

683-3, Yubang-Dong, Yongin-Si, Kyunggi-Do, Korea. 449-080 Tel: +82-31-321-2664 Fax: +82-31-321-1664

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SAR TEST REPORT

Dates of Tests: April 27~28, 2009 Test Report S/N:DR50110905D Test Site: DIGITAL EMC CO., LTD.

FCC ID

R2NSEC-8089

APPLICANT

Epivalley Co., Ltd.

FCC Classification: Licensed Portable Transmitter (PCB)

EUT Type: EVDO 1X USB Modem

Model Name SEC-8089

Test Device Serial No.: Identical prototype

TX Frequency Range: 824.70 ~848.31 MHz (CDMA_CELL)

1851.25~1908.75MHz(CDMA_PCS)

869.70 ~893.31 MHz (CDMA_CELL) RX Frequency Range:

1931.25~1988.75MHz(CDMA_PCS)

Max. SAR Measurement: 1.120 mW/g CDMA _CELL Body SAR

0.261 mW/g CDMA _PCS Body SAR

Max. RF Output Power 0.222W ERP CDMA Cellular Band (23.46 dBm)

0.040W EIRP CDMA PCS Band (15.98 dBm)

Application Type: Certification

Rule Part(s): §2.1093; FCC/OET Bulletin Supplement C[July 2001]

Data of issue: May 07, 2009

This wireless portable device has been shown to be capable of compliance for localized specific absorption rate (SAR) for uncontrolled environment/general population exposure limits specified in ANSI/IEEE Std. C95.1-1992 and had been tested in accordance with the measurement procedures specified in FCC/OET Bulletin 65 Supplement C (2001) and IEEE Std. 1528-2003.

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

EpiValley 6

Tested by:

Reviewed by :

Sun-Kyu Ryu (Engineer) Harvey Sung (Chief Engineering Director)

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1. INTROCUCTION/SAR DEFINITION

In 1974, the International Radiation Protection Association (IRPA) formed a working group on non-ionizing radiation (NIR), which examined the problems arising in the field of Protection against the various types of NIR. At the IRPA Congress in Paris in 1977, this working group because the International Non-Ionizing Radiation Committee (INIRC).

In cooperation with the Environmental Health Division of the World Health Organization (WHO), the IRPA/INIRC developed a number of health criteria documents on NIR as part of WHO'S Environmental Health Criteria Programme, sponsored by the United Nations Environment Programme (UNEP). Each document includes an overview of the physical characteristics, measurement and instrumentation, sources, and applications of NIR, a thorough review of the literature on biological effects, and an evaluation of the health risks of exposure to NIR. These health criteria have provided the scientific database for the subsequent development of exposure limits and codes of practice relating to NIR.

At the Eighth International Congress of the IRPA (Montreal, 18-22 May 1992), a new, independent scientific organization-the International Commission on Non-Ionizing Radiation Protection (ICNIRP)-was established as a successor to the IRPA/INIRC. The functions of the Commission are to investigate the hazards that may be association with the different forms of NIR, develop international guidelines on NIR exposure to static and extremely-low-frequency (ELF) electric and magnetic field have been reviewed by UNEP/WHO/IRPA (1984, 1987). Those publications and a number of others, including UNEP/WHO/IRPA (1993) and Allen et al. (1991), provided the scientific rationale for these guidelines.

A glossary of terms appears in the Appendix.

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. 1.1)

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

Figure 1.1 SAR Mathematical Equation

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

SAR =
$$E^2$$
/ ρ

Where:

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

 ρ = mass density of the tissue-simulant 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.[6]

2. SAR MEASUREMENT SETUP

Robotic System

Measurements are performed using the DASY4 automated dosimetric 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. 2.1).

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.

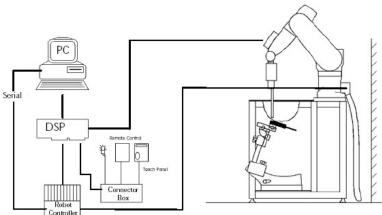


Figure 2.1 SAR Measurement System Setup

System Electronics

The DAE3 consists of a highly sensitive electrometer-grade preamplifier with autozeroing, 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 [7].

3. SAR MEASUREMENT SETUP

Probe Measurement System



The SAR measurements were conducted with the dosimetric probe ET3DV6,designed in the classical triangular configuration [7] (see Fig. 3.2) and 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 multifiber 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 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 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 (see Fig.3.1). The approach is stopped at reaching the maximum.

