FCC ID: WLFSTM-7700

Report No.: DRTFCC1101-0030

Total 27pages

# SAR TEST REPORT

	Test item	: Industrial PDA	
	Model No.	: STM-7700	
	Order No.	: 1012-01459	
	Date of receipt	: 2010-12-29	
	Test duration	2010-12-29 ~ 20	11-01-04
	Date of issue	: 2011-01-27	
	Use of report	: FCC Original Gra	ant
Applicant	: Woongjin System 8	& Technology Co., Ltd.	
			Gasan-dong, Geumcheon-gu,
Test laboratory	: Digital EMC Co., Lt	d.	
	683-3, Yubang-Don	ng, Cheoin-Gu, Yongin-	Si, Kyunggi-Do, 449-080, Korea
	Test specification :	§2.1093, FCC/OET Bul	lletin 65 Supplement C[July 2001]
	Test environment :	See appended test rep	ort
	Test result :	⊠ Pass ☐ Fa	il
			hall not be reproduced except in full,
Tested by:	Witness	sed by:	Reviewed by:
Engineer S.K.Ryu	N/A	2	Technical Director Harvey Sung

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# 1. DESCRIPTION OF DEVICE

Environmental evaluation measurements of specific absorption rate (SAR) distributions in emulated human head and body tissues exposed to radio frequency (RF) radiation from wireless portable devices for compliance with the rules and regulations of the U.S. Federal Communications Commission (FCC).

# **General Information**

Equipment type	Industrial PDA
FCC ID:	WLFSTM-7700
Equipment model name	STM-7700
Equipment add model name	N/A
Equipment serial no.	Identical prototype
Mode(s) of Operation	GSM850, PCS1900
TX Frequency Range	824.2 ~ 848.8 MHz(Cellular Band) 1850.2 ~ 1909.8 MHz(PCS Band)
RX Frequency Range	869.2 ~ 893.8 MHz(Cellular Band) 1930.2 ~ 1989.8 MHz(PCS Band)
Max. SAR Measurement	0.976 mW/g GSM850 GPRS Class 10 Body SAR 0.935 mW/g PCS1900 GPRS Class 10 Body SAR
FCC Equipment Class	Licensed Non-Broadcast Station Transmitter(PCB)
Date(s) of Tests	2010-12-29 ~ 2011-01-04
Antenna Type	Internal antenna

#### 2. INTROCUCTION

The FCC has adopted the guidelines for evaluating the environmental effects of radio frequency radiation in ET Docket 93-62 on Aug. 6, 1996 to protect the public and workers from the potential hazards of RF emissions due to FCC-regulated portable devices. The safety limits used for the environmental evaluation measurements are based on the criteria published by the American National Standards Institute (ANSI) for localized specific absorption rate (SAR) in IEEE/ANSI C95.1-2005 Standard for Safety Levels with Respect to Human Exposure to Radio Frequency Electromagnetic Fields, 3 kHz to 300 GHz. 1992 by the Institute of Electrical and Electronics Engineers, Inc., New York, New York 10017. The measurement procedure described in IEEE/ANSI C95.3-1992 Recommended Practice for the Measurement of Potentially Hazardous Electromagnetic Fields - RF and Microwave is used for guidance in measuring SAR due to the RF radiation exposure from the Equipment Under Test (EUT). These criteria for SAR evaluation are similar to those recommended by the National Council on Radiation Protection and Measurements (NCRP) in Biological Effects and Exposure Criteria for Radio frequency Electromagnetic Fields," NCRP Report No. 86 NCRP, 1986, Bethesda, MD 20814. SAR is a measure of the rate of energy absorption due to exposure to an RF transmitting source. SAR values have been related to threshold levels for potential biological hazards.

#### **SAR Definition**

Specific Absorption Rate (SAR) is defined as the time derivative (rate) of the incremental energy (dU)absorbed by (dissipated in) an incremental mass (dm) contained in a volume element (dV) of a given density () It is also defined as the rate of RF energy absorption per unit mass at a point in an absorbing body (see Fig. 2.1)

$$S A R = \frac{d}{d t} \left( \frac{d U}{d m} \right) = \frac{d}{d t} \left( \frac{d U}{\rho d v} \right)$$

Figure 2.1 SAR Mathematical Equation

SAR is expressed in units of Watts per Kilogram (W/kg).

$$SAR = \frac{\sigma \cdot E^2}{\rho}$$

Where:

σ = conductivity of the tissue - simulating material (S/m)

ρ = mass density of the tissue-simulating material (kg/m3)

E = Total RMS electric field strength (V/m)

NOTE: The primary factors that control rate of energy absorption were found to be the wavelength of the incident field in relations to the dimensions and geometry of the irradiated organism, the orientation of the organism in relation to the polarity of field vectors, the presence of reflecting surfaces, and whether conductive contact is made by the organism with a ground plane.

# **Automated TEST SYSTEM SPECIFICATIONS**

#### **Positioner**

Robot: Stäubli Unimation Corp. Robot Model: RX60L

**Repeatability:** 0.02 mm

No. of axis: 6

#### **Data Acquisition Electronic (DAE) System**

Cell Controller

**Processor:** Pentium 4 CPU

Clock Speed: 3 GHz

Operating System: Window 2000

Data Card: DASY4 PC-Board

**Data Converter** 

Features: Signal, multiplexer, A/D converter. & control logic

Software: DASY4

**Connecting Lines:** Optical downlink for data and status info

Optical uplink for commands and clock

**PC Interface Card** 

**Function:** 24 bit (64 MHz) DSP for real time processing

Link to DAE 3

16 bit A/D converter for surface detection system

serial link to robot

direct emergency stop output for robot

E-Field Probes

**Model:** EX3DV4 S/N: 3643

**Construction:** Triangular core fiber optic detection system

Frequency: 10 MHz to 6 GHz

**Linearity:** ±0.2dB (30MHz to 6GHz)

**Phantom** 

**Phantom:** SAM Twin Phantom (V4.0)

**Shell Material:** Composite **Thickness:**  $2.0 \pm 0.2 \text{ mm}$ 



Figure 2.2 DASY4 Test System

#### 3. DESCRIPTION OF TEST EQUIPMENT

#### 3.1 SAR MEASUREMENT SETUP

#### **Robotic System**

These measurements are performed using the DASY4 automated dos imetric assessment system. The DASY4 is made by Schmid & Partner Engineering AG (SPEAG) in Zurich, Switzerland and consists of high precision robotics system (Staubli), robot controller, Pentium III computer, near-field probe, probe alignment sensor, and the generic twin phantom containing the brain equivalent material. The robot is a six-axis industrial robot performing precise movements to position the probe to the location (points) of maximum electromagnetic field (EMF) (see Fig. 3.1).

