated tissue

The sensitivity of the probe in the dielectric media compared to its sensitivity in the air, is different. Conversion Factor  $(\gamma)$  is defined to determine the degree of the enhancement of sensitivity in the different dielectric media and relate it to its sensitivity in the air.

$$PO_{tot\ tissue} \equiv PO_{tot\ air} \times \gamma$$

Thus,

$$\left|E_{tissue}\right|^{2} = \frac{PO_{tot\_tissue}}{\eta_{E2}} \times \frac{1}{\gamma}$$
 , and  $SAR_{tissue} = \frac{\sigma \times \frac{PO_{tot\_tissue}}{\eta_{E2}} \times \frac{1}{\gamma}}{\rho}$ 

where,

 $\begin{array}{ll} \left|E_{tissue}\right|^2 & RMS \ E\text{-field level } [(V/m)^2] \ induced \ within \ the \ exposed \ tissue \\ PO_{tot\_tissue} & Probe \ voltage \ output \ measured \ in \ the \ simulated \ tissue \ [mV] \\ PO_{tot\_air} & Probe \ voltage \ output \ measured \ in \ the \ air \ (Z_{air} = 377[\Omega]) \ [mV] \\ \eta_{E2} & Sensor \ Factor \ to \ the \ |E|^2, \ an \ arbitrary \ value \ 10.8/3,770 \ [mV/(V/m)^2] \\ \gamma & Conversion \ factor; \ ratio \ of \ sensor \ response \ in \ air \ to \ response \ in \ the \ dielectric \ media \\ \end{array}$ 

The conversion factor ( $\gamma$ ) can be used to scale the E-field in terms of the thermally-derived SAR. It is the quotient of SAR<sub>t</sub>, the SAR determined from temperature measurements in the flat phantom, and PO<sub>tot\_tissue</sub>, the E-field prove output voltage obtained at the same location in the phantom

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$$SAR_{t} = SAR_{tissue}$$

$$\frac{c \cdot \Delta T}{\Delta t} = \frac{\sigma_{@ cal} \times |E_{tissue}|^{2}}{\rho}$$

$$= \frac{\sigma_{@ cal} \times \frac{PO_{tot\_tissue}}{\eta_{E2}} \times \frac{1}{\gamma}}{\rho}$$

Thus,

$$\gamma = \frac{\sigma_{\text{@ cal}}}{\eta_{E2} \times \rho} \times \frac{PO_{tot\_tissue}}{SAR_t} = \frac{\sigma_{\text{@ cal}} \times 3,770}{10.8 \times c \times \rho} \times \frac{PO_{tot\_tissue}}{\Delta T/\Delta t}$$
(Eq. 5.6.4.21)

where,

 $\begin{array}{lll} \gamma & & Conversion factor; ratio of sensor response in air to response in the dielectric media \\ SAR_t & Thermally-derived & SAR [W/Kg] (Eq. 1) \\ |E_{tissue}|^2 & RMS E-field level [(V/m)^2] induced within the exposed tissue \\ PO_{tot_tissue} & Probe voltage output measured in the simulated tissue [mV] \\ \eta_{E2} & Sensor Factor to the |E|^2, an intermediately constant, <math>10.8/3,770 \ [mV/(V/m)^2] \ c & Specific heat capacity of the simulated tissue [J/Kg/°C] \\ \sigma_{@cal} & Conductivity of the simulated tissue during the calibration procedure [S/m] \\ \rho & Actual mass density of the simulated tissue [Kg/m^3] \\ \Delta T/\Delta t & Initial rate of tissue heating, before thermal diffusion takes place [°C /sec] \\ \end{array}$ 

The temperature E-field correlation is illustrated below (for simulated brain tissue) for an example in which the thermal quantities were,

RF power input = 0.5 [W]  $\Delta T = 0.0163 \ [^{\circ}C] \ (from \ thermistor-base \ temperature \ probe)$   $\sigma_{@cal} = 0.97 \ [S/m]$   $\rho = 1,200 \ [Kg/m^3]$   $c = 2,700 \ [J/Kg/^{\circ}C]$   $\Delta t = 30 \ [sec]$ 

The resulting SAR<sub>t</sub> was (Eq. 5.6.4.11)

$$SAR_t = \frac{2,700 \times 0.0163}{30} = 1.467 \text{ [W/Kg]}$$

In this case the output of the E-field probe when at the same position as the thermistor probe was

$$PO_{tot\ tissue} = 28.5 [mV]$$

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The calculation of conversion factor ( $\gamma$ ) from (Eq. 5.6.4.21) follows:

$$\gamma = \frac{0.97}{\frac{10.8}{3.770} \times 1,200} \times \frac{28.5}{1.467} = 5.482$$

#### 5.6.5. Data Acquisition Methodology

#### 5.6.5.1. E-Field Measurement

The probe calibration must be current before starting measurements. Instrumentation amplifier batteries must be charged. This can be monitored by observing DC offset voltages. A daily log of the DC offset voltages should be kept for this purpose.

Measurements in the phantom are automatically calculated for each location by summation of the three dipole outputs. Because each dipole produces an output voltage proportional to the square of the electric field component along the dipole, the sum of dipole voltages represents the RMS values for the total electric field. Thus, taking into consideration the amplifier settings and the DC offset voltages, the total electric field strength at a measurement location is as follows. See Appendix C. PO<sub>tot</sub> is labeled by the software as measure of values (voltages). The SAR for calculations that are derived from the measure of values are discussed below.

At each measurement point, the program records the output of the three channels:

$$|E_i|^2[(V/m)^2] = \frac{PO_i[mV]}{\eta_{E2}[mV/(V/m)^2]} = \frac{U_{L,i} \times AS_i}{\eta_{E2}[mV/(V/m)^2]}$$

$$|E|^{2}[(V/m)^{2}] = \sum_{i}^{3}|E_{i}|^{2} = \frac{1}{\eta_{E2}} \times \sum_{i}^{3}PO_{i} = \frac{PO_{tot}}{\eta_{E2}} = \frac{1}{\eta_{E2}} \times \sum_{i}^{3}(U_{L,i} \times AS_{i})$$

Where,

U<sub>L,i</sub> Linearized probe output for channel i

AS<sub>i</sub> Amplifier Setting for channel i

 $\eta_{E2}$  Sensor Factor to the uniform  $|E|^2$ , an arbitrary value  $10.8/3,770 \text{ [mV/(V/m)}^2]$ 

PO<sub>i</sub> Probe output [mV] of channel i at a measurement point

PO<sub>tot</sub> Total Probe output [mV] at a measurement point

## 5.6.5.2. Sensitivity( $\zeta$ ) of probe in the simulated tissue

The sensitivy( $\zeta$ ) of the probe in the simulated tissue is rendered in terms of Sensor Enhancement Factor in the simulated tissue.

$$\zeta = \frac{\sigma_{\text{@ meas}}}{\eta_{E2} \times \rho \times \gamma} = \frac{\sigma_{\text{@ meas}}}{\frac{10.8}{3.770} \times 1,000 \times \gamma} = \frac{3,770 \times \sigma_{\text{@ meas}}}{10,800 \times \gamma}$$
(Eq. 5.6.5.21)

Where.

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ζ	Sensitivity of the probe in the simulated tissue [W/Kg/mV]
γ	Conversion factor; ratio of sensor response in air to response in the dielectric media
$\eta_{E2}$	Sensor Factor to the $ E ^2$ , an arbitrary value 10.8/3,770 [mV/(V/m) <sup>2</sup> ]
$\sigma_{\text{@meas}}$	Conductivity of the simulated tissue during the measurement [S/m]
ρ	Mass density of the simulated tissue [Kg/m³]; 1,000 [Kg/m3] is conventionally chosen.

Therefore, SAR can be yielded from

$$SAR = \zeta \times PO_{tot tissue}$$
 (Eq. 5.6.5.22)

Where,

 $\begin{array}{ll} \zeta & & \text{Sensitivity of the probe in the simulated tissue } [W/Kg/mV] \\ PO_{tot \ tissue} & & \text{Probe voltage output measured in the simulated tissue } [mV] \end{array}$ 

To continue the example illustrated above,

$$\sigma_{\text{@meas}} = 0.99 \text{ [S/m]}$$

$$PO_{\text{tot\_tissue}} = 11.5 \text{ [mV]}$$

$$\zeta = \frac{3,770 \times \sigma_{\text{@meas}}}{10,800 \times \eta} = \frac{3,770 \times 0.99}{10,800 \times 5.482} = 0.063 \text{ [W/Kg/mV]}$$

$$SAR = \zeta \times PO_{\text{tot\_tissue}} = 0.063 \times 11.5 = 0.725 \text{ [W/Kg]}$$

#### 5.6.5.3. SAR Measurement

The goal of the measurement process is to scan the phantom over a selected area in order to find the region of highest levels of RF energy and then to obtain a single value for the peak spatial-average of SAR over a volume that would contain one gram (in the shape of a cube) of biological tissue. The test procedure, of course, measures SAR in the simulated tissue.

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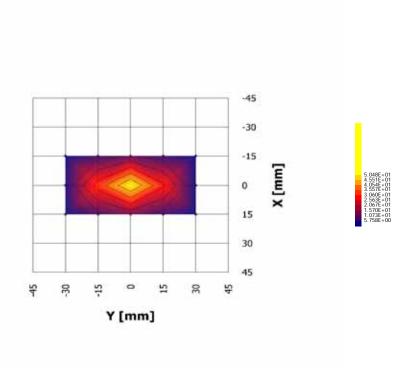


Figure 5.6.5.3 Area scan

The software request the user to move the probe to locations at two extreme corners of a rectangle that encloses the area to be scanned. An arbitrary origin and the spatial resolution for the scan are also specified. Under program control, the scan is performed automatically by the robot-guided probe.

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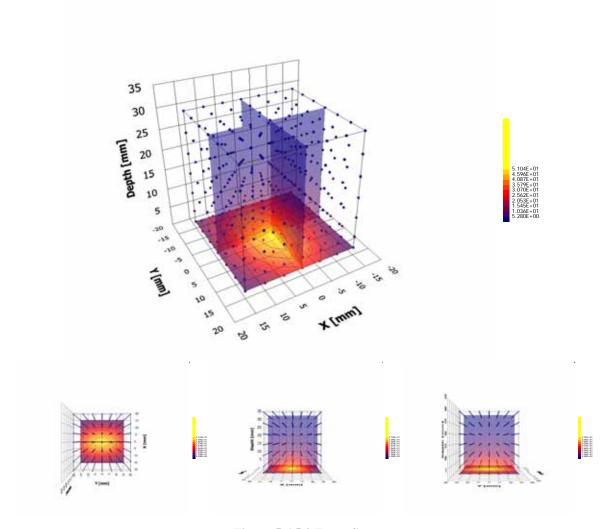


Figure 5.6.5.3 Zoom Scan

The fine resolution volume scan region is centered at the peak SAR locations determined by the interpolated (cubic spline) data from the area scan measurements. The number of measurement point required in a zoom scan is defined to provide an accurate one-gram averaged SAR in terms of both the number of points  $(PT_X \times PT_Y \times PT_Z)$  and the size  $(SZ_X[mm] \times SZ_Y[mm] \times SZ_Z[mm])$  of the cubic. For one-gram SAR,  $(5 \times 5 \times 7)$  and  $(28[mm] \times 28[mm] \times 30[mm])$  is preferred to select below 1 GHz. The zoom scan region extends in each direction for at least 1.5 times the linear dimensions of 1- or 10-gram cube of tissue from each peak. The zoom scan spatial resolution is interpolated down to SAR values on a 1mm grid by using the tri-linear interpolation algorithm.

The peak field values near the surface of a homogeneous phantom are usually not measurable because the sensors in a field probe are located at 2-4 mm behind the tip of the probe and the measurement point is defined at the geometric center of the sensors where the calibration is defined. These SAR values are computed by extrapolating the closest measured points to the surface of the phantom to determine the highest one-gram averaged SAR. The extrapolation

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coefficients are determined with a multi-order curve-fitting algorithm. Generally the 4-th order polynomial least-square fit is sufficient to extrapolate to the surface if the number of the valid measurements, that are non-zero, along the probe axis is greater than 4.