Figure 3.1 DAE System

Probe Specifications

Calibration: In air from 10 MHz to 6.0 GHz

In brain and muscle simulating tissue at

Frequencies of 900 MHz, 1810 MHz, 1950 MHz, 2450 MHz, 2600 MHz

Frequency: 10 MHz to 6 GHz

Linearity : $$\pm 0.2 dB$$ (30 MHz to 6 GHz)

Dynamic: 10 mW/kg to 100 W/kg

Range : Linearity: $\pm 0.2 \text{ dB}$

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 3.1 Triangular Probe Configuration



Figure 3.2 Probe Thick-Film Technique

4. Probe Calibration Process

Dosimetric Assessment Procedure

Each probe is calibrated according to a dosimetric assessment procedure described in [8] with accuracy better than +/- 10%. The spherical isotropy was evaluated with the procedure described in [9] and found to be better than +/-0.25dB. The sensitivity parameters (NormX, NormY, NormZ), the diode compression parameter (DCP) and the conversion factor (ConvF) 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 (see Fig. 4.1), 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.

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 thermistor-based temperature probe is used in conjunction with the E-field probe (see Fig. 4.2).

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

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

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;

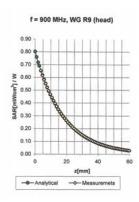


Figure 4.1 E-Field and Temperature Measurements at 900MHz[7]

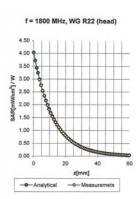


Figure 4.2 E-Field and Temperature Measurements at 1900MHz[7]

5. PHANTOM & EQUIVALENT TISSUES

SAM Phantom



Figure 5.1 SAM Twin 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 [11][12]. 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. 5.1)

Brain & Muscle Simulating Mixture Characterization



The brain and muscle mixtures consist of a viscous gel using hydroxethyl cellullose (HEC) gelling agent and saline solution (see Table 6.1). 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 head tissue dielectric parameters recommended by the IEEE SCC-34/SC-2 have been incorporated in the following table. Other head and body tissue parameters that have not bee specified in P1528 are derived from the issue dielectric parameters computed from he 4-Cole-Cole equations The mixture characterizations used for the brain and muscle tissue simulating liquids are according to the data by C. Gabriel and G. Hartsgrove [13].(see Fig. 5.2)

Figure 5.2 Simulated Tissue

Table 5.1 Composition of the Muscle Tissue Equivalent Matter

INGREDIENTS		SIMULATING TISSUE					
INGREDIENTS		835MHz Brain	835MHz Muscle	1900MHz Brain	1900MHz Muscle		
Mixture Percentage							
WATER		41.45	52.50	54.90	40.40		
DGBE	BE		0.000	44.92	58.00		
SUGAR	GAR		45.00	0.000	0.000		
SALT		1.450	1.400	0.180	0.500		
BACTERICIDE		0.100	0.100	0.000	0.100		
HEC		1.000	1.000	0.000	1.000		
Dielectric Constant	Target	41.50	55.20	40.00	53.30		
Conductivity (S/m) Target		0.900	0.970	1.400	1.520		

Device Holder for Transmitters



Figure 5.2 Mounting Device

In combination with the SAM Twin Phantom V4.0, the Mounting Device (see Fig. 5.2) enables the rotation of the mounted transmitter in spherical coordinates where by the rotation point is the ear opening. The devices can be easily, accurately, and repeatably be positioned according to the FCC specifications. The device holder can be locked at different phantom locations (left head, right head, flat phantom).

 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 [12]. To produce the worst-case condition (the hand absorbs antenna output power), the hand is omitted during the tests.

6. TEST SYSTEM SPECIFICATIONS

Automated Test System Specifications

<u>Positioner</u>

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



Figure 6.1 DASY4 Test System

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: $\pm 0.2 dB(30 MHz to 6 GHz)$

Phantom

Phantom: SAM Twin Phantom (V4.0)

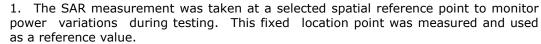
Shell Material: Vivac Composite

Thickness: $2.0 \pm 0.2 \text{ mm}$

7. DOSIMETRIC ASSESSMENT & PHANTOM SPECS

Measurement Procedure

The evaluation was performed using the following procedure:



- 2. The SAR distribution at the exposed side of the head was measured at a distance of 3.9mm from the Inner surface of the shell. The area covered the entire dimension of the head and the horizontal grid spacing was 15mm x 15mm.
- 3. Based on the area scan data, the area of the maximum absorption was determined by spline interpolation. Around this point, a volume of $32\text{mm} \times 32\text{mm} \times 30\text{mm}$ (fine resolution volume scan, zoom scan) was assessed by measuring $5 \times 5 \times 7$ points. On this basis of this data set, the spatial peak SAR value was evaluated with the following procedure (see Fig. 7.1):
- a. The data at the surface was extrapolated, since the center of the dipoles is 2.7 mm away from the tip of the probe and the distance between the surface and the lowest measuring point is 1.2mm. The extrapolation was based on a least square algorithm [15]. 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-for war dalgorithm. 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 splines with the "Not a knot" condition (in x, y, and z directions) [15][16]. The volume was integrated with the trapezoidal algorithm. One thousand points (10 x 10 x 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 procedure #1, was remeasured. If the value changed by more than 5%, the evaluation is repeated.