#### **System Hardware**

A cell controller system contains the power supply, robot controller teach pendant (Joystick), and a remote control used to drive the robot motors. The PC consists of the Micron Pentium IV 500 MHz computer with Windows NT system and SAR Measurement Software DASY4, A/D interface card, monitor, mouse, and keyboard. The Staubli Robot is connected to the cell controller to allow software manipulation of the robot. A data acquisition electronic (DAE) circuit that performs the signal amplification, signal multiplexing, AD-conversion, offset measurements, mechanical surface detection, collision detection, etc. is connected to the Electro-optical coupler (EOC). The EOC performs the conversion from the optical into digital electric signal of the DAE and transfers data to the PC plug-in card.

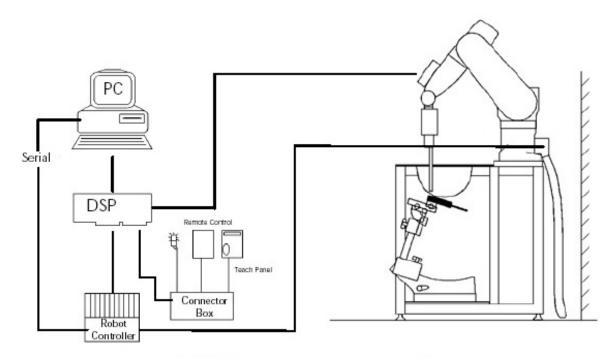


Figure 3.1 SAR Measurement System Setup

#### **System Electronics**

The DAE3 consists of a highly sensitive electrometer-grade preamplifier with auto-zeroing, a channel and gain-switching multiplexer, a fast 16 bit AD-converter and a command decoder and control logic unit. Transmission to the PC-card is accomplished through an optical downlink for data and status information and an optical uplink for commands and clock lines. The mechanical probe mounting device includes two different sensor systems for frontal and sidewise probe contacts. They are also used for mechanical surface detection and probe collision detection. The robot uses its own controller with a built in VME-bus computer. The system is described in detail in.

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#### 3.2 Probe Measurement System

The SAR measurements were conducted with the dosimetric probe EX3DV4, designed in the classical triangular configuration (see Fig. 3.2) and optimized for dosimetric evaluation. The probe is constructed using the thick film technique; with printed resistive lines on ceramic substrates. The probe is equipped with an optical multi fiber line ending at the front of the probe tip. (see Fig. 3.3) It is connected to the EOC box on the robot arm and provides an automatic detection of the phantom surface. Half of the fibers are connected to a pulsed infrared transmitter, the other half to a synchronized receiver. As the probe approaches the surface, the reflection from the surface produces a coupling from the transmitting to the receiving fibers. This reflection increases first during the approach, reaches maximum and then decreases. If the probe is flatly touching the surface, the coupling is zero. The distance of the coupling maximum to the surface is independent of the surface reflectivity and largely independent of the surface to probe angle. The DASY4 software reads the reflection during a software approach and looks for the maximum using a 2nd order fitting. The approach is stopped at reaching the maximum.



**DAE System** 

#### **Probe Specifications**

Calibration: In air from 10 MHz to 6.0 GHz

In brain and muscle simulating tissue at Frequencies of

835 MHz, 1750 MHz, 1900 MHz, 2450 MHz, 2600 MHz, 3500 MHz

Frequency: 10 MHz to 6 GHz

**Linearity:** ±0.2dB (30 MHz to 6 GHz)

**Dynamic:** 10 mW/kg to 100 W/kg

Range: Linearity: ±0.2dB

**Dimensions:** Overall length: 330 mm

**Tip length:** 20 mm

Body diameter: 12 mm

Tip diameter: 2.5 mm

Distance from probe tip to sensor center: 1 mm

Application: SAR Dosimetry Testing

Compliance tests of mobile phones

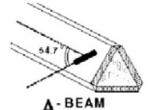


Figure 3.2 Triangular Probe Configurations



Figure 3.3 Probe Thick-Film Technique

#### 3.3 Probe Calibration Process

#### **Dosimetric Assessment Procedure**

Each probe is calibrated according to a dosimetric assessment procedure described in with accuracy better than +/-10%. The spherical isotropy was evaluated with the procedure described in and found to be better than +/-0.25dB. The sensitivity parameters (Norm X, Norm Y, Norm Z), the diode compression parameter (DCP) and the conversion factor (Conv F) of the probe is tested.

#### **Free Space Assessment**

The free space E-field from amplified probe outputs is determined in a test chamber. This is performed in a TEM cell for frequencies below 1 GHz, and in a waveguide above 1GHz for free space. For the free space calibration, the probe is placed in the volumetric center of the cavity at the proper orientation with the field. The probe is then rotated 360 degrees until the three channels show the maximum reading. The power density readings equates to 1 mW/cm<sup>2</sup>.

#### Temperature Assessment \*

E-field temperature correlation calibration is performed in a flat phantom filled with the appropriate simulated brain tissue. The measured free space E-field in the medium, correlates to temperature rise in a dielectric medium. For temperature correlation calibration a RF transparent the rmistor based temperature probe is used in conjunction with the E-field probe

$$SAR = C \frac{\Delta T}{\Delta t}$$

$$SAR = \frac{|E|^2 \cdot \sigma}{\rho}$$

where: where:

 $\Delta t$  = exposure time (30 seconds),  $\sigma$  = simulated tissue conductivity,

C = heat capacity of tissue (brain or muscle),  $\rho$  = Tissue density (1.25 g/cm<sup>3</sup> for brain tissue)

 $\Delta T$  = temperature increase due to RF exposure.