The interpolated and extrapolated SAR values from the zoom scan measurements are integrated in the shape of 1- or 10-gram cube then traversed to determine the highest peak spatial-average SAR in the zoom scan region.

This peak spatial-averaged SAR is reported as SAR [W/kg] for compliance.

#### 5.6.5.4. Data Extrapolation and boundary effect

The distance from the center of the sensor (diode) to the end of the protective tube is called the 'probe offset' or 'sensor offset'. To compensate we use a multi-order polynomial least-square curve fitting to obtain the peak surface value from the voltages measured at the distance from the inner surface of the phantom. The field is measured as close as possible to the phantom's surface and every pre-defined separation distance (1 [mm] to 5 [mm]) along the probe axis (z) for a distance of at least 50 mm until they are not measurable. The appropriate curve is obtained from all the points measured and used to define an exponential decay of the energy density versus depth.

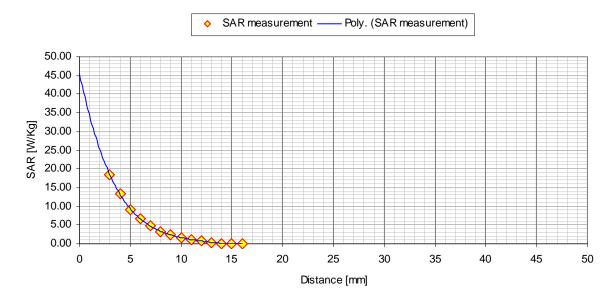


Figure 5.6.5.4 Exponential decay of the energy density versus depth

In general, 4-th order polynomial curve fitting will be applied for extrapolation to the surface, but whenever the 4-th order polynomial is not application from the lack of the input data the lower order polynomial curve fitting will be applied instead.

Boundary effects arise when the tip of an electric field probe approaches the interface between two dielectric media. Under these conditions, the external field is strongly perturbed by the superposition of a scattered field from the probe. The effect of the boundary on the peak spatial-average SAR values strongly depends on the probe dimensions, especially

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the diameter of the tip of the probe. It is known that the error due to boundary effects is very small if the distance between the probe tip and the surface is greater than half the probe diameter. Therefore the first one or two measurements at the vicinity to the phantom surface are excluded for evaluating the exponential decay curve in order to compensate for the boundary effect.

#### **5.6.6.** Determining the Heat Capacity of Simulated Tissue

#### 5.6.6.1. Instruments and Materials

- Calibrated differential thermometer (Vitek or BAT-8 or equivalent)
- Two identical 500 ml containers
- A thermally insulated vessel (thick styrofoam, with a form fitting hole for one container)
- Hot and cold tap water
- Solution under test
- Hot plate
- Temperature vs. time (chart recorder, or data loger)

#### 5.6.6.2. Method

Heat can be propagated by conduction, convection and radiation. In the case of liquids heated from below, gravity convection is the main and predominant heating mechanism of the fluid mass.

Obtain two containers that can be rapidly heated (e.g. glass or suitable plastic). Fill one container with 250 ml of water, the other with the same mass of simulated tissue. The initial temperature of the water should be the same as that of the simulated tissue ( $\pm 1^{\circ}$ C). Since we are dealing with heating by electromagnetic sources at ambient temperature, it is essential that we eliminate the chance of any direct infrared heating of the temperature sensor. To ensure this, position the tip of the sensor 2 mm from the bottom of the center of the container. Turn on the heat source and wait at least 5 minutes for its temperature to stabilize. Record the initial temperature of the water. Place the container of water 5 mm above the center of the hot plate and monitor the temperature increase.

After 30 seconds of heating, the water temperature should have increased by at least 5 °C. Record the time and temperature. Remove the container from the heat source and place it in the thermally insulated vessel. Stir the liquid thoroughly and record the steady state temperature 1-2 minutes after stirring.

Repeat the above procedure using the container of simulated tissue. Ensure that the container is placed on the same area of the hot plate, is heated for the identical length of time, and the steady state temperature is recorded after the identical time interval.

Since the heat capacity of water is  $C_w = 1,000 \text{ [cal/Kg/°C]}$  or 4,189 [J/Kg/°C] with excellent approximation (~1%) in the temperature range of interest, the heat capacity ( $C_s$ ) of the solution is given by:

$$C_s = C_w \cdot \frac{\Delta T_w}{\Delta T_s}$$

where  $\Delta T_w$  is the temperature increase of water and  $\Delta T_s$  the temperature increase of the solution. The ration of the values,  $\Delta T_w/\Delta T_s$ , should be the same (within the sensitivity of the thermometer) at the end of the heating and stirring. This ensures that the liquids have been uniformly heated.

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#### 5.6.6.3. Rationale

$$C \cdot \Delta T = Heat \ Flow \cdot Time = Total \ Heating \ Energy$$

If the heat flow, sample mass, and absorption (heat transfer) are the same for both liquids, then:

$$C_{w} \cdot \Delta T_{w} = C_{s} \cdot \Delta T_{s}$$

The heat flow and total heating are kept constant by using the same source for the same amount of time. If the heat transfer mechanisms for the woe liquids are about the same, with insignificant differences in convective and conductive characteristics, then any differences in temperature increase are a direct measure of the specific heat capacity, C.

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# 5.7. SAR MEASUREMENT SYSTEM VERIFICATION

# 5.7.1. Standard Source

A half-wave dipole is positioned below the bottom of the phantom and centered with its axis parallel to the longest side of the phantom. The distance between the liquid filled phantom bottom surface and the center of the dipole axis, *s*, is chosen as specified IEEE 1528 at the specific test frequency (i.e. 15 mm at 835 MHz). A low loss and low dielectric constant spacer is used to establish the correct distance between the top surface of the dipole and the bottom surface of the phantom.



# 5.7.2. Standard Source Input Power Measurement

The system validation is performed as shown below or in Figure 7.1 in IEEE 1528.

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First the power meter PM1 (including attenuator Att1) is connected to the cable to measure the forward power at the location of the dipole connector (X). The signal generator is adjusted for the desired forward power at the dipole connector (taking into account the attenuation of Att1) as read by power meter PM2. After connecting the cable to the dipole, the signal generator is readjusted for the same reading at power meter PM2. If the signal generator does not allow adjustment in 0.01dB steps, the remaining difference at PM2 must be taken into consideration. PM3 records the reflected power from the dipole to ensure that the value is not changed from the previous value. The reflected power was verified to be at least 20dB below the forward power.

#### 5.7.3. **System Validation Procedure**

A complete 1g-averaged SAR measurement is performed. The measured 1g-averaged SAR value is normalized to a forward power of 1W to a half-wave dipole and compared with the reference SAR value for the reference dipole and flat phantom shown in columns 2 and 3 of Table 7.1 in IEEE 1528.

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## 5.8. POWER MEASUREMENT

Whenever possible, a conducted power measurement is performed. To accomplish this, we utilize a fully charged battery, a calibrated power meter and a cable adapter provided by the manufacturer. The data of the cable and related circuit losses are also provided by the manufacturer. The power measurement is then performed across the operational band and the channel with the highest output power is recorded.

Power measurement is performed before and after the SAR to verify if the battery was delivering full power at the time of testing. A difference in output power would determine a need for battery replacement and to repeat the SAR test.

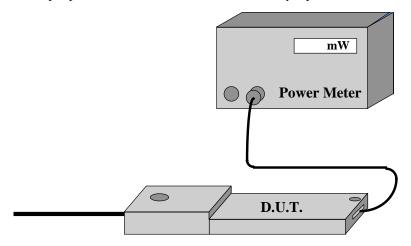


Figure 5.8. 1.Measured Power + Cable and Switching Mechanism Loss

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#### 5.9. POSITIONING OF D.U.T.

The clear SAM phantom shell have been previously marked with a highly visible grid with a defined centre line, so it can easily be seen through the liquid simulated tissue. In the case of testing a cellular phone, this line is connecting the ear channel with the corner of the lips. The D.U.T. is then placed by centering the speaker with the ear channel and the center of the radio width with the corner of the mouth.

For HAND HELD devices (push-to-talk), or any other type of wireless transmitters postioned in front of the face, the D.U.T. will be positioned 2.5cm distance from a flat phantom to simulate the frontal facial position in use. All bodyworn operating configurations are tested using a flat phantom. The length and width of the phantom is at least twice the corresponding dimensions of the test device, including its antenna.

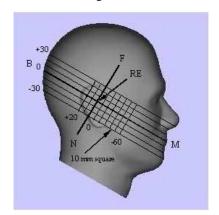


Figure 5.9. 1. Side view of the phantom showing relevant marking

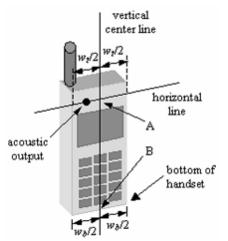


Figure 5.9. 2. Handset vertical and horizontal reference lines – fixed case

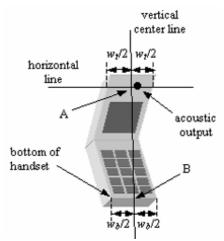


Figure 5.9. . Handset vertical and horizontal reference lines – "clam-shell"

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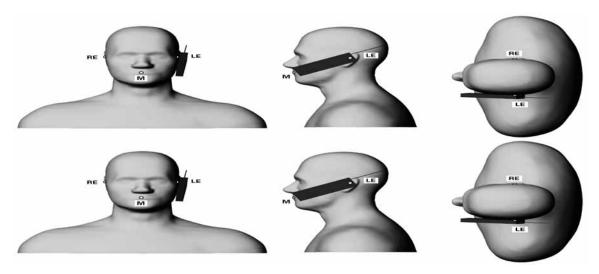


Figure 5.9. 4. Phone position 1, "cheek" or "touch" position. The reference points for the right ear (RE), left ear (LE) and mouth (M), which define the reference plane for phone positioning, are indicated. The shoulders are shown for illustration purposes only.

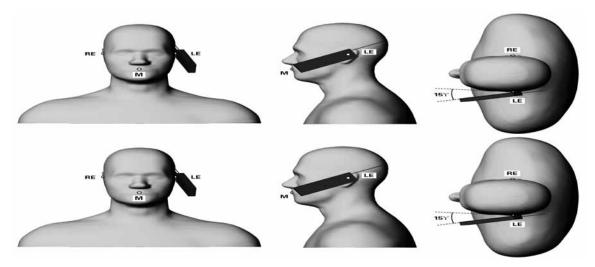


Figure 5.9. 5. Phone position 2, "tilted position." The reference points for the right ear (RE), left ear (LE) and mouth (M), which define the reference plane for phone positioning, are indicated. The shoulders are shown for illustration purposes only.

The handset holder is mainly made of PVC and contains no metallic component at all in order to minimize field perturbation. Velcro and elastic band were used to attach the D.U.T. on the plate of handset holder.

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CM-16, 2.4 GHz Cordeless Phone M/N: CM-16 FCC ID: SDVCM16

## 5.10. SAR MEASUREMENT UNCERTAINTY

This uncertainty analysis covers the 3D-EMC Laboratory test procedure for Specific Absorption Rate (SAR) associated with wireless telephones and similar devices.

#### **Standards Covered Are:**

WGMTE 96/4 - Secretary SC211/B

FCC 96-326, ET Docket No. 93-62

Industry Canada RSS 102

ACA Radiocommunications (Electromagnetic Radiation - Human Exposure) Amendment Standard 2000 (No. 1)

The laboratory test procedure, and this uncertainty analysis, may be used to cover all standards above. It is based on test equipment and procedures specified by 3D-EMC Laboratories, Inc. located in Ft. Lauderdale, Florida.