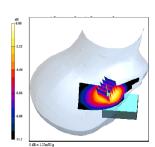


Figure 7.1 Sample Sar Area Scan

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. 7.2). The perimeter sidewalls 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 7.2 SAM Twin Phantom shell

8. DEFINITION OF REFERENCE POINTS

EAR Reference Point

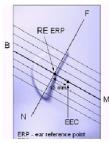


Figure 8.2 Close-up side view of ERPs

Figure 8.1 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 9.2. 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 8.2). 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 [5]



Figure 8.1 Front, back and side view of SAM Twin Phantom

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. 8.3). 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.

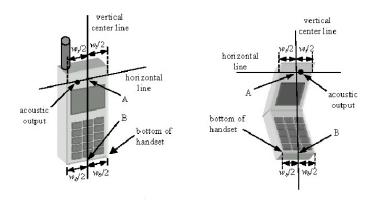


Figure 8.3 Handset Vertical Center & Horizontal Line Reference Points

9. TEST CONFIGURATION POSITIONS (Continued)

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 9.1), 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 9.1 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 9.2)

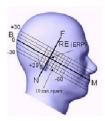


Figure 9.2 Side view w/ relevant markings

9. TEST CONFIGURATION POSITIONS (Continued)

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 9.3).



Figure 9.3 Front, Side and Top View of Ear/15° Tilt Position

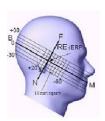


Figure 9.4 Side view w/ relevant markings

9. TEST CONFIGURATION POSITIONS

Body Holster / Belt Clip Configurations

Body-worn operating configurations are tested with the belt-clips and holsters attached to

the device and positioned against a flat phantom in a normal use configuration (see Figure 9.5). A device with a headset output is tested with a headset connected to the device. Body dielectric parameters are used.

Accessories for Body- worn operation configurations are divided into two categories: those that do not contain and those that do contain metallic components 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.





Figure 9.5 Body Belt Clip & Holster Configurations

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 distance between the back of the device and the flat phantom is used.All test position spacings are 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 accessory(ies), 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.

10. ANSI / IEEE C95.1-1992 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, who 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.

Table 10.1.

SAR Human Exposure Specified in ANSI/IEEE C95.1-1992

	HUMAN EXPOSURE LIMITS					
	General Public Exposure Occupational Exposur (W/kg) or (mW/g) (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				

11.IEEE P1528 – MEASUREMENT UNCERTAINTIES

Error Description	Uncertaint	Probability	Divisor	(Ci)	Standard	vi 2 or
Life Description	value ±%	Distribution	DIVISOI	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 %	∞
Physical Parameters						
Phantom Shell	± 4.0	Rectangular	√3	1	± 2.3 %	∞
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 %	∞
Combined Standard Uncertainty					± 10.3 %	330
Expanded Uncertainty (k=2)					± 20.6 %	

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

12. SYSTEM VERIFICATION

Tissue Verification

Table 12.1 Simulated Tissue Verification [5]

MEASURED TISSUE PARAMETERS								
Data(s)	Target Frequency	Dielectric o	constant: ε	Conduct	tivity: σ			
Date(s)	Target Frequency	Target	Measured	Target	Measured			
April.27, 2009	835 MHz Brain	41.5	41.3	0.90	0.921			
April.27, 2009	835 MHz Muscle	55.2	54.0	0.97	0.993			
April.28, 2009	1900 MHz Brain	40.0	39.4	1.40	1.460			
April.28, 2009	1900 MHz Muscle	53.3	52.8	1.52	1.582			

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 12.2 System Validation [5]

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}	Deviation (%)				
April.27, 2009	D-835V2, S/N: 464	835 MHz Brain	2.375	2.44	2.74				
April.28, 2009	D-1900V2, S/N: 5d029	1900V2, S/N: 5d029							

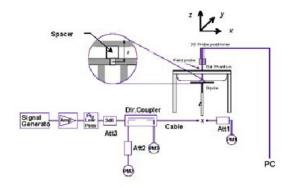




Figure 12.1 Dipole Validation Test Setup

13. FCC 3G SAR MEASUREMENT PROCEDURES - OCT. 2007

FCC 3G MEASUREMENT PROCEDURES

Power measurements were performed using a base station simulator under average power.

13.1 SAR MEASURMENT CONDITIONS FOR CDMA2000

The following procedures were followed according to FCC"SAR Measurements Procedures for 3G Devices" v02, October 2007.