SAR is proportional to  $\Delta T$  /  $\Delta t$  , the initial rate of tissue heating, before thermal diffusion takes place. Now it's possible to quantify the electric field in the simulated tissue by equating the thermally derived SAR to the E- field;

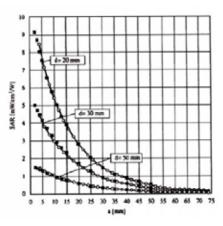


Figure 3.4 E-Field and Temperature Measurements at 900MHz

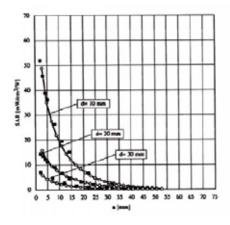


Figure 3.5 E-Field and Temperature Measurements at 1800MHz

#### **Data Extrapolation**

The DASY4 software automatically executes the following procedures to calculate the field units from the microvolt readings at the probe connector. The first step of the evaluation is a linearization of the filtered input signal to account for the compression characteristics of the detector diode. The compensation depends on the input signal, the diode type and the DC-transmission factor from the diode to the evaluation electronics. If the exciting field is pulsed, the crest factor of the signal must be known to correctly compensate for peak power. The formula for each channel can be given like below;

with 
$$V_i$$
 = compensated signal of channel i (i=x,y,z)  
 $U_i$  = input signal of channel i (i=x,y,z)  
 $U_i$  = input signal of channel i (i=x,y,z)  
 $U_i$  = crest factor of exciting field (DASY parameter)  
 $U_i$  = crest factor of exciting field (DASY parameter)

From the compensated input signals the primary field data for each channel can be evaluated:

E-field probes: with 
$$V_i$$
 = compensated signal of channel i (i = x,y,z)  
Norm<sub>i</sub> = sensor sensitivity of channel i (i = x,y,z)  
 $\mu V/(V/m)^2$  for E-field probes  
ConvF = sensitivity of enhancement in solution  
 $E_i$  = electric field strength of channel i in V/m

The RSS value of the field components gives the total field strength (Hermetian magnitude):

$$E_{tot} = \sqrt{E_x^2 + E_y^2 + E_z^2}$$

The primary field data are used to calculate the derived field units.

$$SAR = E_{txt}^2 \cdot \frac{\sigma}{\rho \cdot 1000}$$
 with  $SAR = local specific absorption rate in W/g = total field strength in V/m = conductivity in [mho/m] or [Siemens/m] p = equivalent tissue density in g/cm3$ 

The power flow density is calculated assuming the excitation field to be a free space field.

$$P_{prop} = \frac{E_{tot}^2}{3770}$$
 with  $P_{prop} = \text{equivalent power density of a plane wave in W/cm}^2$  = total electric field strength in V/m

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#### 3.4 SAM PHANTOM

The SAM Twin Phantom V4.0 is constructed of a fiberglass shell integrated in a wooden table. The shape of the shell is based on data from an anatomical study designed to determine the maximum exposure in at least 90% of all users. It enables the dosimetric evaluation of left and right hand phone usage as well as body mounted usage at the flat phantom region. A cover prevents the evaporation of the liquid. Reference markings on the Phantom allow the complete setup of all predefined phantom positions and measurement grids by manually teaching three points in the robot. (see Fig. 3.6)

#### **Phantom Specification**

**Phantom:** SAM Twin Phantom (V4.0)

Shell Material: Vivac Composite Thickness:  $2.0 \pm 0.2 \text{ mm}$ 

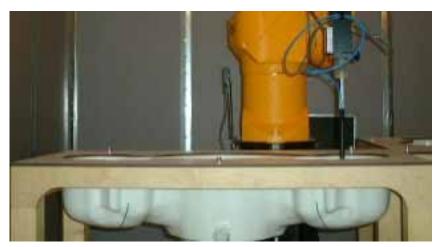


Figure 3.6 SAM Twin Phantom

#### 3.5 Device Holder for Transmitters

In combination with the SAM Twin Phantom V4.0 the Mounting Device (see Fig. 3.7), enables the rotation of the mounted transmitter in spherical coordinates whereby the rotation point is the ear opening. The devices can be easily, accurately, and repeat ably be positioned according to the FCC, CENELEC, IEC and IEEE specifications. The device holder can be locked at different phantom locations (left head, right head, flat phantom).



Figure 3.7 Mounting Device

Note: A simulating human hand is not used due to the complex anatomical and geometrical structure of the hand that may produce infinite number of configurations. To produce the worst-case condition (the hand absorbs antenna output power), the hand is omitted during the tests.

#### 3.6 Brain & Muscle Simulating Mixture Characterization



**Simulated Tissue** 

The brain and muscle mixtures consist of a viscous gel using hydrox-ethyl cellulose (HEC) gelling agent and saline solution (see Table 3.1). Preservation with a bacteriacide is added and visual inspection is made to make sure air bubbles are not trapped during the mixing process. The mixture is calibrated to obtain proper dielectric constant (permittivity) and conductivity of the desired tissue. The mixture characterizations used for the brain and muscle tissue simulating liquids are according to the data by C. Gabriel and G. Harts grove.

**Table 3.1 Composition of the Tissue Equivalent Matter** 

INGREDIENTS	835MHz Brain	835MHz Muscle	1800MHz Brain	1800MHz Muscle	1900MHz Brain	1900MHz Muscle	2450MHz Brain	2450MHz Muscle
WATER	40.19%	50.75%	55.24%	69.04%	55.24%	70.23%	71.88%	73.4%
SUGAR	57.90%	48.21%	-	-	-	-	-	-
SALT	1.48%	0.94%	0.31%	2.72%	0.31%	0.29%	0.16%	0.06%
DGBE	-	-	44.45%	28.24%	44.45%	29.48%	7.99%	26.54%
Triton X-100	-	-	-	-	-	-	19.97%	-
BACTERIACIDE	0.18%	0.10%	-	-	-	-	-	-
HEC	0.25%	-	-	-	-	-	-	-
Dielectric Constant Target	41.5	55.2	40	53.3	40	53.3	39.2	52.7
Conductivity Target (S/m)	0.9	0.97	1.4	1.52	1.4	1.52	1.8	1.95

Salt: 99 % Pure Sodium Chloride Sugar: 98 % Pure Sucrose

Water: De-ionized, 16M resistivity HEC: Hydroxyethyl Cellulose

DGBE: 99 % Di(ethylene glycol) butyl ether,[2-(2-butoxyethoxy) ethanol]

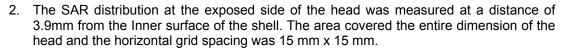
Triton X-100(ultra pure): Polyethylene glycol mono [4-(1,1,3,3-tetramethylbutyl)phenyl]

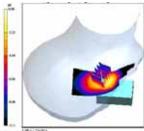
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#### 4. SAR MEASUREMENT PROCEDURE

The evaluation was performed using the following procedure:

1. The SAR measurement was taken at a selected spatial reference point to monitor power variations during testing. This fixed location point was measured and used as a reference value.





