

in accordance with the requirements of FCC Report and Order: ET Docket 93-62, and OET Bulletin 65 Supplement C

for

#### **DSSS TOUCH PANEL TRANSCEIVER**

**MODEL: CF-VDW07** 

FCC ID: N/A

**OCTOBER 11, 2001** 

**REPORT NO: 01U1008-1** 

Prepared for

MATSUSHITA ELECTRIC INDUSTRIAL CO., LTD. 1006 OAZA KADOMA KADOMA OSAKA JAPAN 571

Prepared by

COMPLIANCE CERTIFICATION SERVICES 561F MONTEREY ROAD, MORGAN HILL, CA 95037, USA TEL: (408) 463-0885



# **CERTIFICATE OF COMPLIANCE (SAR EVALUATION)**

Dates of Tests: October 10-11, 2001 Report No: 01U1008-1

APPLICANT: MATSUSHITA ELECTRIC INDUSTRIAL CO., LTD.

1006 OAZA KADOMA

KADOMA OSAKA JAPAN 571

TRADE NAME: PANASONIC MODEL: CF-VDW07

SERIAL NUMBER: N/A (PRE-PRODUCTION)

FCC ID: N/A

CATEGORY: MOBILE/PORTABLE DSSS DEVICE

Test Sample is a: Pre-Production Unit
Tx Frequency: 2412 – 2462 MHz (CDMA)

Max. RF Output Power: 13mW based on Manufacture's Information

FCC Classification: Unlicensed Intentional Radiator RF Exposure environment: General Population/Uncontrolled

Application Type: Certification FCC Rule Part(s): § 15.247

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 65 Supplement C (released on 6/29/2001 see Test Report).



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.

**Steve Cheng** 

**EMC Engineering Manager** 

NVLAP accreditation does not constitute any product endorsement by NVLAP or any agency of the United States Government. CCS certifies that no party to this application has been denied the FCC benefits pursuant to Section 5301 of the Anti-Drug Abuse Act of 1988, 21 U.S.C. 853(a).

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#### 1. EUT DESCRIPTION

**APPLICANT:** MATSUSHITA ELECTRIC INDUSTRIAL CO., LTD.

DATE: October 11.2001

1006 OAZA KADOMA

KADOMA OSAKA JAPAN 571

TRADE NAME: **PANASONIC** MODEL: CF-VDW07

SERIAL NUMBER: N/A (PRE-PRODUCTION)

FCC ID: N/A

**CATEGORY:** MOBILE/PORTABLE DSSS DEVICE

Test Sample is a: Pre-production Unit

EUT Type: Mobile/Portable PC with 2.4GHz Direct Sequence Spread Spectrum Transceiver

Trade Name: **PANASONIC** Model(s): CF-VDW07

FCC IDENTIFIER: N/A

S/N: N/A (Pre-Production) Tx Frequency: 2412 - 2462 MHz (CDMA)

Application Type: Certification

FCC Classification: Unlicensed Intentional Radiator

Modulation(s): **CDMA** FCC Rule Part(s): § 15.247

Max. RF Output Power: 13mW Based on Manufacture Information

Antenna Type: Built-in Antenna Dimensions: N/A

Dates of Tests: October 11, 2001 Report Project No.: 01U1008-1



<sup>1</sup> Specific Absorption Rate (SAR) is a measure of the rate of energy absorption due to exposure to an RF transmitting source (wireless portable device).

<sup>&</sup>lt;sup>2</sup> IEEE/ANSI Std. C95.1-1992 limits are used to determine compliance with FCC ET Docket 93-62.