Output Power Verification

See 3GPP2 C.S0011/TIA-98-E as recommended by "SAR Measurement Procedures for 3G Devices", June 2006. Maximum output power is verified on the High, Middle and Low channels according to procedures in section 4.4.5.2 of 3GPP2 C.S0011/TIA-98-E. SO55 tests were measured with power control bits in "All Up" condition.

- 1. If the mobile station (MS) supports Reverse TCH RC 1 and Forward TCH RC 1, set up a call using Fundamental Channel Test Mode 1 (RC=1/1) with 9600 bps data rate only.
- 2. Under RC1, C.S0011 Table 4.4.5.2-1, Table 13-1 parameters were applied.
- 3. If the MS supports the RC 3 Reverse FCH, RC3 Reverse SCH0 and demodulation of RC 3,4, or 5, set up a call using Supplemental Channel Test Mode 3 (RC 3/3) with 9600 bps Fundamental Channel and 9600 bps SCH0 data rate.
- 4. Under RC3, C.S0011 Table 4.4.5.2-2, Table 13-2 was applied.
- 5. FCHs were configured at full rate for maximum SAR with "All Up" power control bits.

Table 13-1
Parameters for Max. Power for RC1

Parameter	Units	Value
Îor	dBm/1.23 MHz	-104
$\frac{\text{Pilot E}_{c}}{\text{I}_{or}}$	dB	-7
Traffic E _c	dB	-7.4

Table 13-2
Parameters for Max. Power for RC3

Parameter	Units	Value		
Îor	dBm/1.23 MHz	-86		
$\frac{\text{Pilot E}_{\text{c}}}{\text{I}_{\text{or}}}$	dB	-7		
$\frac{\text{Traffic E}_{c}}{\text{I}_{or}}$	dB	-7.4		

13.2 Body SAR Measurements

SAR is measured using FTAP/RTAP and FETAP/RETAP respectively for Rev. 0 and Rev. A devices. The AT is tested with a Reverse Data Channel rate of 153.6 kbps in Subtype 0/1 Physical Layer Configurations; and a Reverse Data Channel payload size of 4096 bits and Termination Target of 16 slots in Subtype 2 Physical Layer Configurations. Both FTAP and FETAP are configured with a Forward Traffic Channel data rate corresponding to the 2-slot version of 307.2 kbps with the ACK Channel transmitting in all slots. AT power control should be in All Bits Up conditions for TAP/ETAP.

Body SAR is measured using Subtype 0/1 Physical Layer configurations for Rev. 0. SAR for Subtype 2 Physical Layer configurations is not required for Rev. A when the maximum average output of each RF channels is less than that measured in Subtype 0/1 Physical layer configurations. Otherwise, SAR is measured on the maximum output channel for Rev. A using the exposure configuration that results in the highest SAR for that RF channels in Rev 0. Head SAR is required for EV-DO devices that support operations next to the ear; for example, with VOIP, using Subtype 2 Physical Layer configurations according to the required handset test configurations.

13.3 1x RTT Support

For EV-DO devices that also support 1xRTT voice and/or data operations, SAR is not required for 1xRTT when the maximum average output of each channel is less than 1/4 dB higher than that measured in Subtype 0/1 Physical Layer configurations for Rev. 0. Otherwise, the 'Body SAR Measurements' procedures in the 'CDMA-2000 1x Handsets' section should be applied

13.4 SAR Measurement Procedures for USB Dongle Transmitters

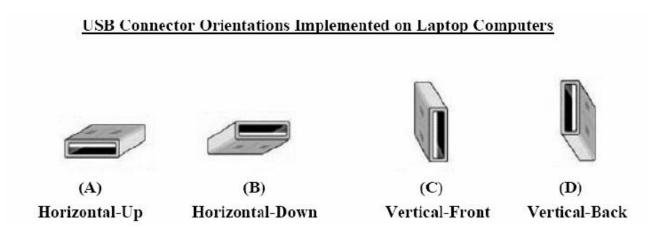
Certain clarifications to the SAR measurement procedures for USB dongle transmitters given in KDB447498 were provided during the TCB workshop on October 7, 2008. These procedures are intended for USB dongle transmitters with internal antennas. If the dongle has a built-in external antenna or one that can swivel, there could be more than four orientations that will require testing. If the USB connector on the dongle can swivel, there are also other conditions that need consideration.