Sample SAR Area Scan

- 3. Based on the area scan data, the area of the maximum absorption was determined by sp line interpolation. Around this point, a volume of 32 mm x32 mm x 30 mm (fine resolution volume scan, zoom scan) was assessed by measuring 5 x 5 x 7 points. On this basis of this data set, the spatial peak SAR value was evaluated with the following procedure (see Sample Sar Area Scan):
  - a. The data at the surface was extrapolated, since the center of the dipoles is 2.5 mm away from the tip of the probe and the distance between the surface and the lowest measuring point is 1.2mm. The extrapolation was based on a least square algorithm. A polynomial of the fourth order was calculated through the points in z-axes. This polynomial was then used to evaluate the points between the surface and the probe tip.
  - b. The maximum interpolated value was searched with a straight-forward algorithm. Around this maximum the SAR values averaged over the spatial volumes (1g or 10g) were computed using the 3D-Spline interpolation algorithm. The 3D-spline is composed of three one-dimensional sp lines with the "Not a knot" condition (in x, y, and z directions). The volume was integrated with the trapezoidal algorithm. One thousand points (10 x 10x 10) were interpolated to calculate the average.
  - c. All neighboring volumes were evaluated until no neighboring volume with a higher average value was found.
- 4. The SAR reference value, at the same location as step 1, was re-measured. If the value changed by more than 5%, the evaluation is repeated.

#### Specific Anthropomorphic Mannequin (SAM) Specifications

The phantom for handset SAR assessment testing is a low-loss dielectric shell, with shape and dimensions derived from the anthropometric data of the 90th percentile adult male head dimensions as tabulated by the US Army. The SAM Twin Phantom shell is bisected along the mid-sagittal plane into right and left halves (see Fig. 4.1). The perimeter side walls of each phantom halves are extended to allow filling with liquid to a depth that is sufficient to minimized reflections from the upper surface. The liquid depth is maintained at a minimum depth of 15cm to minimize reflections from the upper surface.



Figure 4.1 Sam Twin Phantom shell

#### 5. DEFINITION OF REFERENCE POINTS

#### **EAR Reference Point**

Figure 5.1 shows the front, back and side views of the SAM Twin Phantom. The point "M" is the reference point for the center of the mouth, "LE" is the left ear reference point (ERP), and "RE" is the right ERP. The ERPs are 15mm posterior to the entrance to the Ear canal (EEC) along the B-M line (Back-Mouth), as shown in Figure 5.5. The plane Passing, through the two ear canals and M is defined as the Reference Plane. The line N-F (Neck- Front) is perpendicular to the reference plane and passing through the RE (or LE) is called the Reference Pivoting Line (see Figure 5.2). Line B-M is perpendicular to the N-F line. Both N-F and B-M lines are marked on the external phantom shell to facilitate handset positioning.



Figure 5.1 Front, back and side view of SAM Twin Phantom

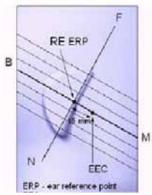


Figure 5.2 Close-up side view of ERPs

#### **Handset Reference Points**

Two imaginary lines on the handset were established: the vertical centerline and the horizontal line. The test device was placed in a normal operating position with the "test device reference point" located along the "vertical centerline" on the front of the device aligned to the "ear reference point" (See Fig. 5.3). The "test device reference point" was than located at the same level as the center of the ear reference point. The test device was positioned so that the "vertical centerline" was bisecting the front surface of the handset at it's top and bottom edges, positioning the "ear reference point" on the outer surface of the both the left and right head phantoms on the ear reference point.

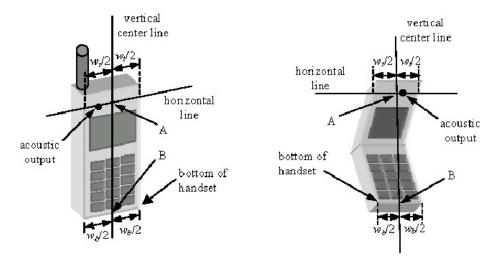


Figure 5.3 Handset Vertical Center & Horizontal Line Reference Points

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#### **5.1 TEST CONFIGURATION POSITIONS**

#### **Positioning for Cheek/Touch**

1. The test device was positioned with the handset close to the surface of the phantom such that point A is on the (virtual) extension of the line passing through points RE and LE on the phantom (see Figure 5.4), such that the plane defined by the vertical center line and the horizontal line of the phone is approximately parallel to the sagittal plane of the phantom.



Figure 5.4 Front, Side and Top View of Cheek/Touch Position

- 2. The handset was translated towards the phantom along the line passing through RE & LE until the handset touches the ear.
- 3. While maintaining the handset in this plane, the handset was rotated around the LE-RE line until the vertical centerline was in the plane normal to MB-NF including the line MB (reference plane).
- 4. The phone was hen rotated around the vertical centerline until the phone (horizontal line) was symmetrical was respect to the line NF.
- 5. While maintaining the vertical centerline in the reference plane, keeping point A on the line passing through RE and LE, and maintaining the phone contact with the ear, the handset was rotated about the line NF until any point on the handset made contact with a phantom point below the ear (cheek). (See Figure 5.5)

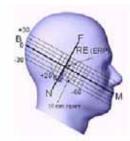


Figure 5.5 Side view w/ relevant markings

#### Positioning for Ear / 15 ° Tilt

With the test device aligned in the "Cheek/Touch Position":

- 1. While maintaining the orientation of the phone, the phone was retracted parallel to the reference plane far enough to enable a rotation of the phone by 15degree.
- 2. The phone was then rotated around the horizontal line by 15 degree.
- 3. While maintaining the orientation of the phone, the phone was moved parallel to the reference plane until any part of the phone touches the head. (In this position, point A was located on the line RE-LE). The tilted position is obtained when the contact is on the pinna. If the contact was at any location other than the pinna, the angle of the phone would then be reduced. The tilted position was obtained when any part of the phone was in contact of the ear as well as a second part of the phone was in contact with the head (see Figure 5.6).