# 2. REQUIREMENTS FOR COMPLIANCE TESTING DEFINED BY THE **FCC**

The US Federal Communications Commission has released the report and order "Guidelines for Evaluating the Environmental Effects of RF Radiation", ET Docket No. 93-62 in August 1996 [1]. The order requires routine SAR evaluation prior to equipment authorization of portable transmitter devices, including portable telephones. For consumer products, the applicable limit is 1.6 mW/g as recommended by the ANSI/IEEE standard C95.1-1992 [6] for an uncontrolled environment (Paragraph 65). According to the Supplement C of OET Bulletin 65 "Evaluating Compliance with FCC Guide-lines for Human Exposure to Radio frequency Electromagnetic Fields", released on Jun 29, 2001 by the FCC, the device should be evaluated at maximum output power (radiated from the antenna) under "worst-case" conditions for normal or intended use, incorporating normal antenna operating positions, device peak performance frequencies and positions for maximum RF energy coupling.

#### 3. DOSIMETRIC ASSESSMENT SETUP

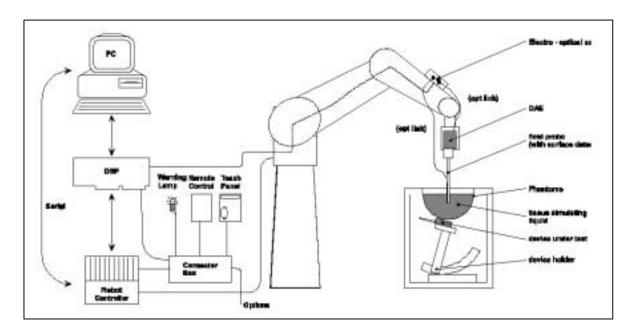
These measurements were performed with the automated near-field scanning system DASY3 from Schmid & Partner Engineering AG (SPEAG). The system is based on a high precision robot (working range greater than 0.9 m) which positions the probes with a positional repeatability of better than ± 0.02 mm. Special E- and H-field probes have been developed for measurements close to material discontinuity, the sensors of which are directly loaded with a Schottky diode and connected via highly resistive lines to the data acquisition unit. The system is described in detail in [3].

The SAR measurements were conducted with the dosimetric probe ET3DV6 SN: 1577 (manufactured by SPEAG), designed in the classical triangular configuration [3] and optimized for dosimetric evaluation. The probe has been calibrated according to the procedure described in [7] with accuracy of better than ±10%. The spherical isotropy was evaluated with the procedure described in [8] and found to be better than ±0.25 dB.

The phantom used was the \Generic Twin Phantom" described in [4]. The ear was simulated as a spacer of 4 mm thickness between the earpiece of the phone and the tissue simulating liquid. The Tissue simulation liquid used for each test is in according with the FCC OET65 supplement C as listed below.

Ingredients	Frequency (MHz)									
(% by weight)	45	50	835		915		1900		2450	
Tissue Type	Head	Body	Head	Body	Head	Body	Head	Body	Head	Body
Water	38.56	51.16	41.45	52.4	41.05	56.0	54.9	40.4	62.7	73.2
Salt (NaCl)	3.95	1.49	1.45	1.4	1.35	0.76	0.18	0.5	0.5	0.04
Sugar	56.32	46.78	56.0	45.0	56.5	41.76	0.0	58.0	0.0	0.0
HEC	0.98	0.52	1.0	1.0	1.0	1.21	0.0	1.0	0.0	0.0
Bactericide	0.19	0.05	0.1	0.1	0.1	0.27	0.0	0.1	0.0	0.0
Triton X-100	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	36.8	0.0
DGBE	0.0	0.0	0.0	0.0	0.0	0.0	44.92	0.0	0.0	26.7
Dielectric Constant	43.42	58.0	42.54	56.1	42.0	56.8	39.9	54.0	39.8	52.5
Conductivity (S/m)	0.85	0.83	0.91	0.95	1.0	1.07	1.42	1.45	1.88	1.78