13.5 SAR compliance test considerations

Test all USB orientations (see chart below – A: Horizontal-Up, B: Horizontal-Down, C: Vertical-Front, and D: Vertical-Back) with a device to phantom separation distance of 5mm or less, according to KDB 447498 requirements. Current generation laptop computers should be used to ensure proper measurement distances. The same test separation distance should be used for all frequency bands and modes in each USB orientation. The typical Horizontal-Up USB connection (A), found in the majority of laptop computers, must be tested using an appropriate laptop computer. A laptop with either Vertical-Front (C) or Vertical-Back (D) USB connection should be used to test one of the vertical USB orientations. If laptop computers are not available for testing the Horizontal-Down (B) or the remaining Vertical USB orientations, a short and high quality USB cable (12 inches or less) may be used for testing these other orientations. It should be ensured that the USB cable does not affect device radiating characteristics and output power of the dongle.

These test orientations are intended to cover the exposure conditions found on typical laptop computers with either horizontal or vertical USB connector configurations at various locations in the keyboard section of laptop computers. Depending on the design of an individual dongle, if the antenna is not located at the very end of the dongle or there are no other swiveling mechanisms that could increase the exposure potential, the tip of the dongle typically would not require SAR testing. Dongle tip testing is generally determined by the design of the individual device. For swivel connectors or antennas, the test orientations and configurations will need to be considered on a case-by-case basis because there could be various swiveling combinations and locking mechanisms involved that can affect test considerations. The grantee or its test lab should submit a KDB inquiry to determine the test requirements before conducting certification to support the test procedures.

USB dongles have a rather small footprint; therefore, smaller SAR scan resolutions may be necessary, as compared to the typically larger resolutions used for cellphones, to keep the uncertainty of the interpolation, USB dongle will need to be embedded in several cm of Styrofoam to position the dongle for SAR testing.



14. TEST CONFIGURATION POSITIONS AND POWER TABLE

Summary of SAR Test Configuration

DUT Orientation	Host PC or USB Cable (Length)	Separation [mm]	Remark
Horizontal-Up	Toshiba M100	5.0	FCC DoC Approved
Horizontal-Down	Extension Cable (12 cm)	5.0	N/A
Vertical-Front	Extension Cable (12 cm)	5.0	N/A
Vertical-Back	Extension Cable (12 cm)	5.0	N/A
Bottom	Extension Cable (12 cm)	5.0	N/A
Тор	Extension Cable (12 cm)	5.0	N/A

Max. Power Output Table for SEC-8089 (W/ Short USB Cable)

				1X RRT		EvDo		EvDo		
		RC1	RC1	RC3	RC3	RC3	(Re	v.0)	(Rev.A)	
Band	Channel					SO32				
		SO2	SO55	SO2	SO55	(TDS	FTAP	RTAP	FETAP	RETAP
						0)				
	1013	23.86	23.89	23.70	23.78	23.72	23.75	23.88	23.83	23.88
Cellular	0384	24.72	24.86	24.66	24.64	24.65	24.67	24.72	24.62	24.72
	0777	22.76	22.83	22.73	22.66	22.63	22.92	23.02	23.12	22.94
	0025	24.46	24.47	24.39	24.40	24.39	24.42	24.40	24.44	24.39
PCS	0600	24.95	24.96	24.86	24.90	24.93	24.42	24.82	24.04	24.05
	1175	23.33	23.54	23.42	23.27	23.38	23.46	23.42	23.39	23.13

NOTE1: There was no power deviation between the power when using the short USB cable and the power when direct plug into USB port of the Laptop PC.

Mixture Type: 835 MHz Body PC: Toshiba Laptop

15.1 MEASUREMENT RESULTS (CDMA Cellular Body SAR)							
Application	FREQUENCY		Begin Power	End Power	Device Test	SAR	
Application	MHz	Ch	(dBm)	(dBm)	Position	(W/kg)	
	836.52	384	24.67	24.34	5.0mm [H_UP]	0.707	
	824.70	1013	23.88	23.87	5.0mm [H-DOWN]	0.966	
	836.52	384	24.72	24.61	5.0mm [H-DOWN]	0.760	
E)/DO D O	848.31	777	23.02	23.01	5.0mm [H-DOWN]	0.313	
EVDO Rev. O	836.52	384	24.72	24.46	5.0 mm [V-Front]	0.516	
	836.52	384	24.72	24.71	5.0 mm [V-Back]	0.347	
	836.52	384	24.72	24.68	5.0mm [Bottom]	0.100	
	836.52	384	24.72	24.45	5.0mm [Top]	0.103	
	Body 1.6 W/kg (mW/g) averaged over 1 gram						

- 1. Test procedures used are according to FCC/OET Bulletin 65, Supp.C [July 2001] and FCC 3G SAR MEASUREMENT PROCEDURES - OCT. 2007.
- 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.Test Signal Call Mode
- □ Continuous Tx On □ Manu.Test Codes
- BaseStation Simulator
- 6. Tissue parameters and temperatures are listed on the SAR plots.
- 7. Liquid tissue depth is 15.0 cm.±0.1
- 8. Laptop M/N: Toshiba M100 S/N: 57271209K