Figure 5.6 Front, Side and Top View of Ear/15° Tilt Position

#### **Body Holster /Belt Clip Configurations**

Body-worn operating configurations are tested with the belt-clips and holsters attacked to 3the device and positioned against a flat phantom in a normal use configuration (see Figure 5.7). A device with a headset output is tested with a headset connected to the device. Body dielectric parameters are used.





Figure 5.7 Body Belt Clip & Holstor Configurations

Accessories for Body-worn operation configurations are divided into two categories: those that do not contain metallic components and those that contain metallic components. When multiple accessories that do not contain metallic components are supplied with the device, the device is tested with only the accessory that dictates the closest spacing to the body. Then multiple accessories that contain metallic components are supplied with the device, the device is tested with each accessory that contains a unique metallic component. If multiple accessories share an identical metallic component (i.e. 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 is tested.

Body-worn accessories may not always be supplied or available as options for some Devices intended to be authorized for body-worn use. In this case, a test configuration where a separation distances between the back of the device and the flat phantom is used. All test position spacing is documented.

Transmitters that are designed to operate in front of a person's face, as in push-to-talk configurations, are tested for SAR compliance with the front of the device positioned to face the flat phantom. For devices that are carried next to the body such as a shoulder, waist or chest-worn transmitters, SAR compliance is tested with the accessories, including headsets and microphones, attached to the device and positioned against a flat phantom in a normal use configuration.

In all cases SAR measurements are performed to investigate the worst-case positioning. Worst-case positioning is then documented and used to perform Body SAR testing.

In order for users to be aware of the body-worn operating requirements for meeting RF exposure compliance, operating instructions and cautions statements are included in the user's manual.

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#### 6. ANSI / IEEE C95.1-2005 RF EXPOSURE LIMITS

#### **Uncontrolled Environment**

UNCONTROLLED ENVIRONMENTS are defined as locations where there is the exposure of individuals who have no knowledge or control of their exposure. The general population/uncontrolled exposure limits are applicable to situations in which the general public may be exposed or in which persons who are exposed as a consequence of their employment may not be made fully aware of the potential for exposure or cannot exercise control over their exposure. Members of the general public would come under this category when exposure is not employment-related; for example, in the case of a wireless transmitter that exposes persons in its vicinity.

#### **Controlled Environment**

CONTROLLED ENVIRONMENTS are defined as locations where there is exposure that may be incurred by persons who are aware of the potential for exposure, (i.e. as a result of employment or occupation). In general, occupational/controlled exposure limits are employment, which have been made fully aware of the potential for exposure and can exercise control over their exposure. This exposure category is also applicable when the exposure is of a transient nature due to incidental passage through a location where the exposure levels may be higher than the general population/uncontrolled limits, but the exposed person is fully aware of the potential for exposure and can exercise control over his or her exposure by leaving the area or by some other appropriate means.

	HUMAN EXPO	HUMAN EXPOSURE LIMITS				
	General Public Exposure (W/kg) or (mW/g)	Occupational Exposure (W/kg) or (mW/g)				
Whole-Body average SAR (W/kg)	0.08	0.40				
Localized SAR (head and trunk) (W/kg)	1.60	8.00				
Localized SAR (limbs) (W/kg)	4.00	20.0				

Table 6.1.SAR Human Exposure Specified in ANSI / IEEE C95.1-2005

#### NOTES:

- \* The Spatial Peak value of the SAR averaged over any 1 g of tissue (defined as a tissue volume in the shape of a cube) and over the appropriate averaging time.
- \*\* The Spatial Average value of the SAR averaged over the whole-body.
- \*\*\* The Spatial Peak value of the SAR averaged over any 10 g of tissue (defined as a tissue volume in the shape of a cube) and over the appropriate averaging time.

Uncontrolled Environments are defined as locations where there is the exposure of individuals who have no knowledge or control of their exposure.

Controlled Environments are defined as locations where there is exposure that may be incurred by persons who are aware of the potential for exposure, (i.e.as a result of employment or occupation).

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# 7. IEEE P1528 -MEASUREMENT UNCERTAINTIES

Eman Description	Uncertaint	Probability	Divisor	(Ci)	Standard	vi 2 or
Error Description	value ±%	Distribution	DIVISOR	1g	(1g)	Veff
Measurement System						
Probe calibration	± 4.8	Normal	1	1	± 4.8 %	∞
Axial isotropy	± 4.7	Rectangular	√3	0.7	± 1.9 %	∞
Hemispherical isotropy	± 9.6	Rectangular	√3	0.7	± 3.9 %	∞
Boundary Effects	± 1.0	Rectangular	√3	1	± 0.6 %	∞
Probe Linearity	± 4.7	Rectangular	√3	1	± 2.7 %	∞
Detection limits	± 1.0	Rectangular	√3	1	± 0.6 %	∞
Readout Electronics	± 1.0	Normal	1	1	± 1.0 %	∞
Response time	± 0.8	Rectangular	√3	1	± 0.5 %	∞
Integration time	± 2.6	Rectangular	√3	1	± 1.5 %	∞
RF Ambient Conditions	± 3.0	Rectangular	√3	1	± 1.7 %	∞
Probe Positioner	± 0.4	Rectangular	√3	1	± 0.2 %	∞
Probe Positioning	± 2.9	Rectangular	√3	1	± 1.7 %	∞
Algorithms for Max. SAR Eval.	± 1.0	Rectangular	√3	1	± 0.6 %	∞
Test Sample Related						
Device Positioning	± 2.9	Normal	1	1	± 2.9 %	145
Device Holder	± 3.6	Normal	1	1	± 3.6 %	5
Power Drift	± 5.0	Rectangular	√3	1	± 2.9 %	8
Physical Parameters						
Phantom Shell	± 4.0	Rectangular	√3	1	± 2.3 %	8
Liquid conductivity (Target)	± 5.0	Rectangular	√3	0.64	± 1.8 %	∞
Liquid conductivity (Meas.)	± 2.5	Normal	1	0.64	± 1.6 %	∞
Liquid permittivity (Target)	± 5.0	Rectangular	√3	0.6	± 1.7 %	∞
Liquid permittivity (Meas.)	± 2.5	Normal	1	0.6	± 1.5 %	8
CombinedStandard Uncertainty					± 10.3 %	330
Expanded Uncertainty (k=2)					± 20.6 %	