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IEEE C95.1-1991, FCC OET Bulletin 65 (Supplement C), Industry Canada RSS-102(Issue 1) and ACA Radiocommunications (Electromagnetic Radiation – Human Exposure) Amendment Standard 2000 (No. 1)

CM-16, 2.4 GHz Cordeless Phone M/N: CM-16 FCC ID: SDVCM16

# **5.10.1.** Measurement Uncertainty

## 5.10.1.1. Measurement Uncertainty evaluation for handset SAR test

							h =	i =	
a	b	с	d	e = f(d,k)	F	g	cxf/e	c x g / e	k
Uncertainty		Tol.	Prob.		$c_i$	$c_i$	1-g	10-g	
Component		(± %)	Dist.		(1-g)	(10-g)	$\boldsymbol{u}_i$	$u_i$	
•	Sec.			Div.	, 0,	, ,	(±%)	(±%)	$v_i$
Measurement System									
Probe Calibration	E1.1	3.0	N	1	1	1	3.0	3.0	∞
Axial Isotropy	E1.2	5.0	R	√3	0.7	0.7	2.0	2.0	∞
Hemispherical Isotropy	E1.2	8.0	R	√3	1	1	4.6	4.6	×
Boundary Effect	E1.3	10.0	R	√3	1	1	5.8	5.8	∞
Linearity	E1.4	4.2	R	√3	1	1	2.4	2.4	∞
System Detection Limits	E1.5	2.0	R	√3	1	1	1.2	1.2	∞
Readout Electronics	E1.6	1.0	N	1	1	1	1.0	1.0	∞
Response Time	E1.7	1.5	R	√3	1	1	0.9	0.9	∞
Integration Time	E1.8	2.0	R	√3	1	1	1.2	1.2	∞
RF Ambient Conditions	E5.1	3.0	R	√3	1	1	1.7	1.7	∞
Probe Positioner Mechanical Tolerance	E5.2	1.0	R	√3	1	1	0.6	0.6	$\infty$
Probe Positioning with respect to Phantom Shell	E5.3	3.0	R	√3	1	1	1.7	1.7	∞
Extrapolation, interpolation and Integration Algorithms for Max. SAR Evaluation	E4.2	3.5	R	√3	1	1	2.0	2.0	∞
Test sample Related									
Test Sample Positioning	E3.2.1	7.5	N	1	1	1	7.5	7.5	11
Device Holder Uncertainty	E3.1.1	6.5	N	1	1	1	6.5	6.5	8
Output Power Variation - SAR drift measurement	5.6.2	5.0	R	√3	1	1	2.9	2.9	∞
Phantom and Tissue Parameters									
Phantom Uncertainty (shape and thickness tolerances)	E2.1	4.0	R	√3	1	1	2.3	2.3	∞
Liquid Conductivity Target - tolerance	E2.2	5.0	R	√3	0.7	0.5	2.0	1.4	× ×
Liquid Conductivity - measurement uncertainty	E2.2	4.0	R	√3	0.7	0.5	1.6	1.2	× ×
Liquid Permittivity Target tolerance	E2.2	5.0	R	√3	0.6	0.5	1.7	1.4	oc
Liquid Permittivity - measurement uncertainty	E2.2	4.0	R	√3	0.6	0.5	1.4	1.2	oc
Combined Standard Uncertainty			RSS				14.3	14.2	
Expanded Uncertainty							• • •	•••	
(95% confidence interval)							28.5	28.3	

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CM-16, 2.4 GHz Cordeless Phone M/N: CM-16

#### 5.10.1.2. Measurement Uncertainty for System Performance Check

							h =	i =	
а	b	С	d	e = f(d,k)	f	g	cxf/e	cxg/e	k
Uncertainty		Tol.	Prob.		$c_i$	$c_i$	1-g	10-g	$v_i$
		(1.0/)	D: 4		(1)	(10 )			
Component	Sec.	(± %)	Dist.	Div.	(1-g)	(10-g)	<i>u<sub>i</sub></i> (±%)	<i>u<sub>i</sub></i> (±%)	or v <sub>eff</sub>
Measurement System				Div.			(± /0)	(± /0)	
Probe Calibration	E1.1	3.0	N	1	1	1	3.0	3.0	oc
Axial Isotropy	E1.2	5.0	R	√3	0.7	0.7	2.0	2.0	- oo
Hemispherical Isotropy	E1.2	8.0	R	√3	1	1	4.6	4.6	8
Boundary Effect	E1.3	10.0	R	√3	1	1	5.8	5.8	oc .
Linearity	E1.4	4.2	R	√3	1	1	2.4	2.4	∞
System Detection Limits	E1.5	2.0	R	√3	1	1	1.2	1.2	oc .
Readout Electronics	E1.6	1.0	N	1	1	1	1.0	1.0	8
Response Time	E1.7	1.5	R	√3	1	1	0.9	0.9	8
Integration Time	E1.8	2.0	R	√3	1	1	1.2	1.2	8
RF Ambient Conditions	E5.1	3.0	R	√3	1	1	1.7	1.7	8
Probe Positioner Mechanical Tolerance	E5.2	0.4	R	√3	1	1	0.2	0.2	8
Probe Positioning with respect to Phantom Shell	E5.3	3.0	R	√3	1	1	1.7	1.7	∞
Extrapolation, interpolation and Integration Algorithms for Max. SAR Evaluation	E4.2	3.5	R	√3	1	1	2.0	2.0	∞
Dipole									
Dipole Axis to Liquid Distance	7, X3.2	2.0	R	√3	1	1	1.2	1.2	oc
Input Power and SAR Drift Measurement	7, 5.6.2	3.0	R	√3	1	1	1.7	1.7	∞
Phantom and Tissue Parameters									
Phantom Uncertainty - shell thickness tolerance	E2.1	4.0	R	√3	1	1	2.3	2.3	∞
Liquid Conductivity – deviation from target values	E2.2	5.0	R	√3	0.7	0.5	2.0	1.4	$\infty$
Liquid Conductivity - measurement uncertainty	E2.2	4.0	R	√3	0.7	0.5	1.6	1.2	∞
Liquid Permittivity – deviation from target values	E2.2	5.0	R	√3	0.6	0.5	1.7	1.4	∞
Liquid Permittivity - measurement uncertainty	E2.2	4.0	R	√3	0.6	0.5	1.4	1.2	∞
Combined Standard Uncertainty			RSS				10.0	9.9	
Expanded Uncertainty									
(95% confidence interval)							20.1	19.8	

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# **EXHIBIT 6. SAR PRESCANS**

# **6.1.1.** Body Configuration

# 6.1.1.1. Test configurations used

Body-worn operating configurations should be tested with the belt-clips and holsters attached to the device and positioned against a flat phantom in normal use configurations. Devices with a headset output should be tested with a headset connected to the device. The D.U.T. was placed against the phantom and tested in its appropriate holster as would normally be used by the end user. If the SAR measured at the middle channel for each test is at least 3.0 dB lower than the SAR limit, testing at the high and low channels is optional for such test configuration(s).

If the transmission band of the test device is less than 10 MHz, testing at the high and low frequency channels is optional.

When multiple accessories that do not contain metallic components are supplied with the device, the device may be tested with only the accessory that dictates the closest spacing to the body. When multiple accessories that contain metallic components are supplied with the device, the device must be tested with each accessory that contains a unique metallic component. If multiple accessories share an identical metallic component (e.g., the same metallic belt-clip used with different holsters with no other metallic components), only the accessory that dictates the closest spacing to the body must be tested.

Body-worn accessories may not always be supplied or available as options for some devices that are intended to be authorized for body-worn use. A separation distance of 1.5 cm between the back of the device and a flat phantom is recommended for testing body-worn SAR compliance under such circumstances. Other separation distances may be used, but they should not exceed 2.5 cm. In these cases, the device may use body-worn accessories that provide a separation distance greater than that tested for the device provided however that the accessory contains no metallic components..

#### 6.1.1.2. Equipment permutation investigated for each orientation

N/A

#### 6.1.1.3. Comments on non-tested configurations

N/A

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#### 6.2. RECOMMENDED CAUTION STATEMENTS TO BE INCLUDED IN USERS MANUAL

In order for users to be aware of the body-worn operating requirements for meeting RF exposure compliance, operating instructions and caution statements should be included in the manual. The information should allow users to make informed decisions on the type of body-worn accessories and operating configurations that are appropriate for the device. The following are *examples* of typical statements that provide end-users with the necessary information about body-worn accessories:

Example 1. For a product that has the potential to be used in a body worn configuration and has been tested and certified with a specific accessory device(s):

"For body worn operation, this phone has been tested and meets the FCC RF exposure guidelines when used with the (*manufacturer name*) accessories supplied or designated for this product. Use of other accessories may not ensure compliance with FCC RF exposure guidelines."

Example 2. For a product that has the potential to be used in a body worn configuration and has not been certified with a specific accessory device(s):

"For body worn operation, this phone has been tested and meets FCC RF exposure guidelines when used with an accessory that contains no metal and that positions the handset a minimum of (specified distance) from the body. Use of other accessories may not ensure compliance with FCC RF exposure guidelines."

Example 3. For a product that has the potential to be used in a body worn configuration with future manufacturer designed accessories:

"For body worn operation, this phone has been tested and meets the FCC RF exposure guidelines when used with a (manufacturer name) accessory designated for this product or when used with an accessory that contains no metal and that positions the handset a minimum of (specified distance) from the body."

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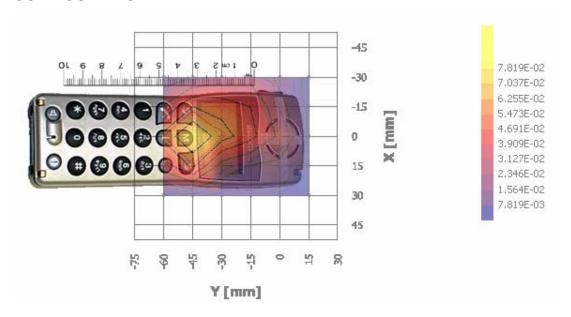
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# **EXHIBIT 7. SAR MEASUREMENT**

# 7.1. HEAD CONFIGURATION



The center of the ear pieces was set as the origin point (0, 0) of the plot for all the measurements in head configuration.

# **7.1.1.** Left Head\*

#	Configuration	Antenna Position	Frequency [MHz]	Channel	SAR <sub>local</sub> Before [W/Kg]	SAR <sub>local</sub> After [W/Kg]	MAX SAR <sub>1g</sub> [W/Kg]
*	General population/Uncontrolled Exposure Category I	Limit					1.6
01	Left Head		2401.056	Lowest			-
02	Cheek Position	Integrated	2440.800	Middle	0.01	0.01	0.04
03			2482.272	Highest			-
04	Left Head		2401.056	Lowest			-
05	Tilt Position	Integrated	2440.800	Middle	0.01	0.01	0.03
06			2482.272	Highest			-

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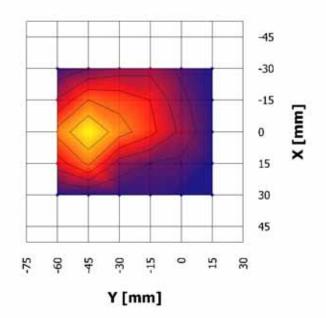
<sup>\*</sup> If the SAR measured at the middle channel for each test configuration is at least 3.0 dB lower than the SAR limit, testing at the high and low channels is optional for such test configuration(s).