Mixture Type: 835 MHz Body PC: Toshiba Laptop

15.2 MEASUR	15.2 MEASUREMENT RESULTS (CDMA Cellular Body SAR)								
Application	FREQU	JENCY	Begin Power	End Power	Device Test	SAR			
Application	MHz	Ch	(dBm)	(dBm)	Position	(W/kg)			
	824.70	1013	23.72	23.73	5.0mm [H_UP]	0.971			
	836.52	384	24.65	24.52	5.0mm [H_UP]	0.965			
	848.31	777	22.63	22.67	5.0mm [H_UP]	0.304			
	824.70	1013	23.72	23.76	5.0mm [H-DOWN]	1.120			
	836.52	384	24.65	24.66	5.0mm [H-DOWN]	1.030			
CDMA 2000	848.31	777	22.63	22.63	5.0mm [H-DOWN]	0.461			
	836.52	384	24.65	24.45	5.0 mm [V-Front]	0.602			
	836.52	384	24.65	24.35	5.0 mm [V-Back]	0.498			
	836.52	384	24.65	24.59	5.0mm [Bottom]	0.150			
	836.52	384	24.65	24.39	5.0mm [Top]	0.123			
	Body 1.6 W/kg (mW/g) averaged over 1 gram								

- 1. Test procedures used are according to FCC/OET Bulletin 65, Supp.C [July 2001] and FCC 3G SAR MEASUREMENT PROCEDURES OCT. 2007.
- 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.Test Signal Call Mode
- Continuous Tx On
- Manu.Test Codes
- BaseStation Simulator
- 6. Tissue parameters and temperatures are listed on the SAR plots.
- 7. Liquid tissue depth is 15.0 cm.±0.1
- 8. Laptop M/N: Toshiba M100 S/N: 57271209K
- 9. Body SAR for CDMA mode was tested under RC3/SO32

Mixture Type: 835 MHz Body PC: Toshiba Laptop

15.3 MEASUREMENT RESULTS (CDMA Cellular Body SAR)						
Application	FREQUENCY		Begin Power	End Power	Device Test	SAR
	MHz	Ch	(dBm)	(dBm)	Position	(W/kg)
	824.70	1013	23.88	23.81	5.0mm [H-DOWN]	1.090
EVDO Rev. A	836.52	384	24.72	24.71	5.0mm [H-DOWN]	0.717
	848.31	777	22.94	22.95	5.0mm [H-DOWN]	0.347
	Body 1.6 W/kg (mW/g) averaged over 1 gram					

- 1. Test procedures used are according to FCC/OET Bulletin 65, Supp.C [July 2001] and FCC 3G SAR MEASUREMENT PROCEDURES OCT. 2007.
- 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.Test Signal Call Mode

 Continuous Tx On

 Manu.Test Codes

 BaseStation Simulator
- 6. Tissue parameters and temperatures are listed on the SAR plots.
- 7. Liquid tissue depth is 15.0 cm.±0.1
- 8. Laptop M/N: Toshiba M100 S/N: 57271209K
- 9. The SAR tests were repeated with the worst case configuration of Rev.0 and low, middle, high channels.

Mixture Type: 1900 MHz Body PC: Toshiba Laptop

15.4 MEASUREMENT RESULTS (CDMA PCS Body SAR)						
Application	FREQUENCY		Begin Power	End Power	Device Test	SAR
	MHz	Ch	(dBm)	(dBm)	Position	(W/kg)
EVDO Rev. O	1880.0	600	24.82	24.81	5.0mm [H_UP]	0.094
	1851.25	25	24.40	24.39	5.0mm [H-DOWN]	0.189
	1880.0	600	24.82	24.83	5.0mm [H-DOWN]	0.117
	1908.75	1175	23.42	23.41	5.0mm [H-DOWN]	0.261
	1880.0	600	24.82	24.83	5.0 mm [V-Front]	0.043
	1880.0	600	24.82	24.68	5.0 mm [V-Back]	0.069
	1880.0	600	24.82	24.56	5.0mm [Bottom]	0.00507
	1880.0	600	24.82	24.84	5.0mm [Top]	0.00879
	Body 1.6 W/kg (mW/g) averaged over 1 gram					

- 1. Test procedures used are according to FCC/OET Bulletin 65, Supp.C [July 2001] and FCC 3G SAR MEASUREMENT PROCEDURES - OCT. 2007.
- 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.Test Signal Call Mode
- □ Continuous Tx On □ Manu.Test Codes
- BaseStation Simulator
- 6. Tissue parameters and temperatures are listed on the SAR plots.
- 7. Liquid tissue depth is 15.0 cm.±0.1
- 8. Laptop M/N: Toshiba M100 S/N: 57271209K

