FCC ID: ZK9STM-8800

Report No.: DRTFCC1105-0203

Total 26pages

SAR TEST REPORT

	Test item	: Industi	rial PDA	
	Model No.	: STM-8	800	
	Order No.	: 1103-0	0274	
	Date of receipt	: 2011-0	3-03	
	Test duration	: 2011-0	5-09 ~ 2011-05-11	
	Date of issue	: 2011-0	5-17	
	Use of report	: FCC C	riginal Grant	
Applicant	: Woongjin Holdings	Co., Ltd.		
	3F. Kukdong Bldg.	Chungmuro	3-ga, Jung-gu, Seoul, Korea	
Test laboratory			u, Yongin-Si, Kyunggi-Do, 449)-080, Korea
	Test specification :	§2.1093, FC	C/OET Bulletin 65 Supplemen	t C[July 2001]
	Test environment :	See appende	ed test report	
	Test result :	□ Pass	☐ Fail	
	report is inhibited other than it without the written	s purpose. This	ly to the sample supplied by applicant test report shall not be reproduced e TAL EMC CO., LTD.	
Tested by:	Witnes	ssed by:	Reviewed by:	

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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:	ZK9STM-8800
Equipment model name	STM-8800
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	0.859 mW/g GSM850 Body SAR 0.798 mW/g PCS1900 Body SAR
FCC Equipment Class	PCS Licensed Transmitter
Date(s) of Tests	2011-05-09 ~ 2011-05-11
Antenna Type	Internal antenna

Note. This report is re-issued according to request of applicant.

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 (ρ) 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.

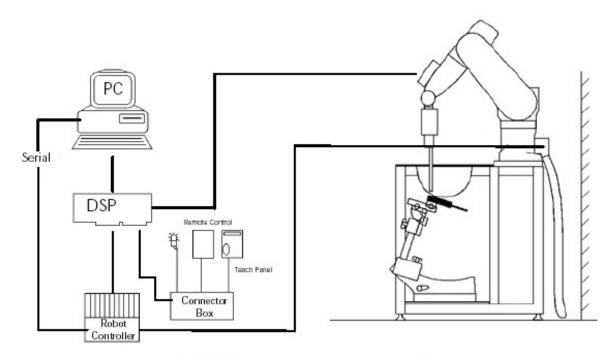


Figure 3.1 SAR Measurement System Setup

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 450 MHz, 835 MHz, 1750 MHz, 1900 MHz

2450 MHz, 2600 MHz, 3500 MHz, 5200 MHz, 5300 MHz, 5600 MHz, 5800 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

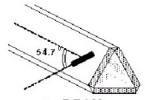
Body diameter: 12 mm

Tip diameter: 2.5 mm

Distance from probe tip to sensor center: 1 mm

Application: SAR Dosimetry Testing

Compliance tests of mobile phones



∆-BEAM

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/cm².

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:

 Δt = exposure time (30 seconds), σ = simulated tissue conductivity,

C = heat capacity of tissue (brain or muscle), ρ = Tissue density (1.25 g/cm³ for brain tissue)

 ΔT = temperature increase due to RF exposure.

SAR is proportional to ΔT / Δt , 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;

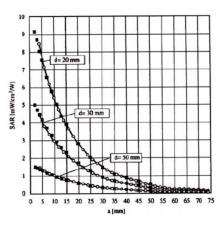


Figure 3.4 E-Field and Temperature Measurements at 900MHz

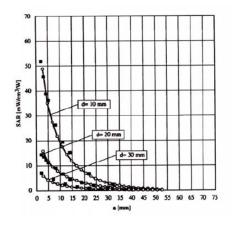


Figure 3.5 E-Field and Temperature Measurements at 1800MHz

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 = diode compression point (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_i = 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.

$$P_{pwe} = \frac{E_{tot}^2}{3770}$$
 with $P_{pwe} = \text{equivalent power density of a plane wave in W/cm}^2$ = 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 ± 0.2 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]

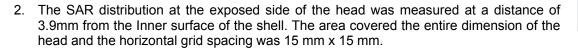
Triton X-100(ultra pure): Polyethylene glycol mono [4-(1,1,3,3-tetramethylbutyl)phenyl]

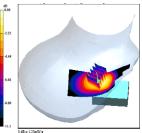
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4. SAR MEASUREMENT PROCEDURE

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

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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.