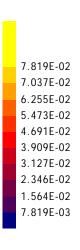
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CM-16, 2.4 GHz Cordeless Phone M/N: CM-16

#### 7.1.1.1. Left Cheek Position; #02

Test date [MM/DD/YYYY]	08/25/2004
Test by	JaeWook Choi
Room temperature [°C]	24
Room humidity [%]	40
Simulated tissue temperature [°C]	24
Separation distance, d [mm]	0
Test frequency [MHz]	2440.800
E-field Probe	M/N: ET20, S/N:03JUN-0028, Sensor Offset: 2.0 mm
Sensor Factor $(\eta_{Pd})_{[mV/(mW/cm)]}^2$	10.8
Amplifier Settings (AS <sub>1</sub> , AS <sub>2</sub> , AS <sub>3</sub> )	0.0055211224, 0.0059622436, 0.0056122209
Tissue Type	Brain
Measured conductivity [S/m]	1.78 (-1.0 %)
Measured dielectric constant	38.2 (-2.6 %)
Conversion Factor (γ)	4.539
Sensitivity (ζ) [W/Kg/mV]	1.369E-01
Measurement Area Specification (X × Y )	$75 _{\text{mm}} \times 60 _{\text{mm}}$ ; Resolution: $15 _{\text{mm}} \times 15 _{\text{mm}}$
Measurement Volume Specification $(X \times Y \times Z)$	$7_{\text{pts}} \times 7_{\text{pts}} \times 7_{\text{pts}}$ , $30_{\text{mm}} \times 30_{\text{mm}} \times 30_{\text{mm}}$ ; Resolution: $5_{\text{mm}} \times 5_{\text{mm}} \times 5_{\text{mm}}$
SAR <sub>1g</sub> [W/Kg]	0.04





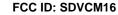
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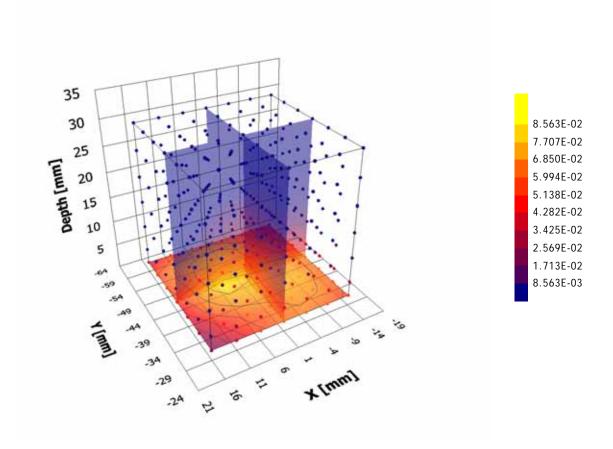
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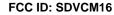
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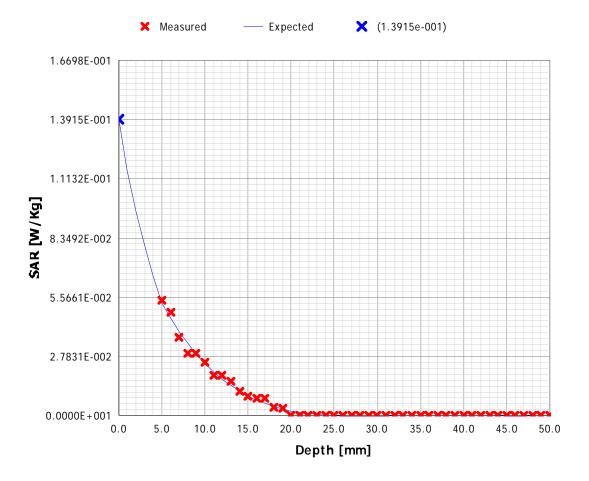
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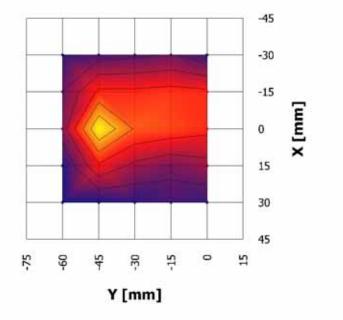
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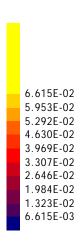
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CM-16, 2.4 GHz Cordeless Phone M/N: CM-16

#### 7.1.1.2. Left Tilt Position; #05

Test date [MM/DD/YYYY]	08/25/2004
Test by	JaeWook Choi
Room temperature [°C]	24
Room humidity [%]	40
Simulated tissue temperature [°C]	24
Separation distance, d [mm]	0
Test frequency [MHz]	2440.800
E-field Probe	M/N: ET20, S/N:03JUN-0028, Sensor Offset: 2.0 mm
Sensor Factor $(\eta_{Pd})_{[mV/(mW/cm)]}^2$	10.8
Amplifier Settings (AS <sub>1</sub> , AS <sub>2</sub> , AS <sub>3</sub> )	0.0055211224, 0.0059622436, 0.0056122209
Tissue Type	Brain
Measured conductivity [S/m]	1.78 (-1.0 %)
Measured dielectric constant	38.2 (-2.6 %)
Conversion Factor (γ)	4.539
Sensitivity (ζ) [W/Kg/mV]	1.369E-01
Measurement Area Specification (X × Y )	$60_{\text{mm}} \times 60_{\text{mm}}$ ; Resolution: $15_{\text{mm}} \times 15_{\text{mm}}$
Measurement Volume Specification $(X \times Y \times Z)$	$7_{\text{pts}} \times 7_{\text{pts}} \times 7_{\text{pts}}$ , $30_{\text{mm}} \times 30_{\text{mm}} \times 30_{\text{mm}}$ ; Resolution: $5_{\text{mm}} \times 5_{\text{mm}} \times 5_{\text{mm}}$
SAR <sub>1g</sub> [W/Kg]	0.04





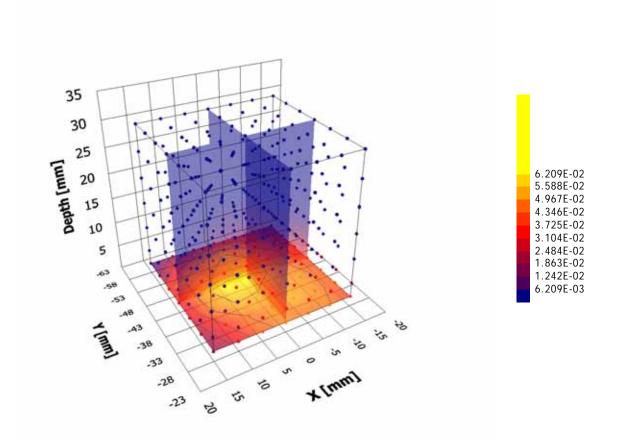
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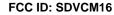
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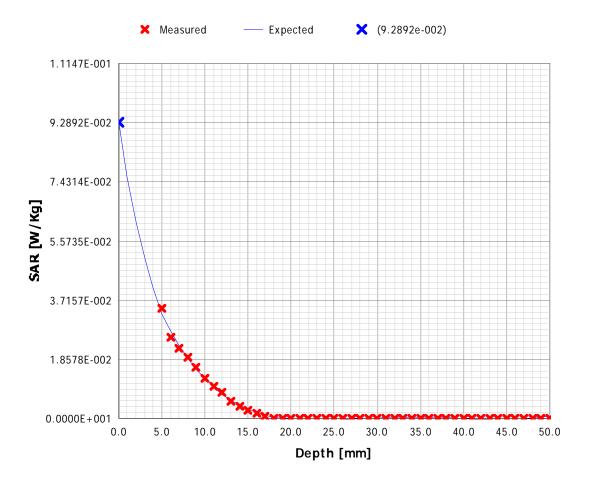
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# 7.1.2. Right Head\*

#	Configuration	Antenna Position	Frequency [MHz]	Channel	SAR <sub>local</sub> Before [W/Kg]	SAR <sub>local</sub> After [W/Kg]	MAX SAR <sub>1g</sub> [W/Kg]
*	General population/Uncontrolled Exposure Category I	Limit					1.6
07	Right Head		2401.056	Lowest			-
08	Cheek Position	Integrated	2440.800	Middle	0.02	0.02	0.05
09			2482.272	Highest			-
10	Right Head		2401.056	Lowest			-
11	Tilt Position	Integrated	2440.800	Middle	0.02	0.01	0.04
12			2482.272	Highest	· ·		-

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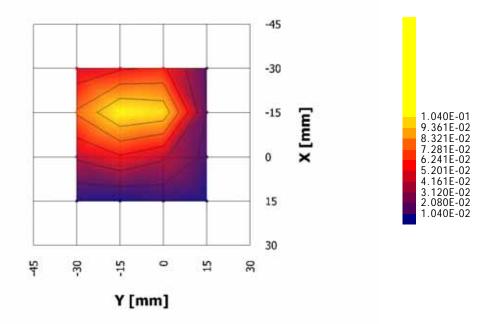
<sup>\*</sup> If the SAR measured at the middle channel for each test configuration is at least 3.0 dB lower than the SAR limit, testing at the high and low channels is optional for such test configuration(s).

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CM-16, 2.4 GHz Cordeless Phone M/N: CM-16

## 7.1.2.1. Right Cheek Position; #08

Test date [MM/DD/YYYY]	08/25/2004
Test by	JaeWook Choi
Room temperature [°C]	24
Room humidity [%]	40
Simulated tissue temperature [°C]	24
Separation distance, d [mm]	0
Test frequency [MHz]	2440.800
E-field Probe	M/N: ET20, S/N:03JUN-0028, Sensor Offset: 2.0 mm
Sensor Factor $(\eta_{Pd})_{[mV/(mW/cm)]}^2$	10.8
Amplifier Settings (AS <sub>1</sub> , AS <sub>2</sub> , AS <sub>3</sub> )	0.0055211224, 0.0059622436, 0.0056122209
Tissue Type	Brain
Measured conductivity [S/m]	1.78 (-1.0 %)
Measured dielectric constant	38.2 (-2.6 %)
Conversion Factor (γ)	4.539
Sensitivity (ζ) [W/Kg/mV]	1.369E-01
Measurement Area Specification (X × Y )	$45 _{\text{mm}} \times 45 _{\text{mm}}$ ; Resolution: $15 _{\text{mm}} \times 15 _{\text{mm}}$
Measurement Volume Specification $(X \times Y \times Z)$	$7_{\text{pts}} \times 7_{\text{pts}} \times 7_{\text{pts}}$ , $30_{\text{mm}} \times 30_{\text{mm}} \times 30_{\text{mm}}$ ; Resolution: $5_{\text{mm}} \times 5_{\text{mm}} \times 5_{\text{mm}}$
SAR <sub>1g</sub> [W/Kg]	0.05



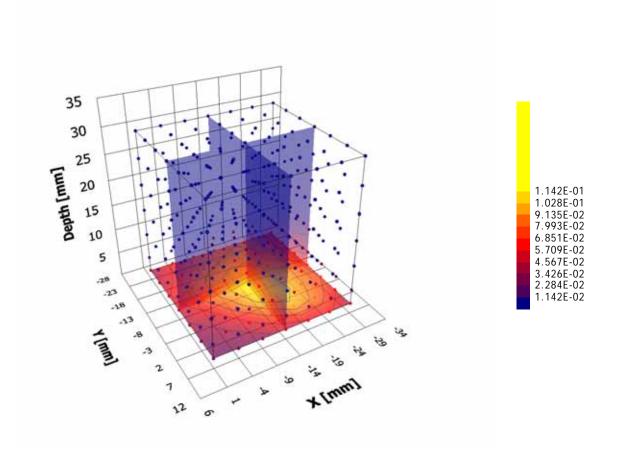
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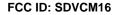
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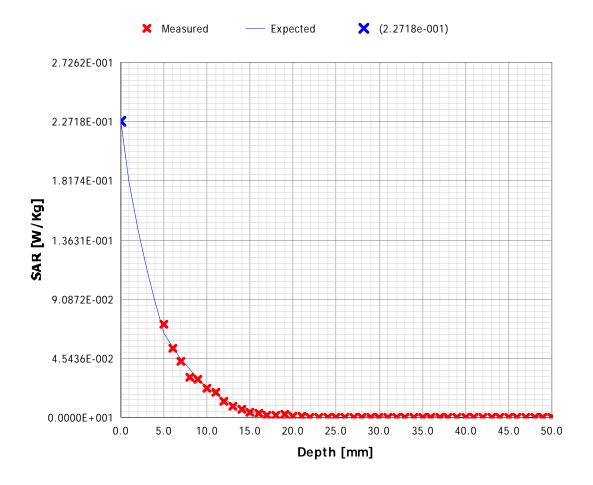
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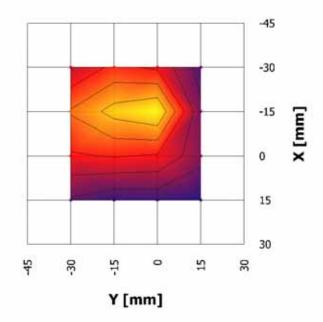
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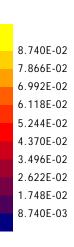
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CM-16, 2.4 GHz Cordeless Phone M/N: CM-16