Mixture Type: 1900 MHz Body PC: Toshiba Laptop

15.5 MEASUREMENT RESULTS (CDMA PCS Body SAR)						
Application	FREQUENCY		Begin Power	End Power	Device Test	SAR
	MHz	Ch	(dBm)	(dBm)	Position	(W/kg)
	1880.0	600	24.93	24.73	5.0mm [H_UP]	0.093
	1851.25	25	24.39	24.38	5.0mm [H-DOWN]	0.188
	1880.0	600	24.93	24.78	5.0mm [H-DOWN]	0.100
CDMA 2000	1908.75	1175	23.38	23.24	5.0mm [H-DOWN]	0.217
CDMA 2000	1880.0	600	24.93	24.91	5.0 mm [V-Front]	0.043
	1880.0	600	24.93	24.72	5.0 mm [V-Back]	0.067
	1880.0	600	24.93	24.79	5.0mm [Bottom]	0.00666
	1880.0	600	24.93	24.93	5.0mm [Top]	0.00876
	Body 1.6 W/kg (mW/g) averaged over 1 gram					

- 1. Test procedures used are according to FCC/OET Bulletin 65, Supp.C [July 2001] and FCC 3G SAR MEASUREMENT PROCEDURES - OCT. 2007.
- 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.Test Signal Call Mode
- □ Continuous Tx On □ Manu.Test Codes
- BaseStation Simulator
- 6. Tissue parameters and temperatures are listed on the SAR plots.
- 7. Liquid tissue depth is 15.0 cm.±0.1
- 8. Laptop M/N: Toshiba M100 S/N: 57271209K
- 9. Body SAR for CDMA mode was tested under RC3/SO32

Mixture Type: 1900 MHz Body PC: Toshiba Laptop

15.6 MEASUREMENT RESULTS (CDMA PCS Body SAR)						
Application	FREQUENCY		Begin Power	End Power	Device Test	SAR
	MHz	Ch	(dBm)	(dBm)	Position	(W/kg)
EVDO Rev. A	1851.25	25	24.44	24.42	5.0mm [H-DOWN]	0.184
	1880.0	600	24.04	24.00	5.0mm [H-DOWN]	0.095
	1908.75	1175	23.39	23.14	5.0mm [H-DOWN]	0.242
	Body 1.6 W/kg (mW/g) averaged over 1 gram					

NOTE:

- 1. Test procedures used are according to FCC/OET Bulletin 65, Supp.C [July 2001] and FCC 3G SAR MEASUREMENT PROCEDURES OCT. 2007.
- 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.Test Signal Call Mode

 Continuous Tx On

 Manu.Test Codes
- 6. Tissue parameters and temperatures are listed on the SAR plots.
- 7. Liquid tissue depth is 15.0 cm.±0.1
- 8. Laptop M/N: Toshiba M100 S/N: 57271209K
- 9. The SAR tests were repeated with the worst case configuration of Rev.0 and low, middle, high channels.

BaseStation Simulator

16. SAR TEST EQUIPMENT

Table 16.1 Test Equipment Calibration

EQUIPMENT SPECIFICATIONS					
Туре	Calibration Date	Next	Serial Number		
		Calibration Date			
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	N/A	N/A	21/4		
Celeron ,Window 2000			N/A		
Data Acquisition Electronics	November 12, 2008	November 12, 2009	519		
Dosimetric E-Field Probe	January 14, 2009	January 14, 2010	3643		
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	January 28, 2006	January 28, 2010	464		
SPEAG Validation Dipole D1900 MHz	March 17, 2009	March 17 2011	5d029		
Head/Body Equivalent Matter(835 MHz)	January 2009	January 2010	N/A		
Head/Body Equivalent Matter(1900 MHz)	January 2009	January 2010	N/A		
HP EPM-442A Power Meter	March 13, 2009	March 13, 2010	GB37170267		
HP E4421A Signal Generator	July 09, 2008	July 09, 2009	US37230529		
Attenuator (10dB)	January 19, 2009	January 19 2010	BP4387		
Attenuator (3dB)	July 15, 2008	July 15, 2009	MY39260700		
Low pass filter (1.5GHz)	February 03, 2009	February 03, 2010	N/A		
Low pass filter (3.0GHz)	October 01, 2008	October 01, 2009	N/A		
Dual Directional Coupler	February 03, 2009	February 03, 2010	50228		
Amplifier	February 02, 2009	February 02, 2010	1020 D/C 0221		
Network Analyzer	March 13, 2009	March 13, 2010	8753D		
HP85070D Dielectric Probe Kit	N/A	N/A	LISO1440118		
SEMITEC Engineering	N/A	N/A	Shield Room		

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.