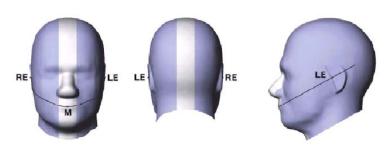


Figure 5.1 Front, back and side view of SAM Twin Phantom

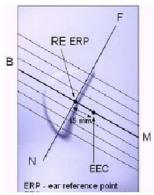


Figure 5.2 Close-up side view of ERPs

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.

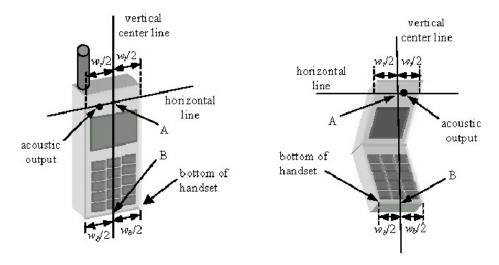


Figure 5.3 Handset Vertical Center & Horizontal Line Reference Points

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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)

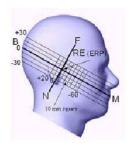


Figure 5.5 Side view w/ relevant markings

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 & Holster 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 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).

7. IEEE P1528 - MEASUREMENT UNCERTAINTIES

Face Description	Uncertaint	Probability	Distant	(Ci)	Standard	vi 2 or
Error Description	value ±%	Distribution	Divisor	1g	(1g)	Veff
Measurement System						
Probe calibration	± 4.8	Normal	1	1	± 4.8 %	8
Axial isotropy	± 4.7	Rectangular	√3	0.7	± 1.9 %	8
Hemispherical isotropy	± 9.6	Rectangular	√3	0.7	± 3.9 %	8
Boundary Effects	± 1.0	Rectangular	√3	1	± 0.6 %	∞
Probe Linearity	± 4.7	Rectangular	√3	1	± 2.7 %	8
Detection limits	± 1.0	Rectangular	√3	1	± 0.6 %	8
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 %	8
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 %	8
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 %	8
Liquid conductivity (Meas.)	± 2.5	Normal	1	0.64	± 1.6 %	8
Liquid permittivity (Target)	± 5.0	Rectangular	√3	0.6	± 1.7 %	8
Liquid permittivity (Meas.)	± 2.5	Normal	1	0.6	± 1.5 %	∞
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 (%)	
May 11, 2011	835 MHz Muscle	55.2	55.3	0.18	0.970	0.947	-2.37	
May 09, 2011	1900 MHz Muscle	53.3	53.1	-0.38	1.520	1.550	1.97	

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 _{1g} (mW/g) Measured SAR _{1g} (mW/g) Deviat							
May 11, 2011	D-835V2, S/N: 464	835 MHz Muscle	2.44	2.48	1.64		
May 09, 2011	D-1900V2, S/N: 5d029	1900 MHz Muscle	9.85	10.0	1.52		

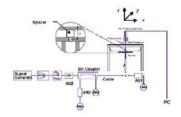




Figure 8.1 Dipole Validation Test Setup

About KDB Publication 450824-2

The following are the recommended FCC procedures for SAR dipole calibration.