## 7.1.2.2. Right Tilt Position; #11

Test date [MM/DD/YYYY]	08/25/2004
Test by	JaeWook Choi
Room temperature [°C]	24
Room humidity [%]	40
Simulated tissue temperature [°C]	24
Separation distance, d [mm]	0
Test frequency [MHz]	2440.800
E-field Probe	M/N: ET20, S/N:03JUN-0028, Sensor Offset: 2.0 mm
Sensor Factor $(\eta_{Pd})_{[mV/(mW/cm)]}^2$	10.8
Amplifier Settings (AS <sub>1</sub> , AS <sub>2</sub> , AS <sub>3</sub> )	0.0055211224, 0.0059622436, 0.0056122209
Tissue Type	Brain
Measured conductivity [S/m]	1.78 (-1.0 %)
Measured dielectric constant	38.2 (-2.6 %)
Conversion Factor (γ)	4.539
Sensitivity (ζ) [W/Kg/mV]	1.369E-01
Measurement Area Specification (X × Y )	$45 _{\text{mm}} \times 45 _{\text{mm}}$ Resolution: $15 _{\text{mm}} \times 15 _{\text{mm}}$
Measurement Volume Specification $(X \times Y \times Z)$	$7_{\rm pts} \times 7_{\rm pts} \times 7_{\rm pts}$ , $30_{\rm mm} \times 30_{\rm mm} \times 30_{\rm mm}$ ; Resolution: $5_{\rm mm} \times 5_{\rm mm} \times 5_{\rm mm}$
SAR <sub>1g</sub> [W/Kg]	0.04





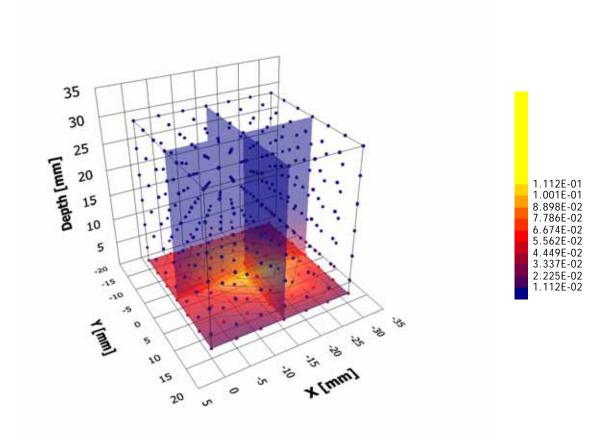
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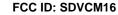
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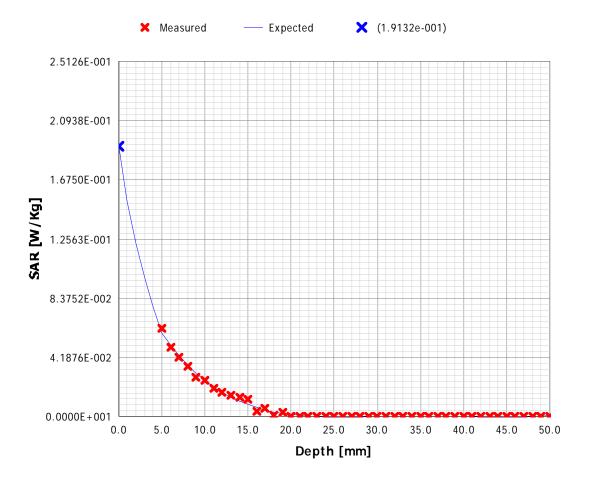
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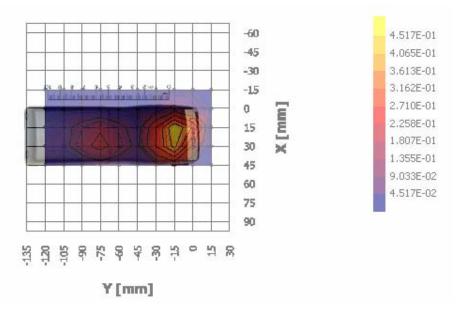
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## 7.2. BODY CONFIGURATION



The left top corner of the DUT in the rear-view was set as origin point (0, 0) for all the measurements in body configuration. The second peak was not evaluated since its reading was found to be at least 3 dB lower than the first peak reading.

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# 7.2.1. Back-side of DUT parallel to the phantom and in contact\*

#	Configuration	Antenna Position	Frequency [MHz]	Channel	SAR <sub>local</sub> Before [W/Kg]	SAR <sub>local</sub> After [W/Kg]	$\begin{array}{c} MAX \\ SAR_{1g} \\ {}_{[W/Kg]} \end{array}$
*	General population/Uncontrolled Exposure Category Limit					1.6	
13	Headset #1		2401.056	Lowest			-
14	Back-side of DUT parallel to the phantom and in	Integrated	2440.800	Middle	0.12	0.10	0.29
15	contact		2482.272	Highest			-
16	Headset #2		2401.056	Lowest			-
17	Back-side of DUT parallel to the phantom and in	Integrated	2440.800	Middle	0.16	0.15	0.31
18	contact		2482.272	Highest			-
19	Headset #3		2401.056	Lowest			-
20	Back-side of DUT parallel to the phantom and in	Integrated	2440.800	Middle	0.15	0.15	0.33
21	contact		2482.272	Highest			-

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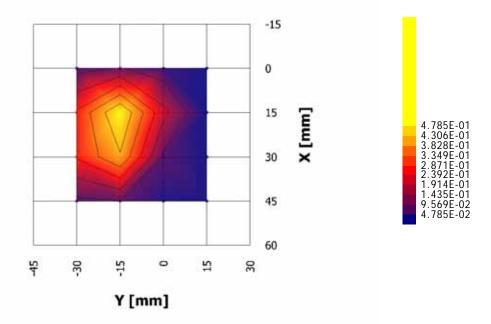
<sup>\*</sup> If the SAR measured at the middle channel for each test configuration is at least 3.0 dB lower than the SAR limit, testing at the high and low channels is optional for such test configuration(s).

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CM-16, 2.4 GHz Cordeless Phone M/N: CM-16

#### 7.2.1.1. D.U.T. perpendicular to the phantom and in contact, Generic Headset 1; #14

Test date [MM/DD/YYYY]	08/25/2004
Test by	JaeWook Choi
Room temperature [°C]	24
Room humidity [%]	40
Simulated tissue temperature [°C]	24
Separation distance, d [mm]	0
Test frequency [MHz]	2440.800
E-field Probe	M/N: ET20, S/N:03JUN-0028, Sensor Offset: 2.0 mm
Sensor Factor $(\eta_{Pd})_{[mV/(mW/cm)]}^2$	10.8
Amplifier Settings (AS <sub>1</sub> , AS <sub>2</sub> , AS <sub>3</sub> )	0.0055211224, 0.0059622436, 0.0056122209
Tissue Type	Muscle
Measured conductivity [S/m]	1.94 (-0.6 %)
Measured dielectric constant	50.2 (-4.8 %)
Conversion Factor (γ)	4.068
Sensitivity ( $\zeta$ ) [W/Kg/mV]	1.665E-01
Measurement Area Specification (X × Y )	$45 _{\text{mm}} \times 45 _{\text{mm}}$ ; Resolution: $15 _{\text{mm}} \times 15 _{\text{mm}}$
Measurement Volume Specification $(X \times Y \times Z)$	$7_{\text{pts}} \times 7_{\text{pts}} \times 7_{\text{pts}}$ , $30_{\text{mm}} \times 30_{\text{mm}} \times 30_{\text{mm}}$ ; Resolution: $5_{\text{mm}} \times 5_{\text{mm}} \times 5_{\text{mm}}$
SAR <sub>1g</sub> [W/Kg]	0.29



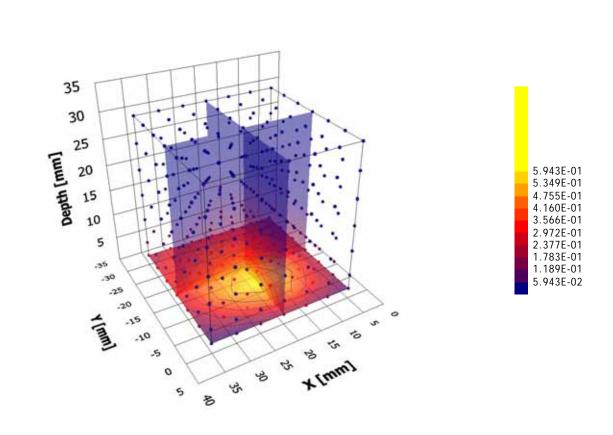
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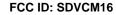
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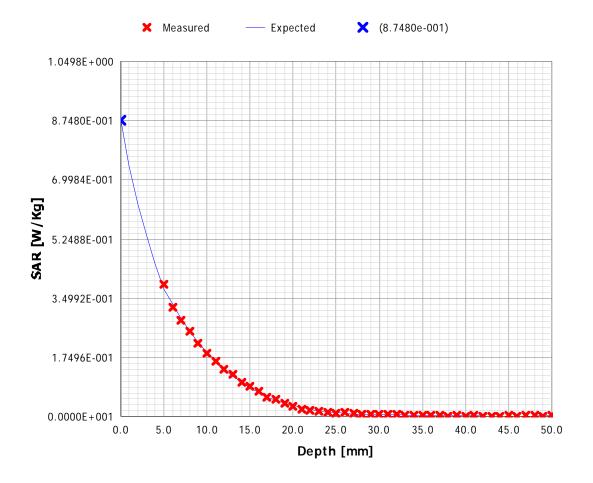
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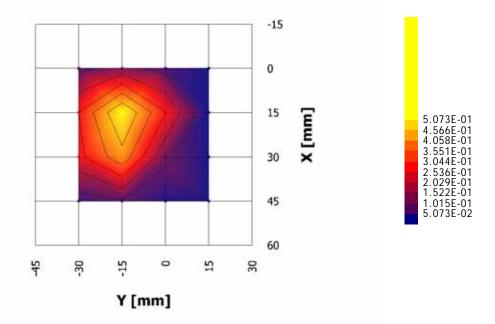
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CM-16, 2.4 GHz Cordeless Phone M/N: CM-16

#### 7.2.1.2. D.U.T. perpendicular to the phantom and in contact, Generic Headset 2; #17

Test date [MM/DD/YYYY]	08/25/2004
Test by	JaeWook Choi
Room temperature [°C]	24
Room humidity [%]	40
Simulated tissue temperature [°C]	24
Separation distance, d [mm]	0
Test frequency [MHz]	2440.800
E-field Probe	M/N: ET20, S/N:03JUN-0028, Sensor Offset: 2.0 mm
Sensor Factor $(\eta_{Pd})_{[mV/(mW/cm)]}^2$	10.8
Amplifier Settings (AS <sub>1</sub> , AS <sub>2</sub> , AS <sub>3</sub> )	0.0055211224, 0.0059622436, 0.0056122209
Tissue Type	Muscle
Measured conductivity [S/m]	1.94 (-0.6 %)
Measured dielectric constant	50.2 (-4.8 %)
Conversion Factor (γ)	4.068
Sensitivity (ζ) [W/Kg/mV]	1.665E-01
Measurement Area Specification (X × Y )	$45 _{\text{mm}} \times 45 _{\text{mm}}$ ; Resolution: $15 _{\text{mm}} \times 15 _{\text{mm}}$
Measurement Volume Specification $(X \times Y \times Z)$	$7_{\text{pts}} \times 7_{\text{pts}} \times 7_{\text{pts}}$ , $30_{\text{mm}} \times 30_{\text{mm}} \times 30_{\text{mm}}$ ; Resolution: $5_{\text{mm}} \times 5_{\text{mm}} \times 5_{\text{mm}}$
SAR <sub>1g</sub> [W/Kg]	0.31