#### 3.1. MEASUREMENT SYSTEM DIAGRAM



#### The DASY3 system for performing compliance tests consist of the following items:

- 1. A standard high precision 6-axis robot (Stäubli RX family) with controller and software.
- 2. An arm extension for accommodating the data acquisition electronics (DAE).
- 3. A dosimetric probe, i.e., an isotropic E-field probe optimized and calibrated for usage in tissue simulating liquid. The probe is equipped with an optical surface detector system.
- 4. A data acquisition electronic (DAE), which performs the signal amplification, signal multiplexing, AD-conversion, offset measurements, mechanical surface detection, collision detection, etc. The unit is battery powered with standard or rechargeable batteries. The signal is optically transmitted to the EOC.
- 5. A unit to operate the optical surface detector, which is connected to the EOC. The Electro-optical coupler (EOC) performs the conversion from the optical into a digital electric signal of the DAE. The EOC is connected to the PC plug-in card. The functions of the PC plug-in card based on a DSP is to perform the time critical task such as signal filtering, surveillance of the robot operation fast movement interrupts.
- 6. A computer operating Windows 95 or larger
- 7. DASY3 software
- 8. Remote control with teaches pendant and additional circuitry for robot safety such as warning lamps, etc.
- The generic twin phantom enabling testing left-hand and right-hand usage.
- 10. The device holder for handheld EUT.
- 11. Tissue simulating liquid mixed according to the given recipes (see Application Note).
- 12. System validation dipoles to validate the proper functioning of the system.

#### 3.2. SYSTEM COMPONENTS

#### **ET3DV5 Probe Specification**

Construction Symmetrical design with triangular core Built-in optical fiber for surface detection System Built-in shielding against static charges Calibration In air from 10 MHz to 2.5 GHz In brain and muscle simulating tissue at Frequencies of 450 MHz, 900 MHz and 1.8 GHz (accuracy  $\pm\,8\%)$ 

Frequency 10 MHz to > 6 GHz; Linearity:  $\pm$  0.2 dB

(30 MHz to 3 GHz)

Directivity  $\pm\,0.2$  dB in brain tissue (rotation around probe axis)

 $\pm$  0.4 dB in brain tissue (rotation normal probe axis)

Dynamic 5 mW/g to > 100 mW/g;

Range Linearity: ± 0.2 dB

Surface  $\pm\,0.2$  mm repeatability in air and clear liquids

Detection over diffuse reflecting surfaces. Dimensions Overall length: 330 mm

Tip length: 16 mm Body diameter: 12 mm Tip diameter: 6.8 mm

Distance from probe tip to dipole centers: 2.7 mm Application General dosimetric up to 3 GHz

Compliance tests of mobile phones

Fast automatic scanning in arbitrary phantoms

The SAR measurements were conducted with the dosimetric probe ET3DV6 designed in the classical triangular configuration 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. 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 DASY3 software reads the reflection during a software approach and looks for the maximum using a 2 nd order fitting. The approach is stopped when reaching the maximum.



Figure. Photograph of the probe



Figure. Inside view of ET3DV6 E-field Probe

#### **E-Field Probe Calibration Process**

Each probe is calibrated according to a dosimetric assessment procedure described in [6] with accuracy better than +/- 10%. The spherical isotropy was evaluated with the procedure described in [7] 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 are tested.

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The free space E-field from amplified probe outputs is determined in a test chamber. This is performed in a TEM cell for frequencies bellow 1 GHz, and in a waveguide above 1 GHz for free space. For the free space calibration, the probe is placed in the volumetric center of the cavity and at the proper orientation with the field. The probe is then rotated 360 degrees.

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 dielectric medium. For temperature correlation calibration a RF transparent thermistor-based temperature probe is used in conjunction with the E-field probe.

#### **Data Evaluation**

Media parameters:

The DASY3 software automatically executes the following procedures to calculate the field units from the microvolt readings at the probe connector. The parameters used in the evaluation are stored in the configuration modules of the software:

Probe parameters: - Sensitivity Norm<sub>i</sub>, a<sub>i0</sub>, a<sub>i1</sub>, a<sub>i2</sub>

> - Conversion factor ConvF<sub>i</sub> - Diode compression point Dcp<sub>i</sub>

Device parameters: - Frequency

> Crest factor cf - Conductivity σ

- Density

These parameters must be set correctly in the software. They can either be found in the component documents or be imported into the software from the configuration files issued for the DASY3 components. In the direct measuring mode of the multi-meter option, the parameters of the actual system setup are used. In the scan visualization and export modes, the parameters stored in the corresponding document files are used.