- 1) The phantom configuration, tissue dielectric parameters, dipole positioning requirements, dielectric spacer and other electrical and mechanical details should be clearly specified in the dipole calibration report. Dipoles must be recalibrated at least once every three years; however, immediate re-calibration is required for the following conditions. The test laboratory must ensure that the required supporting information and documentation have been included in the SAR report to qualify for the extended 3-year calibration interval; otherwise, the IEEE Standard 1528-2003 recommended annual calibration is expected.
- When the most recent return-loss, measured at least annually, deviates by more than 20% from the previous measurement (i.e. 0.2 of the dB value) or not meeting the required -20 dB return-loss specification

Antenna Parameters

Date(s)	Frequency	Return loss (Head)	Return loss (Body)
March 21.2011	835MHz	-30.5 dB	-25.1 dB
March 21.2011	1900MHz	-36.8 dB	-25.8 dB

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-g 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 _{Ref} and antenna is > 5.0 cm from other antennas o output ≤ P _{Ref} and antenna is > 2.5 cm from other antennas, each either output power output ≤ P _{Ref} 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	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: ZK9STM-8800

BT Max. RF output power: 2.12 dBm(1.629 mW)
W-LAN Max. RF output power: 8.44 dBm(6.982 mW)
Antenna separation distance between GSM and BT: 97 mm
Antenna separation distance between GSM and W-LAN: 109 mm

- Note 1: unlicensed transmitters stand alone SAR is not required when following condition.
 - > Output power ≤ P_{Ref} , antenna distance from other antennas > 2.5cm each with either output power ≤ P_{Ref} or 1-g SAR < 1.2 W/kg

Therefore Bluetooth and W-LAN stand alone SAR is not required.

- Note 2 : SAR For Simultaneous transmission
 - ➤ (GSM850_{sar} + B/T_{sar} + W-LAN_{sar}) < 1.6 W/kg, so simultaneous transmission is not performed.
 - → (PCS1900_{sar} + B/T_{sar}+ W-LAN_{sar}) < 1.6 W/kg, so simultaneous transmission is not performed.
 </p>

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.

			- V	т.	"De	fault Test	Channel	s"
Mo	Mode		Channel	Turbo Channel	§15	.247	ID	пт
TO COMPANY OF THE PARTY OF THE		e d	Channel		802.11b	802.11g	UNII	
		2.412	1"		V	∇		
802.1	1 b/g	2.437	6	6	V	∇		
	8	2.462	11#		1	∇	120	
		5.18	36		28	6	1	
		5.20	40	42 (5.21 GHz)	4	20		
		5.22	44	42 (J.21 GHZ)	i.	20		
		5.24	48	50 (5.25 GHz)	<i>a</i> :	20	1	
	73.6	5.26	52	30 (3.23 0112)		V	1	
		5.28	56	58 (5.29 GHz)		and the same of		*
	200	5.30	60	30 (3.23 GHZ)			1 10	*
		5.32	64				1	
	1	5.500	100		(a)			
	UNII	5.520	104				1	
		5.540	108					
802.11a	The same	5.560	112		and the same of th			
	-	5.580	116			-	1	
-		5.600	120	Unknown	45			
1		5.620	124		-		1	
100		5.640	128			1		*
		5.660	132			_		
		5.680	136		_		1	
1		5.700	140			a - 6		*
	UNII	5.745	149	100 10 00 000	1	8 8	√	2
	or	5.765	153	152 (5.76 GHz)		*		*
	§15.247	5.785	157	1.00.45.00.077.3	1	8 8	-	*
		5.805	161	160 (5.80 GHz)		*	√	a e
	§15.247	5.825	165		1	K s		

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-8800

		Test Result(dBm)								
Band	Channel	GSM	GPRS 1 TX Slot	GPRS 2 TX Slot	GPRS 3 TX Slot	GPRS 4 TX Slot	EDGE 1TX Slot	EDGE 2TX Slot	EDGE 3TX Slot	EDGE 4TX Slot
0014	128	32.5	32.3	32.3	N/A	N/A	N/A	N/A	N/A	N/A
GSM 850	190	32.4	32.2	32.2	N/A	N/A	N/A	N/A	N/A	N/A
650	251	32.2	32.0	32.0	N/A	N/A	N/A	N/A	N/A	N/A
0014	512	29.2	29.2	29.1	N/A	N/A	N/A	N/A	N/A	N/A
GSM 1900	661	29.2	29.2	29.1	N/A	N/A	N/A	N/A	N/A	N/A
	810	29.3	29.1	29.2	N/A	N/A	N/A	N/A	N/A	N/A