The above measurement uncertainties are according to IEEE P1528 (2003)

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# 8. SYSTEM VERIFICATION

# **Tissue Verification**

**Table 8.1 Simulated Tissue Verification** 

MEASURED TISSUE PARAMETERS								
Date(s)	Torget Frequency	Dielectric constant: ε			Conductivity: σ			
	Target Frequency	Target	Measured	Deviation (%)	Target	Measured	Deviation (%)	
December 29, 2010	835 MHz Muscle	55.2	53.3	-3.44	0.970	0.967	-0.31	
January 04, 2011	1900 MHz Muscle	53.3	51.8	-2.81	1.520	1.580	3.95	

# **Test System Validation**

Prior to assessment, the system is verified to the  $\pm 10\%$  of the specifications at 835 MHz and 1900 MHz by using the system validation kit(s). (Graphic Plots Attached)

**Table 8.2 System Validation** 

SYSTEM DIPOLE VALIDATION TARGET & MEASURED (835 MHz / 1900 MHz values are normalized to a forward power of 1/4 W)									
Date(s)	System Validation Kit:	Target Frequency	Targeted SAR <sub>1g</sub> (mW/g)	Measured SAR <sub>1g</sub> (mW/g)	Deviation (%)				
December 29, 2010	D-835V2, S/N: 464	835 MHz Muscle	2.44	2.49	2.05				
January 04, 2011	D-1900V2, S/N: 5d029	1900 MHz Muscle	9.85	10.0	1.52				

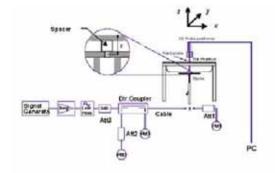




Figure 8.1 Dipole Validation Test Setup

# 9. Multiple TRANSMITTERS SAR CONSIDERATIONS

The following procedures adopted from "FCC SAR Evaluation Considerations for Handsets with Multiple Transmitters" v01r05 #648474 on September 2008 are applicable to handsets with built-in unlicensed transmitters such as 802.11 a/b/g and Bluetooth devices which may simultaneously transmit with the licensed transmitter.

	2.45	5.15-5.35	5.47-5.85	GHz
P Ref	12	6	5	mW

Device output power should be rounded to the nearest mW to compare with values specified in this table

**Table 9.1 Output Power Thresholds for Unlicensed Transmitters** 

	Individual Transmitter	Simultaneous Transmission
Licensed Transmitters	Routine evaluation required	SAR not required: <u>Unlicensed only</u> o when stand-alone 1-q SAR is not
Unlicensed Transmitters	When there is no simultaneous transmission — o output < 60/f: SAR not required o output ≥ 60/f: stand-alone SAR required  When there is simultaneous transmission — Stand-alone SAR not required when  O output ≤ 2.P <sub>Ref</sub> and antenna is > 5.0 cm from other antennas o output ≤ P <sub>Ref</sub> and antenna is > 2.5 cm from other antennas, each either output power output ≤ P <sub>Ref</sub> or 1-g SAR < 1.2 W/Kg  Otherwise stand-alone SAR is required  When stand-alone SAR is required o test SAR on highest output channel for each wireless mode and exposure condition  o if SAR for highest output channel is > 50% of SAR limit, evaluate all channels according to normal procedures	o when stand-alone 1-g SAR is not required and antenna is > 5 cm from other antennas  Licensed & Unlicensed  o when the sum of the 1-g SAR is <1.6 W/kg for all simultaneous transmitting antennas o when SAR to antenna separation ratio of simultaneous transmitting antenna pair is < 0.3  SAR required:  Licensed & Unlicensed  antenna pairs with SAR to antenna separation ratio ≥ 0.3; test is only required for the configuration that results in the highest SAR in standalone configuration for each wireless mode and exposure condition  Note: simultaneous transmission exposure conditions for head and body can be different for different style phones; therefore, different test requirements may apply

Table 9.2 SAR Evaluation Requirements for Cell phones with Multiple Transmitters

FCC ID: WLFSTM-7700

BT Max. RF output power: 2.093 dBm(1.619 mW)
W-LAN Max. RF output power: 16.64 dBm (46.132mW)
Antenna separation distance between GSM and BT: 170 mm
Antenna separation distance between GSM and W-LAN: 200 mm

- Note 1: unlicensed transmitters stand alone SAR is not required when following condition.
  - Output power ≤ P<sub>Ref</sub>, antenna distance from other antennas > 2.5cm each with either output power ≤ P<sub>Ref</sub> or 1-g SAR < 1.2 W/kg</p>
- Note 2 : SAR For Simultaneous transmission
  - ➤ (GSM850<sub>sar</sub> + B/T<sub>sar</sub> + W-LAN<sub>sar</sub>) < 1.6 W/kg, so simultaneous transmission is not performed.
  - ► (PCS1900<sub>sar</sub> + B/T<sub>sar</sub>+ W-LAN<sub>sar</sub>) < 1.6 W/kg, so simultaneous transmission is not performed.

# 10. Configuring 802.11 a/b/g Transmitters for SAR Measurement

#### SAR Testing with IEEE 802.11 a/b/g Transmitters

Normal network operating configurations are not suitable for measuring the SAR of 802.11 a/b/g transmitters. Unpredictable in network traffic and antenna diversity conditions can introduce undesirable variations in SAR results. The SAR for these devices should be measured using chipset based test mode software to ensure the results are consistent and reliable.

#### **General Device Setup**

Chipset based test mode software is hardware dependent and generally varies among manufacturers. The device operating parameters established in test mode for SAR measurements must be identical to those programmed in production units, including output power levels, amplifier gain settings and other RF performance tuning parameters. The test frequencies should correspond to actual channel frequencies defined for domestic use. SAR for devices with switched diversity should be measured with only one antenna transmitting at a time during each SAR measurement, according to a fixed modulation and data rate. The same data pattern should be sued for all measurements.