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 as:

$$V_i = U_i + U_i^2 \cdot \frac{cf}{dcp_i}$$

= compensated signal of channel i with  $V_i$ (i = x, y, z)

= input signal of channel i  $U_i$ (i = x, y, z)

cf = crest factor of exciting field (DASY parameter) (DASY parameter) = diode compression point

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

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 $E_i = \sqrt{\frac{V_i}{Norm_i \cdot ConvF}}$ E-field probes:

 $H_i = \sqrt{Vi} \cdot \frac{a_{i0} + a_{i1}f + a_{i2}f^2}{f}$ H-field probes:

V<sub>i</sub> = compensated signal of channel i (i = x, y, z) with

Norm<sub>i</sub> = sensor sensitivity of channel i (i = x, y, z)

μV/(V/m)2 for E-field Probes

ConvF = sensitivity enhancement in solution

= sensor sensitivity factors for H-field probes

= carrier frequency [GHz]

= electric field strength of channel i in V/m  $E_i$ = magnetic field strength of channel i in A/m

The RSS value of the field components gives the total field strength (Hermitian 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.

$$SAR = E_{tot}^{2} \cdot \frac{\sigma}{\rho \cdot 1000}$$

SAR = local specific absorption rate in mW/g with

= total field strength in V/m

= conductivity in [mho/m] or [Siemens/m]

= equivalent tissue density in g/cm<sup>3</sup>

Note that the density is normally set to 1 (or 1.06), to account for actual brain density rather than the density of the simulation liquid.

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

$$P_{pwe} = \frac{E_{tot}^2}{3770}$$
 or  $P_{pwe} = H_{tot}^2 \cdot 37.7$ 

= equivalent power density of a plane wave in mW/cm<sup>2</sup> with

= total electric field strength in V/m = total magnetic field strength in A/m

#### **Generic Twin Phantom**

The Generic Twin Phantom 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 [9][10]. 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 allows the complete setup of all predefined phantom positions and measurement grids by manually teaching three points in the robot. See Figure 8. Shell Thickness  $2 \pm 0.1$  mm Filling Volume Approx. 20 liters Dimensions 810 x 1000 x 500 mm (H x L x W)



Fig. Generic Twin Phantom

#### **Device Holder for Transmitters**

In combination with the Generic Twin Phantom V3.0, the Mounting Device enables the rotation of the mounted transmitter in spherical coordinates whereby the rotation points is the ear opening. The devices can be easily, accurately, and repeatedly positioned according to the FCC and CENELEC 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 produced infinite number of configurations [10]. To produce the worst-case condition (the hand absorbs antenna output power), the hand is omitted during the tests.



Fig. Device Holder

#### 3.3. EUT ARRANGEMENT

# **Body Carrying Configuration**

The body carry configuration is used for body-worn devices that have a body-carrying accessory. Typically, a holster or carrying case is provided or available as an accessory item for supporting portable operations. SAR may vary depending on the body separation distance provided by the type of carrying accessory. Proper usage of the carrying case restricts the antenna to a specified distance away from the surface of the body. For this test the EUT is placed directly on the device holder and the device holder is positioned against the flat portion of the phantom in a normal operating position.



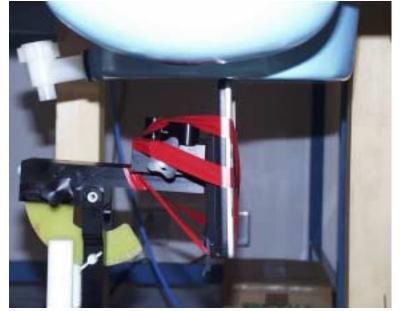


Figure. Normal operation position



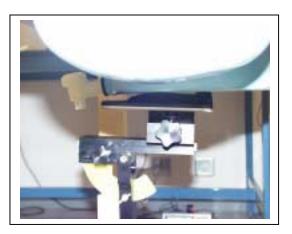


Figure. Abnormal test position for reference only (When EUT bottom side face the stomach)



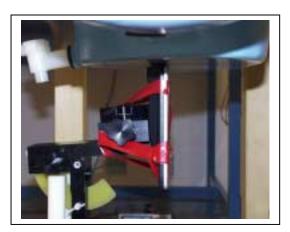


Figure. Abnormal position for reference only (Worst case when antenna side face the stomach)

# **Measurement Uncertainty**

The uncertainty budget has been determined for the DASY3 measurement system according to the NIS81 [13] and the NIST1297 [14] documents and is given in the following Table.