Max. Power Output Table for STM-8800 (W-LAN)

Mode	Frequency (MHz)	Channel No.	Output Power (dBm) Using the Average Power Meter
	2412	1	7.54
802.11b	2437	6	8.44
	2462	11	7.44
	2412	1	7.34
802.11g	2437	6	8.24
	2462	11	7.34

Table 11.2 Max. Power Output Table for STM-8800 (Bluetooth)

Frequency (MHz)	Channel No.	Measured Data (dBm)	(mW)
2402	0	1.73	1.489
2441	39	2.12	1.629
2480	78	1.97	1.574

12. SAR TEST DATA SUMMARY

12.1 Measurement Results (GSM850 Body SAR)

FREQUENCY		Begin Power	Drift Power	Mode	Device Test	Antenna	SAR	
MHz	Ch	(dBm)	(dB)	Wode	Position	Position	(W/kg)	
836.6	190	32.2	-0.058	GPRS Class 10	0 mm [Rear]	Internal	0.221	
824.2	128	32.3	0.013	GPRS Class 10	0 mm [Front]	Internal	0.831	
836.6	190	32.2	0.159	GPRS Class 10	0 mm [Front]	Internal	0.847	
848.8	251	32.0	0.050	GPRS Class 10	0 mm [Front]	Internal	0.859	
836.6	190	32.2	-0.077	GPRS Class 8	0 mm [Front]	Internal	0.415	
836.6	190	32.4	0.020	GSM850	0 mm [Front]	Internal	0.447	
ANSI / IEEE C95.1 2005 – SAFETY LIMIT Spatial Peak Uncontrolled Exposure/ General Population Exposure								

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 2nd 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).
- 10. In case of the front body SAR test, It was performed test because user can be a belt clip or handset case in front and rear both sides.

12.2 Measurement Results (PCS1900 Body SAR)

FREQUENCY		Begin Power	Drift Power	Mode	Device Test	Antenna	SAR	
MHz	Ch	(dBm)	(dB)	Wode	Position	Position	(W/kg)	
1880.0	661	29.1	0.198	GPRS Class 10	0 mm [Rear]	Internal	0.315	
1850.2	512	29.1	0.038	GPRS Class 10	0 mm [Front]	Internal	0.798	
1880.0	661	29.1	0.356	GPRS Class 10	0 mm [Front]	Internal	0.734	
1909.8	810	29.2	0.071	GPRS Class 10	0 mm [Front]	Internal	0.657	
1880.0	661	29.2	0.241	GPRS Class 8	0 mm [Front]	Internal	0.359	
1880.0	661	29.2	0.158	PCS1900	0 mm [Front]	Internal	0.389	
ANSI / IEEE C95.1 2005 – SAFETY LIMIT Spatial Peak Uncontrolled Exposure/ General Population Exposure								

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 2nd 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).
- 10. In case of the front body SAR test, It was performed test because user can be a belt clip or handset case in front and rear both sides.