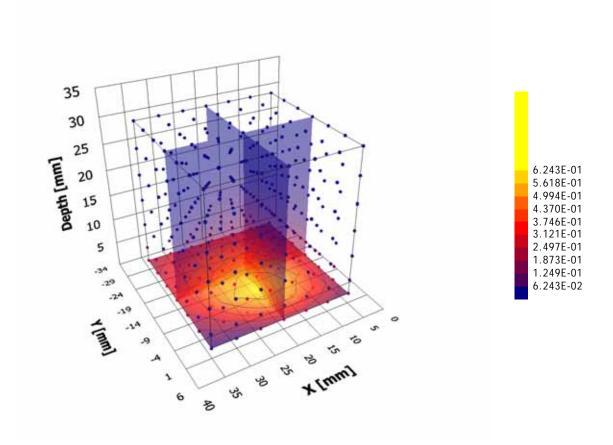
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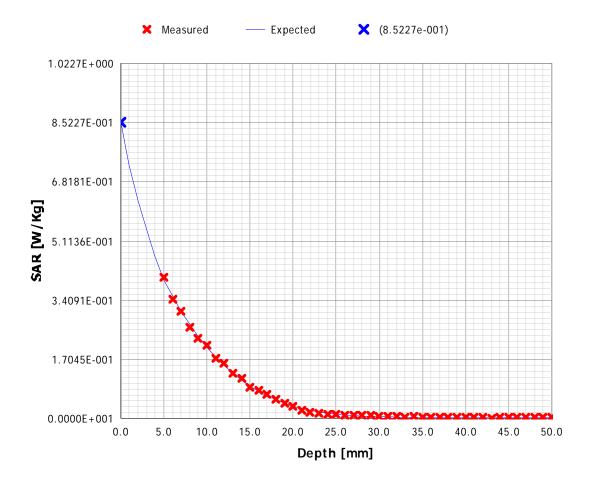
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FCC ID: SDVCM16



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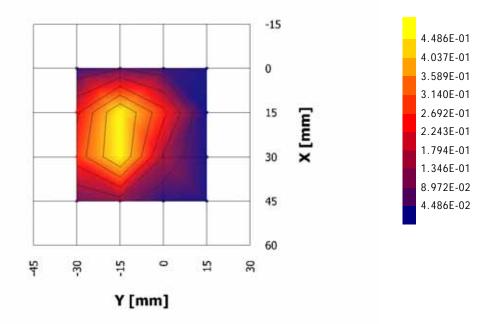
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### 7.2.1.3. D.U.T. perpendicular to the phantom and in contact, Generic Headset 3; #20

Test date [MM/DD/YYYY]	08/25/2004
Test by	JaeWook Choi
Room temperature [°C]	24
Room humidity [%]	40
Simulated tissue temperature [°C]	24
Separation distance, d [mm]	0
Test frequency [MHz]	2440.800
E-field Probe	M/N: ET20, S/N:03JUN-0028, Sensor Offset: 2.0 mm
Sensor Factor $(\eta_{Pd})_{[mV/(mW/cm)]}^2$	10.8
Amplifier Settings (AS <sub>1</sub> , AS <sub>2</sub> , AS <sub>3</sub> )	0.0055211224, 0.0059622436, 0.0056122209
Tissue Type	Muscle
Measured conductivity [S/m]	1.94 (-0.6 %)
Measured dielectric constant	50.2 (-4.8 %)
Conversion Factor (γ)	4.068
Sensitivity (ζ) [W/Kg/mV]	1.665E-01
Measurement Area Specification (X × Y )	$45 _{\text{mm}} \times 45 _{\text{mm}}$ , Resolution: $15 _{\text{mm}} \times 15 _{\text{mm}}$
Measurement Volume Specification $(X \times Y \times Z)$	$7_{\rm pts} \times 7_{\rm pts} \times 7_{\rm pts}$ , $30_{\rm mm} \times 30_{\rm mm} \times 30_{\rm mm}$ ; Resolution: $5_{\rm mm} \times 5_{\rm mm} \times 5_{\rm mm}$
SAR <sub>1g</sub> [W/Kg]	0.33



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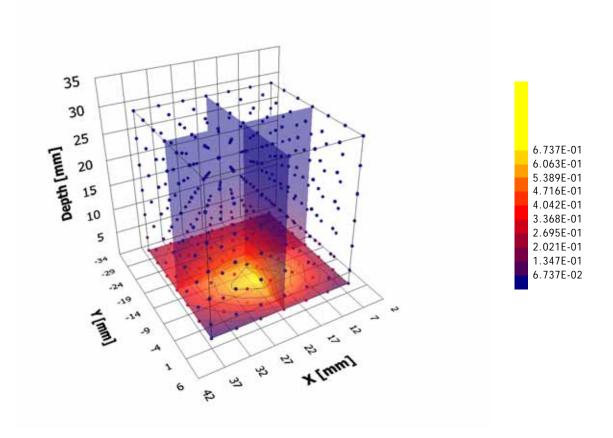
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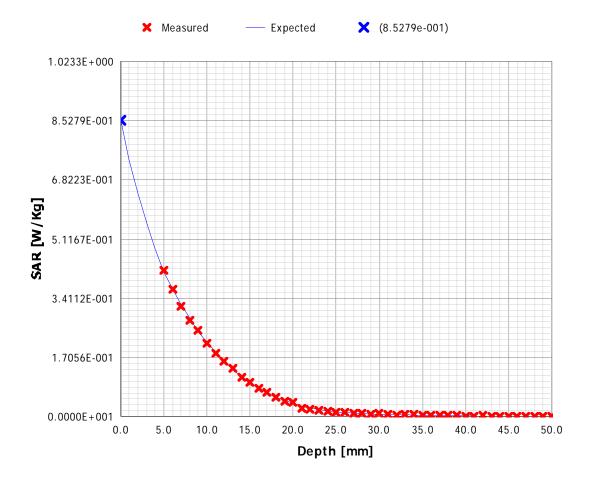
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# **EXHIBIT 8. TISSUE DIELECTRIC PARAMETER CALIBRATION**

# 8.1. SIMULATED TISSUE AT 2450 MHZ

Tissue calibration type	HP Dielectric Strength Probe Syste	m (M/N: 85070C)
Tissue calibration date [MM/DD/YYYY]	08/25/2004	08/25/2004
Tissue calibrated by	JaeWook Choi	JaeWook Choi
Room temperature [°C]	24	24
Room humidity [%]	40	40
Simulated tissue temperature [°C]	24	24
Tissue calibration frequency [MHz]	2450	2450
Tissue Type	Brain	Muscle
Target conductivity [S/m]	1.80	1.95
Target dielectric constant	39.2	52.7
Composition (by weight) [%]	DI Water (59.70 %)	DI Water (76.19 %)
	DGBE (4.76 %)	DGBE (9.52 %)
	Triton X-100 (35.54 %)	Triton X-100 (14.29 %)
Measured conductivity [S/m]	1.78 (-1.0 %)	1.94 (-0.6 %)
Measured dielectric constant	38.2 (-2.6 %)	50.2 (-4.8 %)
Penetration depth (plane wave excitation) [mm]	18.7	19.6

#### **8.1.1. 2450 MHz Brain Tissue**

	Meas. after 5min		DI	DI Water at 20°C			Init. Meas.		
Frequency [GHz]	ε'	ε"	σ [S/m]	ε'	ε"	σ [S/m]	ε'	ε"	σ [S/m]
2.400000000	38.3018	13.0468	1.74	78.6228	10.6185	1.42	38.2627	12.9997	1.74
2.425000000	38.2638	13.0468	1.76	78.5962	10.7410	1.45	38.2242	13.0518	1.76
2.450000000	38.1947	13.0755	1.78	78.5606	10.8424	1.48	38.1710	13.0733	1.78
2.475000000	38.1014	13.1150	1.81	78.5354	10.9339	1.51	38.1013	13.0987	1.80
2.500000000	38.0058	13.1555	1.83	78.4954	11.0600	1.54	37.9732	13.1232	1.83

### 8.1.2. 2450 MHz Muscle Tissue

	Meas. after 5min		DI	DI Water at 20°C			Init. Meas.		
Frequency [GHz]	e'	e"	s [S/m]	e'	e"	s [S/m]	e'	e''	s [S/m]
2.400000000	50.3128	14.0880	1.88	78.7453	10.7677	1.44	50.2424	14.1529	1.89
2.425000000	50.2702	14.1671	1.91	78.7189	10.8718	1.47	50.1996	14.2223	1.92
2.450000000	50.1692	14.2259	1.94	78.6689	10.9878	1.50	50.1170	14.2959	1.95
2.475000000	50.1107	14.2946	1.97	78.6308	11.1000	1.53	50.0374	14.3587	1.98
2.500000000	50.0365	13.3609	1.86	78.5972	11.1927	1.56	49.9656	14.4266	2.01

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# **EXHIBIT 9. SAR SYSTEM CALIBRATION**

# 9.1. GENERAL INFORMATION OF THE PROBE

Probe Type	E-Field Triangle, Isotropic
Model Number	ET20
Serial Number	02FEB-0003
Manufacturer	EMF Safety
Manufactured Date	February 2002
Probe Length [mm]	270
Probe offset [mm]	2.0
Probe Tip diameter [mm]	4.0
Sensor Factor $(\eta_{Pd})_{[mV/(mW/cm])}^2$	10.8
Sensor Factor $(\eta_{E2})_{[mV/(V/m)]}^2$	10.8 / 3770

# 9.2. PROBE LINEARITY AND DYNAMIC RANGE

# 9.2.1. Diode Compression Potential

DCP <sub>1</sub>	66459
DCP <sub>2</sub>	67796
DCP <sub>3</sub>	69561

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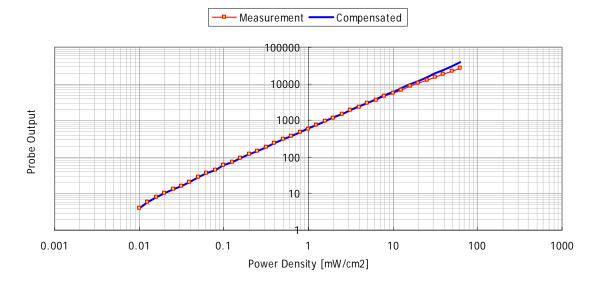
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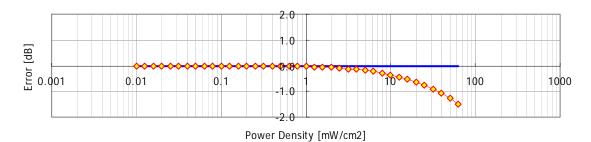
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# **9.2.2.** Channel 1





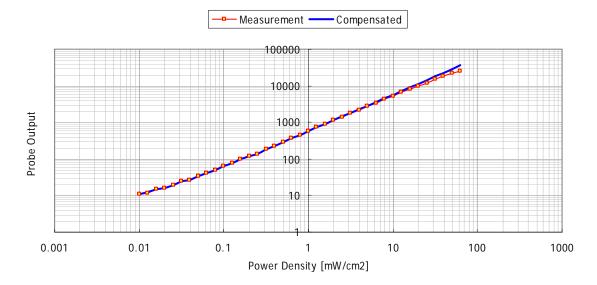
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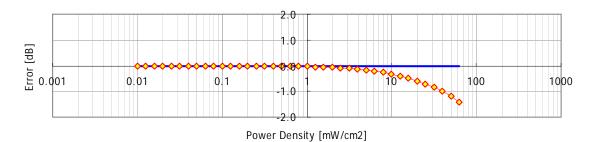
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# 9.2.3. Channel 2





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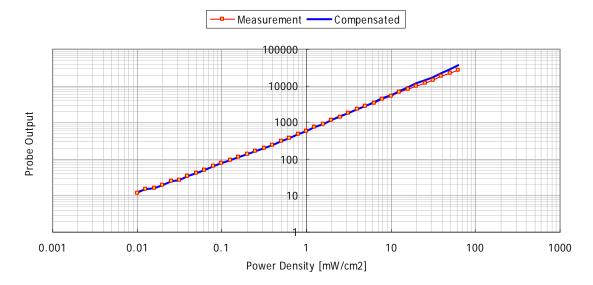
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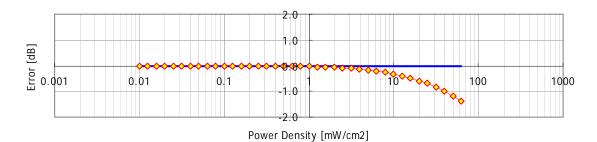
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# **9.2.4.** Channel 3





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# 9.3. PROBE FREE SPACE CALIBRATION