Uncertainty Description	Error	Distrib.	Weight	Std. Dev.	Offset						
	Probe Uncertainty										
Axial isotropy	± 0.2 dB	U-shape	0.5	±2.4 %							
Spherical isotropy	±0.4 dB	U-shape	0.5	±4.8 %							
Isotropy from gradient	±0.5 dB	U-shape	0								
Spatial resolution	±0.5 %	Normal	1	±0.5 %							
Linearity error	±0.2 dB	Rectangle	1	±2.7 %							
Calibration error	±3.3 %	Normal	1	± 3.3 %							
	SAR Evaluation	Uncertainty									
Data acquisition error	±1%	Rectangle	1	±0.6 %							
ELF and RF disturbances	±0.25 %	Normal	1	±0.25 %							
Conductivity assessment	±10 %	Rectangle	1	± 5.8 %							
Spatia	al Peak SAR Eva	aluation Uncert	ainty								
Extrapol boundary effect	±3%	Normal	1	±3%	± 5%						
Probe positioning error	±0.1 mm	Normal	1	± 1%							
Integrat. and cube orient	±3%	Normal	1	±3%							
Cube shape inaccuracies	±2%	Rectangle	1	±1.2 %							
Device positioning	±6%	Normal	1	± 6%							
Combined Uncertainties			1	±11.7 %	± 5%						
Extended uncertainty (K = 2)				± 23.5 %.							

#### 4. EUT TUNE-UP PROCEDURE

The following procedures had been used to prepare the EUT for the SAR test as directed by the Manufacture.

- 1. Connect Display (CF-VDW07) with Host PC (IBM) by Serial cable
- 2. Connect AC adopter (02K6542) with Host PC and start it up.
- 3. Click "CF-VDW07" icon on screen of Host PC.
- Hyper Terminal will be started.
- 4. Press "MENU/CAPTURE/R-BUTTON" and connect AC adaptor with Display (CF-VDW07) simultaneously. "DBG>" will be appeared on the screen of Host PC, but nothing will be shown on the Display (CF-VDW07).
- 5. Input "INI" and press ENTER key. (Initialize)
- 6. Input "Mod 11" and press ENTER key (Set TX Rate at the maximum, 11 Mbs )
- "Current tx rate:11" is displayed.
- 7. Input "cch X" and press ENTER key (X will be replaced by any number from 1 to 11)
- Current channel (chosen channel) will be displayed.
- 8. Input "CSD" and press ENTER key.
- CW signal will start.
- 9. "CSD Start!!!" will be displayed on the screen of Host PC, but nothing will be on the Display (CF-VDW07).
- 10. To stop CW signal, please input "STP "and press ENTER key.

Note: Please perform the testing after serial cable is disconnected from Host PC (IBM) and Display (CF-VDW07).

COMPLIANCE CERTIFICATION SERVICES

#### 5. EVALUATION PROCEDURE

#### 5.1. SIMULATED TISSUE LIQUID PARAMETER CONFIRMATION

The dielectric parameters were checked prior to assessment using the HP85070A dielectric probe kit. The dielectric parameters measured are reported in each correspondent section:

#### 5.2. SYSTEM ACCURACY VERIFICATION

Prior to the assessment, the system validation kit was used to test whether the system was operating within its specifications of  $\pm 10\%$ . The validation results are tabulated below. And also the corresponding SAR plot is attached as well in the SAR plots files.