#### **Frequency Channel Configurations**

802.11 a/b/g and 4.9 GHz operation modes are tested independently according to the service requirements in each frequency band. 802.11 b/g modes are tested on channels 1. 6 and 11. 802.11a is tested for UNII operations on channels 36 and 48 in the 5.15-5.25 GHz Band; channels 52 and 64 in the 5.25-5.35 GHz band; channels 104, 116, 124 and 136 in the 5.470-5.725 GHz BAND; and channel 149 and 161 in the 5.8 GHz band. When 5.8 GHz § 15.247 is also available, channels 149, 157 and 165 should be tested of the UNII channels. 4.9 GHz is tested on channels 1., 10 and 5 or 6, whichever has the higher output power, for 5MHz channels; channels 11, 15 and 19 for 10MHz channels; and channels 21 and 25 for 20MHz channels. These are referred to as the "default test channels". 802.11g mode was evaluated only if the output power was 0.25 dB higher than the 802.11b mode.

		GHz Channel Turbo		"De	fault Test	Channel	s"		
Me	Mode		Channel	Channel	§15.247		ID	UNII	
4400		conserved a	224000000000000000000000000000000000000	Channel	802.11b	802.11g	U	VII.	
		2.412	1"		1	$\nabla$			
802.1	802.11 b/g		6	6	1	$\nabla$		ć.	
	18	2.462	11#		1	$\nabla$			
		5.18	36		22		1		
		5.20	40	42 (5.21 GHz)	ije.				
		5.22	44	42 (J.21 GHZ)			111		
		5.24	48	50 (5.25 GHz)			1		
	73.45	5.26	52	30 (3.23 GHz)			1		
	- 40	5.28	56	58 (5.29 GHz)		-			
	200	5.30	60	36 (3.29 GHz)			1		
		5.32	64			7	1		
		5.500	100						
	UNII	5.520	104				V		
	A	5.540	108						
802.11a	Sec.	5.560	112		and the same of th				
002.11a	-	5.580	116			-	V		
-		5.600	120	Unknown	100				
10		5.620	124		-		1		
M		5.640	128						
100		5.660	132		1				
1		5.680	136		-		<b>\</b>		
The same		5.700	140			65 5	100		
	UNII	5.745	149		1	85 3	4	3	
	or	5.765	153	152 (5.76 GHz)	85 83	*		*	
	§15.247	5.785	157		1	85 3	18	*	
		5.805	161	160 (5.80 GHz)	as 103	*	V	a l	
	§15.247	5.825	165		1				

Table 10.1 802.11 Test channels per FCC Requirements

# 11. SAR TEST DATA SUMMARY AND POWER TABLE

#### 11.1 See Measurement Result Data Pages

#### **Procedures Used To Establish Test Signal**

The EUT was placed into simulated call mode (GSM850, PCS1900) using manufacturers test codes. Such test signals offer a consistent means for testing SAR and are recommended for evaluating SAR. When test modes are not available or inappropriate for testing a EUT, the actual transmission is activated through a base station simulator or similar equipment. See data pages for actual procedure used in measurement.

#### **Device Test Conditions**

The EUT is battery operated. Each SAR measurement was taken with a fully charged battery.

In order to verify that the device was tested at full power, conducted output power measurements were performed before and after each SAR measurement to confirm the output power. If a conducted power deviation of more than 5% occurred, the test was repeated.

#### Max. Power Output Table for STM-7700

		Test Result(dBm)								
Band	Channel	GSM (dBm)	GPRS 1 TX Slot (dBm)	GPRS 2 TX Slot (dBm)	GPRS 3 TX Slot (dBm)	GPRS 4 TX Slot (dBm)	EDGE 1TX Slot (dBm)	EDGE 2TX Slot (dBm)	Slot (dBm)	EDGE 4TX Slot (dBm)
	128	31.7	31.7	31.7	N/A	N/A	N/A	N/A	N/A	N/A
GSM	190	31.8	31.8	31.8	N/A	N/A	N/A	N/A	N/A	N/A
850	251	31.8	31.8	31.8	N/A	N/A	N/A	N/A	N/A	N/A
0014	512	28.9	28.9	28.9	N/A	N/A	N/A	N/A	N/A	N/A
GSM	661	29.1	29.1	29.1	N/A	N/A	N/A	N/A	N/A	N/A
1900	810	28.9	28.9	28.9	N/A	N/A	N/A	N/A	N/A	N/A

#### Max. Power Output Table for STM-7700 (W-LAN)

Mode	Frequency (MHz)	Channel No.	Output Power (dBm) Using the Average Power Meter
	2412	1	16.64
802.11b	2437	6	15.75
	2462	11	15.87
	2412	1	15.31
802.11g	2437	6	14.92
	2462	11	14.74

# Table 11.2 Max. Power Output Table for STM-7700 (Bluetooth)

С	Channel No.	Measured Data (dBm)	(mW)
2402	0	1.863	1.536
2441	39	2.093	1.619
2480	78	2.058	1.606

# 12. SAR TEST DATA SUMMARY

# 12.1 Measurement Results (GSM850 GPRS Body SAR)

FREQU	IENCY	Begin Power	Drift Power	Mode	Device Test	Antenna	SAR
MHz	Ch	(dBm)	(dB)	Wode	Position	Position	(W/kg)
836.6	190	31.8	0.084	GSM850	0 mm [Rear]	Internal	0.535
836.6	190	31.8	0.045	GPRS Class 8	0 mm [Rear]	Internal	0.474
824.2	128	31.7	0.012	GPRS Class 10	0 mm [Rear]	Internal	0.957
836.6	190	31.8	-0.239	GPRS Class 10	0 mm [Rear]	Internal	0.976
848.8	251	31.8	0.189	GPRS Class 10	0 mm [Rear]	Internal	0.904
836.6	190	31.8	-0.070	GPRS Class 10	0 mm [Front]	Internal	0.549
ANSI / IEEE C95.1 2005 – SAFETY LIMIT Spatial Peak Uncontrolled Exposure/ General Population Exposure						Body 1.6 W/kg (mW/g) averaged over 1 gram	

#### NOTE:

- The test data reported are the worst-case SAR value with the antenna-body position set in a typical configuration. Test procedures used are according to FCC/OET Bulletin 65, Supp.C [July 2001].
- 2. All modes of operation were investigated, and worst-case results are reported.
- 3. Prior to testing the conducted output power was measured.
- 4. The EUT is tested 2<sup>nd</sup> hot-spot peak, if it is less than 2dB below the highest peak.
- 5. Battery is fully charged for all readings.
- 6. Test Signal Call Mode 

  Continuous Tx On 

  Manu.Test Codes 

  Base Station Simulator
- 7. Tissue parameters and temperatures are listed on the SAR plots.
- 8. Liquid tissue depth is 15.0cm.±0.1
- 9. Justification for reduced test configurations: per FCC/OET Supplement C (July, 2001), if the SAR measured at the middle channel for each test configuration (Left, right, cheek/touch, tilt/ear, extended and retracted) 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).