# 9.3.1. Calibration Setup at 2450 MHz

Calibration cell type	Waveguide
Model Number	13492
Serial Number	12463-1, 12463-2
Manufacturer	APOLLO
Input Power / Power Density [mW/(mW/cm )]	24.7

# 9.3.2. Amplifier Settings

### 9.3.2.1. Freespace calibration at 2450 MHz

Calibration Date [MM/DD/YYYY]	08/25/2004
Calibrated by	JaeWook Choi
Calibration Frequency [MHz]	2450
Room Temperature [°C]	24
Room Humidity [%]	40
φ [°]	90
$\Phi_1, \Phi_2, \Phi_3$ [°]	54.7, 54.7, 54.7
Pd <sub>[mW/cm]</sub> <sup>2</sup>	2.0
$SUM(U_{L,1}(0^{\circ}),, U_{L,1}(360^{\circ}))$	33,048
$SUM(U_{L,2}(0^{\circ}),, U_{L,2}(360^{\circ}))$	30,603
$SUM(U_{L,3}(0^{\circ}),, U_{L,3}(360^{\circ}))$	32,512
$R_{\rm ISO,1,1}$	1.0000000000
$R_{ISO,2,1}$	1.0798970070
$R_{ISO,3,1}$	1.0165000009
$AS_1$	0.0055211224
$AS_2$	0.0059622436
$AS_3$	0.0056122209

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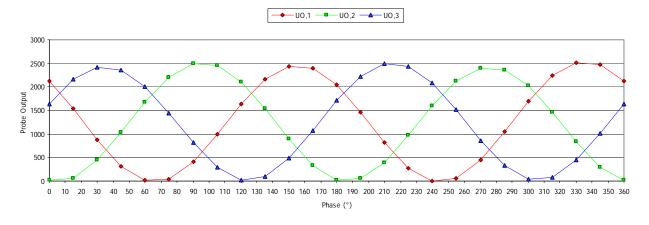
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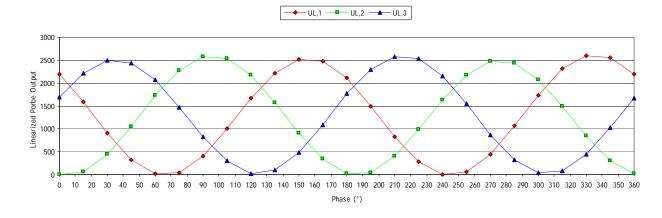
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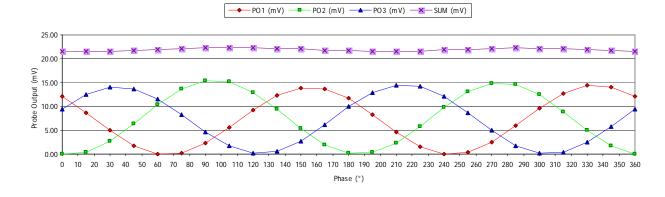
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# 9.3.3. Isotropic response







Isotropy at 2450 MHz: ±0.09 dB

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# 9.4. CALCULABLE WAVEGUIDE CALIBRATION

# 9.4.1. Calibration Setup

# 9.4.1.1. Setup for 2400 MHz

Calibration type	Calculable waveguide calibration
Waveguide Manufacturer	APOLLO
Waveguide dimension (a × b) [mm]	$109.2 \times 54.7$
Dielectric Separator thickness [mm]	12.6
Dielectric Separator permittivity	5.0
Return Loss (at test frequency) [dB]	Better than -20



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# 9.4.2. Simulated Tissue

#### 9.4.2.1. Brain Tissue at 2400 MHz

Tissue calibration type	HP Dielectric Strength Probe System
Tissue calibration date [MM/DD/YYYY]	08/25/2004
Tissue calibrated by	JaeWook Choi
Room temperature [°C]	24
Room humidity [%]	40
Simulated tissue temperature [°C]	24
Tissue calibration frequency [MHz]	2400
Tissue Type	Brain
Target conductivity [S/m]	1.76
Target dielectric constant	39.3
Measured conductivity [S/m]	1.74 (-1.0 %)
Measured dielectric constant	38.2 (-2.6 %)
Penetration depth (plane wave excitation) [mm]	19.1

#### 9.4.2.2. Muscle Tissue at 2400 MHz

Tissue calibration type	HP Dielectric Strength Probe System
Tissue calibration date [MM/DD/YYYY]	08/25/2004
Tissue calibrated by	JaeWook Choi
Room temperature [°C]	24
Room humidity [%]	40
Simulated tissue temperature [°C]	24
Tissue calibration frequency [MHz]	2400
Tissue Type	Muscle
Target conductivity [S/m]	1.90
Target dielectric constant	52.8
Measured conductivity [S/m]	1.88 (-1.0 %)
Measured dielectric constant	50.3 (-4.7 %)
Penetration depth (plane wave excitation) [mm]	20.2

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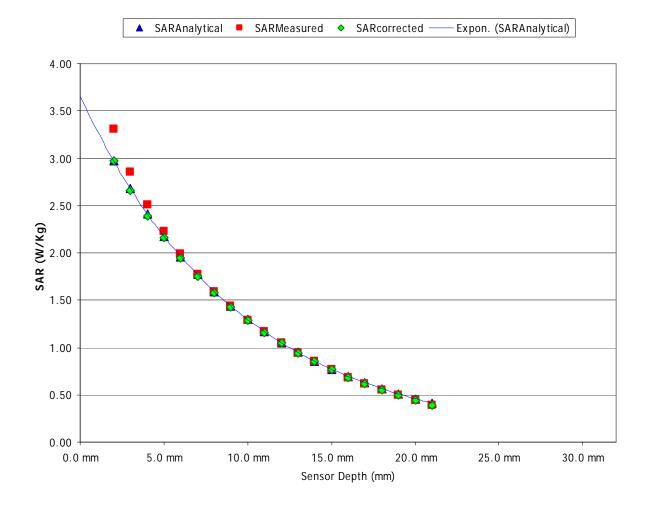
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# 9.4.3. Conversion Factor

#### 9.4.3.1. Calculable waveguide calibration at 2400 MHz for simulated brain tissue

Calibration Date [MM/DD/YYYY]	08/25/2004	
Calibration by	JaeWook Choi	
Calibration Frequency [MHz]	2400	
Room Temperature [°C]	24	
Room Humidity [%]	40	
Simulated Tissue Temperature [°C]	24	
Input Power [mW]	105.02	
Waveguide dimension (a × b) [mm]	$0.1092 \times 0.0547$	
Penetration Depth $\delta_{[m]}$	0.0185	
Conversion Factor (γ)	4.539	



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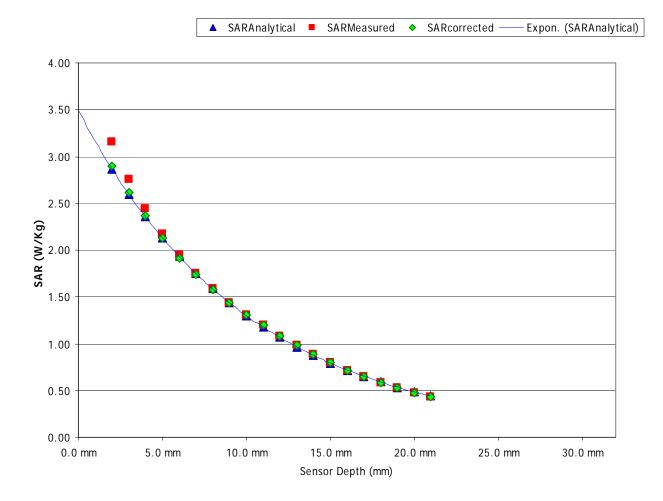
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### 9.4.3.2. Calculable waveguide calibration at 2400 MHz for simulated muscle tissue

Calibration Date [MM/DD/YYYY]	08/25/2004	
Calibration by	JaeWook Choi	
Calibration Frequency [MHz]	2400	
Room Temperature [°C]	24	
Room Humidity [%]	40	
Simulated Tissue Temperature [°C]	24	
Input Power [mW]	105.35	
Waveguide dimension $(a \times b)_{[mm]}$	$0.1092 \times 0.0547$	
Penetration Depth $\delta_{[m]}$	0.0196	
Conversion Factor (γ)	4.068	



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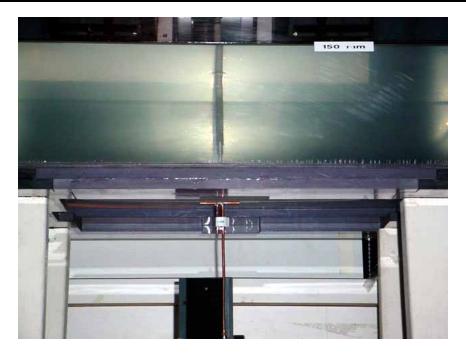
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# **EXHIBIT 10. SAR SYSTEM VERIFICATION**

# 10.1. VERIFICATION SETUP

# 10.1.1. Test setup at 2400 MHz using the dipole reference

Flat phantom dimension (W $\times$ L $\times$ H) $_{[mm]}$	420 × 700 × 200
Flat phantom shell thickness (d <sub>3</sub> ) [mm]	2.0
Flat phantom shell permittivity	2.98
Reference dipole dimension $(L \times h \times d)_{[mm]}$	$51.7 \times 30.4 \times 3.6$
Dipole-to-Phantom (d <sub>2</sub> ) [mm]	8.0
<b>Dipole-to-Liquid</b> $(d_2 + d_3)_{[mm]}$	10.0 (8.0 + 2.0)
Return Loss (at test frequency) [dB]	-24.0



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# 10.2. SIMULATED TISSUE

# 10.2.1. Simulated brain tissue at 2400 MHz

Tissue calibration type	HP Dielectric Strength Probe System
Tissue calibration date [MM/DD/YYYY]	08/25/2004
Tissue calibrated by	JaeWook Choi
Room temperature [°C]	24
Room humidity [%]	40
Simulated tissue temperature [°C]	24
Tissue calibration frequency [MHz]	2400
Tissue Type	Brain
Target conductivity [S/m]	1.76
Target dielectric constant	39.3
Measured conductivity [S/m]	1.74 (-1.0 %)
Measured dielectric constant	38.3 (-2.5 %)
Penetration depth (plane wave excitation) [mm]	19.1

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# 10.3. VERIFICATION RESULT

# 10.3.1. Reference SAR values for simulated brain tissue at 2400 MHz\*

Reference SAR <sub>1g [W/Kg]</sub>	51.1
Reference SAR <sub>s [W/Kg]</sub>	100.9
Measured SAR <sub>1g [W/Kg]</sub>	53.4
Measured SAR <sub>s [W/Kg]</sub>	109.9

#### 10.3.2. Verification result at 2400 MHz

Test date [MM/DD/YYYY]	08/25/2004
Test by	JaeWook Choi
Room temperature [°C]	22
Room humidity [%]	40
Simulated tissue temperature [°C]	22
Test frequency [MHz]	2400
E-field Probe	M/N: ET20, S/N: 02FEB-0003, Sensor Offset: 2.0 <sub>mm</sub>
Sensor Factor $(\eta_{Pd})_{[mV/(mW/cm)]}^2$	10.8
Amplifier Settings (AS <sub>1</sub> , AS <sub>2</sub> , AS <sub>3</sub> )	0.0055211224, 0.0059622436, 0.0056122209
Tissue Type	Brain
Measured conductivity [S/m]	1.74 (-1.0 %)
Measured dielectric constant	38.3 (-2.5 %)
Conversion Factor (γ)	4.539
Sensitivity (ζ) [W/Kg/mV]	1.338E-01
Power [mW]	250
Measurement Volume Specification $(X \times Y \times Z)$	$7_{\rm pts} \times 7_{\rm pts} \times 7_{\rm pts}$ , $30_{\rm mm} \times 30_{\rm mm} \times 30_{\rm mm}$ ; Resolution: $5_{\rm mm} \times 5_{\rm mm} \times 5_{\rm mm}$
$SAR_{1g\ [W/Kg]}$	13.35
SAR <sub>s [W/Kg]</sub>	27.48

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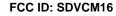
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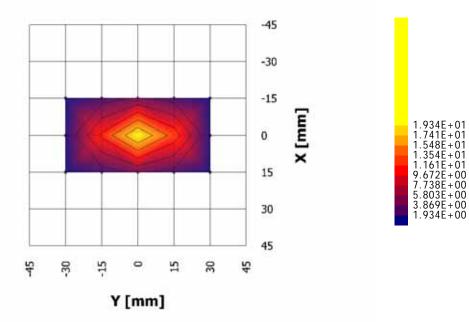
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<sup>\*</sup> All SAR values in 10.3.1 are normalized to a forward power of 1 W.