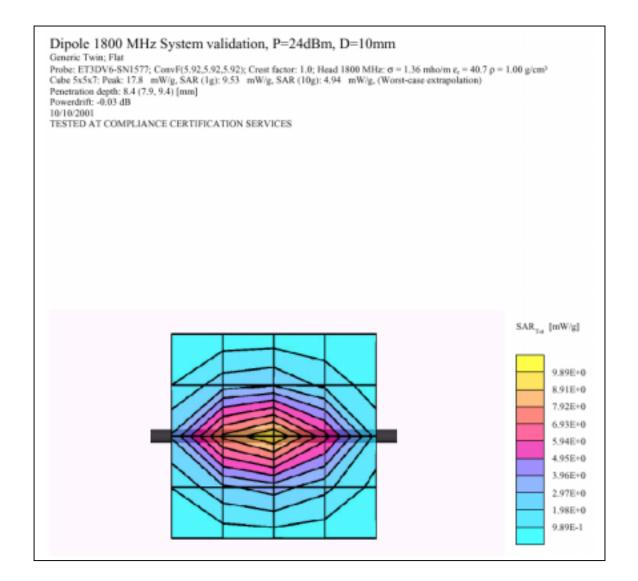
IEEE P1528 recommended reference value

Frequency (MHz)	1 g SAR	10 g SAR	local SAR at surface (above feedpoint)	local SAR at surface (y=2cm offset from feedpoint)
300	3.0	2.0	4.4	2.1
450	4.9	3.3	7.2	3.2
835	9.5	6.2	14.1	4.9
900	10.8	6.9	16.4	5.4
1450	29.0	16.0	50.2	6.5
1800	38.1	19.8	69.5	6.8
1900	39.7	20.5	72.1	6.6
2000	41.1	21.1	74.6	6.5
2450	52.4	24.0	104.2	7.7
3000	63.8	25.7	140.2	9.5

#### System validation result

Simulant	Freq [MHz]	Parameters	Liquid Temp [°C]	Target Value	Measured Value	Deviation [%]	Limit [%]		
	1800			3	22.6	40.0	40.7	+1.75	± 5
Muscle		σ	22.6	1.40	1.36	-2.857	± 5		
		1 g SAR	22.6	38.1	38.12	+0.05	± 10		

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#### 5.3. SAR EVALUATION PROCEDURE

#### The evaluation was performed with the following procedure:

- Step 1: Measurement of the SAR value at a fixed location above the ear point or central position was used as a reference value for assessing the power drop.
- Step 2: The SAR distribution at the exposed side of the head was measured at a distance of 3.9 mm from the inner surface of the shell. The area covered the entire dimension of the head or EUT and the horizontal grid spacing was 20 mm x 20 mm. Based on these data, the area of the maximum absorption was determined by spline interpolation.
- Step 3: Around this point, a volume of 32 mm x 32 mm x 34 mm was assessed by measuring 5 x 5 x 7 points. On the basis of this data set, the spatial peak SAR value was evaluated under the following procedure:
- 1. The data at the surface were extrapolated, since the center of the dipoles is 2.7 mm away from the tip of the probe and the distance between the surface and the lowest measuring point is 1.2 mm. The extrapolation was based on a least square algorithm [11]. 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.
- 2. The maximum interpolated value was searched with a straightforward algorithm. Around this maximum the SAR values averaged over the spatial volumes (1 g or 10 g) were computed by the 3D-Spline interpolation algorithm. The 3D-Spline is composed of three onedimensional splines with the "Not a knot"-condition (in x, y and z-directions) [11], [12]. The volume was integrated with the trapezoidal-algorithm. One thousand points (10 x 10 x 10) were interpolated to calculate the average.
- 3. All neighboring volumes were evaluated until no neighboring volume with a higher average value was found.
- Step 4: Re-measurement of the SAR value at the same location as in Step 1. If the value changed by more than 5%, the evaluation was repeated.