17. 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 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 in possible 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.

18. REFERENCES

- [1] Federal Communications Commission, ET Docket 93-62, Guidelines for Evaluating the Environmental Effects of Radiofrequency Radiation, Aug. 1996.
- [2] ANSI/IEEE C95.1 1991, American National Standard safety levels with respect to human exposure to radio frequency electromagnetic fields, 300kHz to 100GHz, New York: IEEE, Aug. 1992.
- [3] ANSI/IEEE C95.3 1991, IEEE Recommended Practice for the Measurement of Potentially Hazardous Electromagnetic Fields RF and Microwave, New York: IEEE, 1992.
- [4] Federal Communications Commission, OET Bulletin 65 (Edition 97-01), Supplement C (Edition 01-01), Evaluating Compliance with FCC Guidelines for Human Exposure to Radiofrequency Electromagnetic Fields, July 2001.
- [5] IEEE Standards Coordinating Committee 34 IEEE Std. 1528-2003, Recommended Practice for Determining the Peak Spatial-Average Specific Absorption Rate (SAR) in the Human Body Due to Wireless Communications Devices: Experimental Techniques.
- [6] NCRP, National Council on Radiation Protection and Measurements, Biological Effects and Exposure Criteria for Radio Frequency Electromagnetic Fields, NCRP Report No. 86, 1986. Reprinted Feb. 1995.
- [7] T. Schmid, O. Egger, N. Kuster, Automated E-field scanning system for dosimetric assessments, IEEE transaction on Microwave Theory and Techniques, vol. 44, Jan. 1996, pp. 105-113.
- [8] K. Pokovic, T. Schmid, N. Kuster, Robust setup for precise calibration of E-field probes in tissue simulating liquids at mobile communications frequencies, ICECOM97, Oct. 1997, pp. 120-124.
- [9] K. Polovċ,T.Schmid,and N.Kuster ,E-field Probe with improved isotropy in brain simulating liquids. Proceedings of the ELMAR, Zadar, Croatia, June 23-25, 1996, pp. 172-175.
- [10] Schmid & Partner Engineering AG, Application Note: Data Storage and Evaluation, June 1998, p2.
- [11] V. Hombach, K. Meier, M. Burkhardt, E. Kuhn, N. Kuster, The Dependence of EM Energy Absorption upon Human Head Modeling at 900 MHz, IEEE Transaction on Microwave Theory and Techniques, vol. 44 no. 10, Oct. 1996, pp. 1865-1873.
- [12] N. Kuster and Q. Balzano, Energy absorption mechanism by biological bodies in the near field of dipole antennas above 300MHz, IEEE Transaction on Vehicular Technology, vol. 41, no. 1, Feb. 1992, pp. 17-23.
- [13] G. Hartsgrove, A. Kraszewski, A. Surowiec, Simulated Biological Materials for Electromagnetic Radiation Absorption Studies, University of Ottawa, Bioelectromagnetics, Canada: 1987, pp. 29-36.
- [14] Q. Balzano, O. Garay, T. Manning Jr., Electromagnetic Energy Exposure of Simulated Users of Portable Cellular Telephones, IEEE Transactions on Vehicular Technology, vol. 44, no.3, Aug. 1995.
- [15] W. Gander, Computermathematick, Birkhaeuser, Basel, 1992.
- [16] W.H. Press, S.A. Teukolsky, W.T. Vetterling, and B.P. Flannery, Numerical Recepies in C, The Art of Scientific Computing, Second edition, Cambridge University Press, 1992.
- [17] Federal Communications Commission, OET Bulletin 65, Evaluating Compliance with FCC Guidelines for Human Exposure to Radiofrequency Electromagnetic Fields. Supplement C, Dec. 1997.
- [18] N. Kuster, R. Kastle, T. Schmid, Dosimetric evaluation of mobile communications equipment with known precision, IEEE Transaction on Communications, vol. E80-B, no. 5, May 1997, pp. 645-652.
- [19] CENELEC CLC/SC111B, European Prestandard (prENV 50166-2), Human Exposure to Electromagnetic Fields High-frequency: 10kHz-300GHz, Jan. 1995.
- [20] Prof. Dr. Niels Kuster ,ETH,EidgenÖssische Technische Technische Hoschschule Zűrich,Dosimetric Evaluation of the Cellular Phone.
- [21] FCC SAR Measurement Procedures for 3G Devices v2.0, October 2007
- [22] FCC SAR Considerations for Cell Phones with Multiple Transmitters v01r02 #648474, April 2008
- [23] 447498 DO1 Mobile Portable RF Exposure v03r02, Published on: Jul 28 2008
- [24] 447498 DO2 Mobile Portable RF Exposure v03r02, Published on: Dec 02 2008