#### 12.2 Measurement Results (PCS1900 GPRS Body SAR)

FREQU	ENCY	Begin Power	Drift Power	Mode	Device Test	Antenna	SAR
MHz	Ch	(dBm)	(dB)	Wode	Position	Position	(W/kg)
1880.0	661	29.1	-0.023	PCS1900	0 mm [Rear]	Internal	0.461
1880.0	661	29.1	-0.005	GPRS Class 8	0 mm [Rear]	Internal	0.460
1850.2	512	28.9	0.359	GPRS Class 10	0 mm [Rear]	Internal	0.892
1880.0	661	29.1	-0.371	GPRS Class 10	0 mm [Rear]	Internal	0.935
1909.8	810	28.9	-0.128	GPRS Class 10	0 mm [Rear]	Internal	0.851
1880.0	661	29.1	-0.153	GPRS Class 10	0 mm [Front]	Internal	0.684
ANSI / IEEE C95.1 2005 – SAFETY LIMIT Spatial Peak Uncontrolled Exposure/ General Population Exposure						Body 1.6 W/kg (mW/g) averaged over 1 gram	

#### NOTE:

- 1. The test data reported are the worst-case SAR value with the antenna-body position set in a typical configuration. Test procedures used are according to FCC/OET Bulletin 65, Supp.C [July 2001].
- 2. All modes of operation were investigated, and worst-case results are reported.
- 3. Prior to testing the conducted output power was measured.
- 4. The EUT is tested 2<sup>nd</sup> hot-spot peak, if it is less than 2dB below the highest peak.
- 5. Battery is fully charged for all readings.
- 6.Test Signal Call Mode 

  Continuous Tx On 

  Manu.Test Codes 

  BaseStation Simulator
- 7. Tissue parameters and temperatures are listed on the SAR plots.
- 8. Liquid tissue depth is 15.0cm.±0.1
- 9. Justification for reduced test configurations: per FCC/OET Supplement C (July, 2001), if the SAR measured at the middle channel for each test configuration (Left, right, cheek/touch, tilt/ear, extended and retracted) 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).

# 13. SAR TEST EQUOPMENT

**Table 13.1 Test Equipment Calibration** 

EQ	EQUIPMENT SPECIFICATIONS						
Туре	Calibration Date	Next Calibration Date	Serial Number				
Robot	N/A	N/A	F02/5Q85A1/A/01				
Robot Controller	N/A	N/A	F02/5Q85A1/C/01				
Joystick	N/A	N/A	D221340031				
Hicron Computer 1.1GHz Pentium Celeron ,Window 2000	N/A	N/A	N/A				
Data Acquisition Electronics	November 23, 2010	November 23, 2011	520				
Dosimetric E-Field Probe	January 26, 2010	January 26, 2011	3643				
Dosimetric E-Field Probe	November 22, 2010	November 22, 2011	3123				
Dummy Probe	N/A	N/A	N/A				
Sam Phantom	N/A	N/A	N/A				
Probe Alignment Unit LB	N/A	N/A	321				
SPEAG Validation Dipole D835 MHz	March 22, 2010	March 22, 2012	464				
SPEAG Validation Dipole D1900 MHz	March 23, 2010	March 23, 2012	5d029				
Head/Body Equivalent Matter(835MHz)	January 2010	January 2011	N/A				
Head/Body Equivalent Matter(1900MHz)	January 2010	January 2011	N/A				
HP EPM-442A Power Meter	March 12, 2010	March 12, 2011	GB37170267				
HP ESG-3000A Signal Generator	July 01, 2010	July 01, 2011	US37230529				
Attenuator (10dB)	January 11, 2011	January 11, 2012	BP4387				
Attenuator (3dB)	July 01, 2010	July 01,2011	MY39260700				
Low pass filter (1.5GHz)	January 11, 2011	January 11, 2012	N/A				
Low pass filter (3.0GHz)	October 04, 2010	October 04, 2011	N/A				
Dual Directional Coupler	January 11, 2011	January 11, 2012	50228				
Amplifier	October 04, 2010	October 04, 2011	1020 D/C 0221				
Network Analyzer	March 12, 2010	March 12, 2011	3410J01204				
HP85070D Dielectric Probe Kit	N/A	N/A	LISO1440118				
SEMITEC Engineering	N/A	N/A	Shield Room				
8960Series 10 Wireless Comms Test Set	July 02, 2010	July 02, 2011	GB43461134				

#### NOTE:

The E-field probe was calibrated by SPEAG, by temperature measurement procedure. Dipole Validation measurement is performed by Digital EMC. before each test. The brain simulating material is calibrated by Digital EMC using the dielectric probe system and network analyzer to determine the conductivity and permittivity (dielectric constant) of the brain-equivalent material.

# 14. CONCLUSION

#### **Measurement Conclusion**

The SAR measurement indicates that the EUT complies with the RF radiation exposure limits of the FCC. These measurements are taken to simulate the RF effects exposure under the worst-case conditions. Precise laboratory measures were taken to assure repeatability of the tests. The tested device complies with the requirements in respect to all parameters subject to the test. The test results and statements relate only to the item(s) tested.

Please note that the absorption and distribution of electromagnetic energy in the body are very complex phenomena that depend on the mass, shape, and size of the body, the orientation of the body with respect to the field vectors, and the electrical properties of both the body and the environment. Other variables that may play a substantial role impossible biological effects are those that characterize the environment (e.g. ambient temperature, air velocity, relative humidity, and body insulation) and those that characterize the individual (e.g. age, gender, activity level, debilitation, or disease).

Because innumerable factors may interact to determine the specific biological outcome of an exposure to electromagnetic fields, any protection guide shall consider maximal amplification of biological effects as a result of field-body interactions, environmental conditions, and physiological variables.

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