13. SAR TEST EQUOPMENT

Table 13.1 Test Equipment Calibration

	Туре	Manufacturer	Model	Cal.Date	Next.Cal.Date	S/N
	<u> </u>			(dd/mm/yy)	(dd/mm/yy)	
	SEMITEC Engineering	SEMITEC	N/A	N/A	N/A	Shield Room
	Robot Controller	SCHMID	RX90BL CS7MB	N/A	N/A	F02/5Q85A1/A/01
	Robot Controller	SCHMID SCHMID	N/A	N/A N/A	N/A N/A	F02/5Q85A1/C/01
	Joystick Hicron Computer Pentium Celeron					D221340031
	1.1GHz Window2000	N/A	N/A	N/A	N/A	N/A
	Probe Alignment Unit LB	N/A	N/A	N/A	N/A	321
\boxtimes	Mounting Device	SCHMID	Holder	N/A	N/A	N/A
\boxtimes	Sam Phantom	SCHMID	TP1223	N/A	N/A	N/A
\boxtimes	Sam Phantom	SCHMID	TP1224	N/A	N/A	N/A
	Head/Body Equivalent Matter(450MHz)	N/A	N/A	01/01/11	01/01/12	N/A
\boxtimes	Head/Body Equivalent Matter(835MHz)	N/A	N/A	01/01/11	01/01/12	N/A
	Head/Body Equivalent Matter(1800MHz)	N/A	N/A	01/01/11	01/01/12	N/A
	Head/Body Equivalent Matter(1900MHz)	N/A	N/A	01/01/11	01/01/12	N/A
	Head/Body Equivalent Matter(2450MHz)	N/A	N/A	01/01/11	01/01/12	N/A
\boxtimes	Data Acquisition Electronics	SCHMID	DAE3V1	28/01/11	28/01/12	519
	Data Acquisition Electronics	SCHMID	DAE3V1	23/11/10	23/11/11	520
\boxtimes	Dosimetric E-Field Probe	SCHMID	EX3DV4	24/01/11	24/01/12	3643
	Dosimetric E-Field Probe	SCHMID	ET3DV6	06/07/10	06/07/11	1703
	Dummy Probe	N/A	N/A	N/A	N/A	N/A
	450MHz System Validation Dipole	SCHMID	D450V2	24/01/11	24/01/13	1011
\boxtimes	835MHz System Validation Dipole	SCHMID	D835V2	22/03/10	22/03/12	464
	1800MHz System Validation Dipole	SCHMID	D1800V2	16/07/10	16/07/12	2d047
	1900MHz System Validation Dipole	SCHMID	D1900V2	23/03/10	23/03/12	5d029
	2450MHz System Validation Dipole	SCHMID	D2450V2	18/03/10	18/03/12	726
	2600MHz System Validation Dipole	SCHMID	D2600V2	27/05/10	27/05/12	1016
	3500MHz System Validation Dipole	SCHMID	D3500V2	27/05/10	27/05/12	1018
	Network Analyzer	HP	8753D	08/03/11	08/03/12	3410J01204
	Signal Generator	HP	ESG-3000A	01/07/10	01/07/11	US37230529
	Amplifier	EMPOWER	BBS3Q7ELU	04/10/10	04/10/11	1020
	Power Meter	HP	EPM-442A	07/03/11	07/03/12	GB37170267
	Power Sensor	HP	8481A	07/03/11	07/03/12	3318A96566
	Power Sensor	HP	8481A	07/03/11	07/03/12	3318A90918
	Dual Directional Coupler	Agilent	778D-012	11/01/11	11/01/12	50228
	Directional Coupler	HP	773D	01/07/10	01/07/11	2389A00640
\boxtimes	Low Pass Filter 1.5GHz	Micro LAB	LA-15N	11/01/11	11/01/12	N/A
	Low Pass Filter 3.0GHz	Micro LAB	LA-30N	04/10/10	04/10/11	N/A
	Attenuators(3dB)	Agilent	8491B	01/07/10	01/07/11	MY39260700
\boxtimes	Attenuators(10dB)	WEINSCHEL	23-10-34	11/01/11	11/01/12	BP4387
	Step Attenuator	HP	8494A	04/10/10	04/10/11	3308A33341
\boxtimes	Dielectric Probe kit	Agilent	85070D	N/A	N/A	US01440118
\boxtimes	8960 Series 10 Wireless Comms. Test Set	Agilent	E5515C	07/03/11	07/03/12	GB43461134
	Bluetooth Tester F: The E-field probe was calib	TESCOM	TC-3000B	01/07/10	01/07/11	3000B640046

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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