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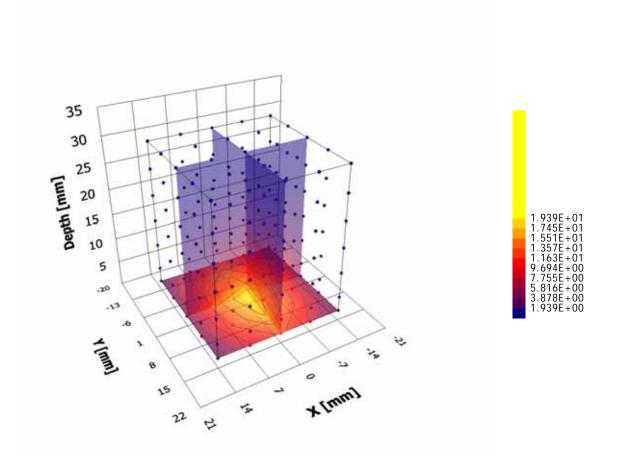
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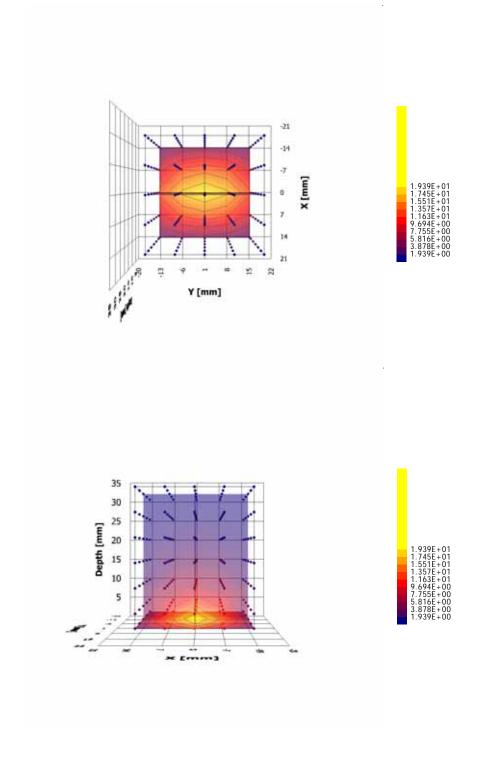


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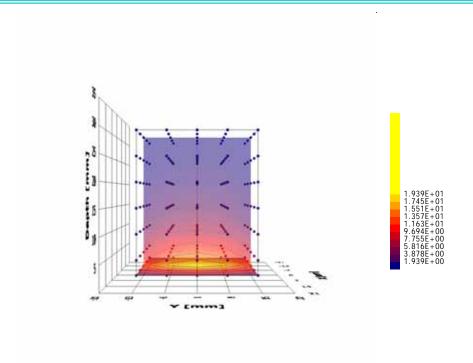
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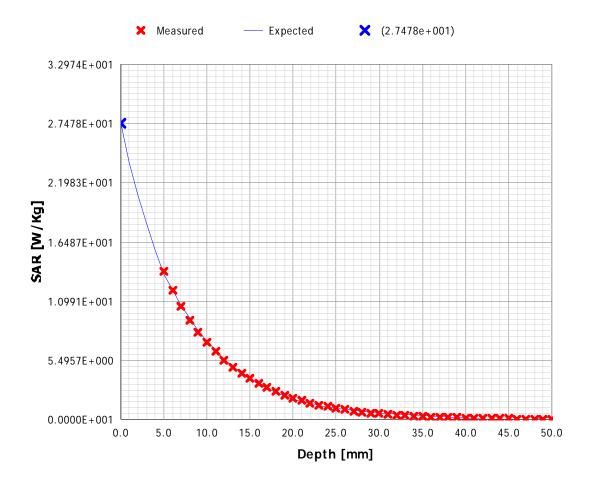
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# **EXHIBIT 11. SAR CALCULATION SUMMARY**

# 11.1. TERMINOLOGY

 $\begin{array}{ll} U_{O,i} & Probe \ raw \ output \ for \ channel \ i \\ U_{L,i} & Linearized \ probe \ output \ for \ channel \ i \end{array}$ 

DCP<sub>i</sub> Diode Compression Potential; Diode specific compression factor for channel i

 $R_{\text{ISO.i.1}}$  Isotropic correction factor normalized to channel 1

AS<sub>i</sub> Amplifier Setting for channel i

 $\eta_{pd}$  Sensor Factor to the uniform power density, an arbitrary value 10.8 [mV/(mW/cm<sup>2</sup>)]

 $\eta_{E2}$  Sensor Factor to the uniform  $|E|^2$ , an arbitrary value 10.8/3,770  $[mV/(V/m)^2]$ 

 $U_{L,1,peak}$  Peak linearized probe output recorded for channel 1 by a rotation about the probe axis with the probe in a test cell

 $\phi$  Smaller angle between the probe axis and the direction of the E-field (i.e.  $90^{\circ}$  when the probe axis is parallel to the

plane of the septum inside TEM cell)

 $\phi_i$  Smaller angle between the probe axis and the dipole sensor axis of the channel i ( $\phi_1 = \phi_2 = 45^\circ$ ,  $\phi_3 = 90^\circ$  for I-beam

probe, and  $\phi_1 = \phi_2 = \phi_3 = 54.7^{\circ}$  for triangular-beam probe)

Pd Well-defined power density [mW/cm<sup>2</sup>] at the calibration point in a test cell

 $\begin{array}{ll} PO_{tot\_solution} & Probe \ output \ [mV] \ in \ the \ simulated \ tissue \\ PO_{tot\_air} & Probe \ output \ [mV] \ in \ the \ air \ (Z_{air} = 377[\Omega]) \\ S.B.T.A.F. & Source-based \ Time-average \ Factor \end{array}$ 

# 11.1.1. Sensor factor( $\eta_{pd}$ and $\eta_{E2}$ ) in the air ( $Z_0 = 377[\Omega]$ )

$$\eta_{Pd} = 10.8[mV/(mW/cm)^2] \equiv \eta_{E2} = \frac{10.8}{3.770}[mV/(V/m)^2]$$

$$Pd[mW/cm^{2}] = \frac{PO_{tot}}{\eta_{Pd}}$$
,  $|E|^{2}[(V/m)^{2}] = \frac{PO_{tot}}{\eta_{E2}}$  and  $SAR[W/Kg] = \frac{\sigma \times \frac{PO_{tot}}{\eta_{E2}}}{\rho}$ 

# 11.1.2. Diode Compensation (Linearity correction)

$$U_{L,i} = U_{O,i} + \frac{{U_{O,i}}^2}{DCP_i}$$

#### 11.1.3. Amplifier settings( $AS_i$ ) and probe output

$$R_{ISO,i,1} = \frac{\sum_{\theta}^{360^{\circ}} U_{L,1}(\theta)}{\sum_{\theta}^{360^{\circ}} U_{L,i}(\theta)}$$

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$$AS_{i} = \eta_{Pd} [mV/(mW/cm^{2})] \times Pd[mW/cm^{2}] \times \cos^{2}(\varphi - \phi_{i}) \times \frac{R_{ISO,i,1}}{U_{L.1,veak}}$$

$$PO_i[mV] = U_{L,i} \times AS_i$$

$$|E_i|^2[(V/m)^2] = \frac{PO_i[mV]}{\eta_{F2}[mV/(V/m)^2]} = \frac{U_{L,i} \times AS_i}{\eta_{F2}[mV/(V/m)^2]}$$

$$|E|^{2}[(V/m)^{2}] = \sum_{i}^{3}|E_{i}|^{2} = \frac{1}{\eta_{E2}} \times \sum_{i}^{3}PO_{i} = \frac{PO_{tot}}{\eta_{E2}} = \frac{1}{\eta_{E2}} \times \sum_{i}^{3}(U_{L,i} \times AS_{i})$$

$$|E|^2[(V/m)^2] = \frac{PO_{tot}[mV]}{\eta_{E2}[mV/(V/m)^2]}$$
, and  $Pd[mW/cm^2] = \frac{PO_{tot}[mV]}{\eta_{Pd}[mV/(mW/m^2)]}$ 

#### 11.1.4. Conversion factor $(\gamma)$ in the simulated tissue

$$\left| E_{solution} \right|^{2} \left[ (V/m)^{2} \right] = \frac{PO_{tot\_solution}[mV]}{\eta_{E2}[mV/(V/m)^{2}]} \times \frac{1}{\gamma}$$

#### 11.1.5. Conversion factor (γ) Calculation #1 – Thermal Trnasfer Calibration (Below 1 GHz)

$$\begin{split} SAR_t &= SAR_{solution} = \frac{\sigma_{@\,cal} \times \left| E_{solution} \right|^2}{\rho_{solution}} \\ \text{,thus } \left| E_{solution} \right|^2 &= \frac{SAR_t \times \rho_{solution}}{\sigma_{@\,cal}} = \frac{PO_{tot\_solution}}{\eta_{E2}} \times \frac{1}{\gamma} \end{split}$$

$$\gamma = \frac{PO_{tot\_solution}}{\eta_{E2}} \times \frac{\sigma_{@cal}}{SAR_t \times \rho_{solution}} = \frac{PO_{tot\_solution}}{\eta_{E2}} \times \frac{\sigma_{@cal}}{\left(c \times \frac{\Delta T}{\Delta t}\right) \times \rho_{solution}}$$

$$= \frac{\sigma_{@ cal}}{\eta_{E2} \times c \times \rho_{solution}} \times \frac{PO_{tot\_solution}}{\left(\frac{\Delta T}{\Delta t}\right)} = \frac{\sigma_{@ cal}}{\eta_{E2} \times c \times \rho_{solution}} \times \frac{\frac{\partial}{\partial P} PO_{tot\_solution}}{\frac{\partial}{\partial P} \left(\frac{\Delta T}{\Delta t}\right)}$$

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CM-16, 2.4 GHz Cordeless Phone M/N: CM-16

# 11.1.6. Conversion factor (γ) Calculation #2 – Calculable Waveguide Calibration (Above 1 GHz)

$$\begin{split} SAR(z) &= SAR_{solution} = \frac{\sigma_{@ cal} \times \left| E_{solution} \right|^{2}}{\rho_{solution}} \\ \text{,thus} \ \left| E_{solution} \right|^{2} &= \frac{SAR(z) \times \rho_{solution}}{\sigma_{@ cal}} = \frac{PO_{tot\_solution}}{\eta_{E2}} \times \frac{1}{\gamma} \\ \gamma &= \frac{PO_{tot\_solution}}{\eta_{E2}} \times \frac{\sigma_{@ cal}}{SAR(z) \times \rho_{solution}} = \frac{PO_{tot\_solution}}{\eta_{E2}} \times \frac{\sigma_{@ cal}}{\frac{4(P_{f} - P_{b})}{\rho_{solution}}} e^{-2z/\delta_{measured}} \times \rho_{solution} \\ &= \frac{PO_{tot\_solution} \times \sigma_{@ cal} \times a \times b \times \delta_{measured}}{\eta_{E2} \times 4(P_{f} - P_{b}) \times e^{-2z/\delta_{measured}}} \end{split}$$

# 11.1.7. Sensitivity ( $\zeta$ ) in the simulated tissue

$$\zeta[W/Kg/mV] = \frac{\sigma_{@meas}[S/m]}{\rho_{head}[Kg/m^{3}]} \times \frac{1}{\eta_{E2}[mV/(V/m)^{2}] \times \gamma}$$

#### 11.1.8. SAR calculation

$$SAR[W / Kg] = \zeta[W / Kg / mV] \times PO_{tot-solution}[mV] \times S.B.T.A.F.$$

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