#### 5.4. EXPOSURE LIMIT

#### (A) Limits for Occupational/Controlled Exposure (W/kg)

Whole-Body	Partial-Body	Hands, Wrists, Feet and Ankles
0.4	8.0	20.0

#### (B) Limits for General Population/Uncontrolled Exposure (W/kg)

Whole-Body	Partial-Body	Hands, Wrists, Feet and Ankles
0.08	1.6	4.0

NOTE 1: Whole-Body SAR is averaged over the entire body, partial-body SAR is averaged over any 1 gram of tissue defined as a tissue volume in the shape of a cube. SAR for hands, wrists, feet and ankles is averaged over any 10 grams of tissue defined as a tissue volume in the shape of a cube.

<u>Population/Uncontrolled Environments</u>: are defined as locations where there is the exposure of individuals who have no knowledge or control of their exposure. <u>Occupational/Controlled Environments</u>: are defined as locations where there is exposure that may be incurred by people who are aware of the potential for exposure, (i.e. as a result of employment or occupation).

# NOTE: POPULATION/UNCONTROLLED ENVIRONMENTS PARTIAL BODY LIMIT 1.6 mW/g APPLIYED TO THIS PRODUCT

REPORT NO: 01U1008-1 DATE: October 11.2001 EUT: DSSS Touch Panel Transceiver

FCC ID: N/A

# 6. RESULTS

This page summarizes the results of the performed dosimetric evaluation. The plots with the corresponding SAR distributions, which reveal information about the location of the maximum SAR with respect to the device could be found in the following pages.

## **SAR TEST DATA SUMMARY**

Ambient TEMPERATURE (°C): 23.5

Relative HUMIDITY (%): 68

Summary of worst case SAR reading

			_	Conducted	Worst case SAR, averaged over 1g [mW/g]			
Mode	Position	Ch	Frequency [MHz]	Power [dBm]	Set-up condition	Measured	Limit	
SS 2450	Body	М	2437	13mW based on manufacture's information	Normal operation position	0.0042	1.6	

# Measurement Results (2450MHz CDMA Body SAR)

Liquid Measurement date: 10/10/2001 By: Steve Cheng

	1									
Simulant	Freq [MHz]	Parameters	Liquid Temp [°C]	Target Value	Measured Value	Deviation [%]	Limit [%]			
Body	2450	3	22.4	52.7	50.5	-4.17	±5			
		σ	22.4	1.95	2.02	+3.59	±5			

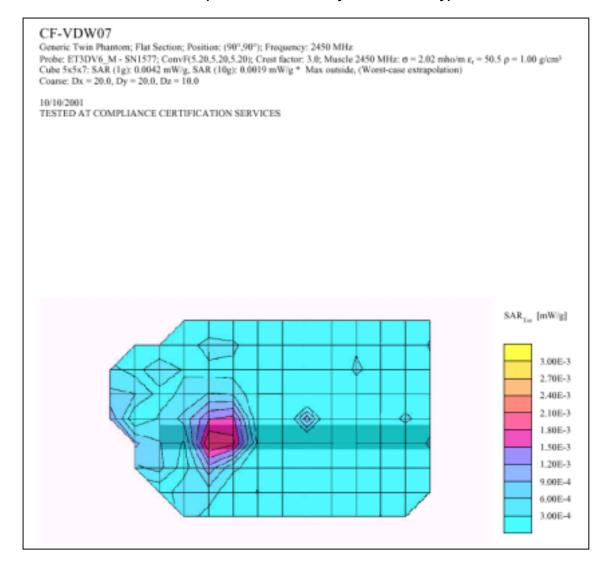
DATE: October 11.2001

#### SAR Measurement result:

			F	Conducted	Worst case SAR, averaged over 1g [mW/g]			
Mode	Position	Ch	Frequency [MHz]	Power [dBm]	Set-up condition	Measured	Limit	
SS 2450	Body	М	2437	13mW based on the manufacture's information	Normal operation position	0.0042	1.6	
SS 2450	Body	М	2437	13mW based on the manufacture's information	Abnormal position for reference only When EUT bottom side face the stomach	0.129	1.6	
SS 2450	Body	М	2437	13mW based on the manufacture's information	Abnormal position for reference only Worst case when ant side face the stomach	0.537	1.6	

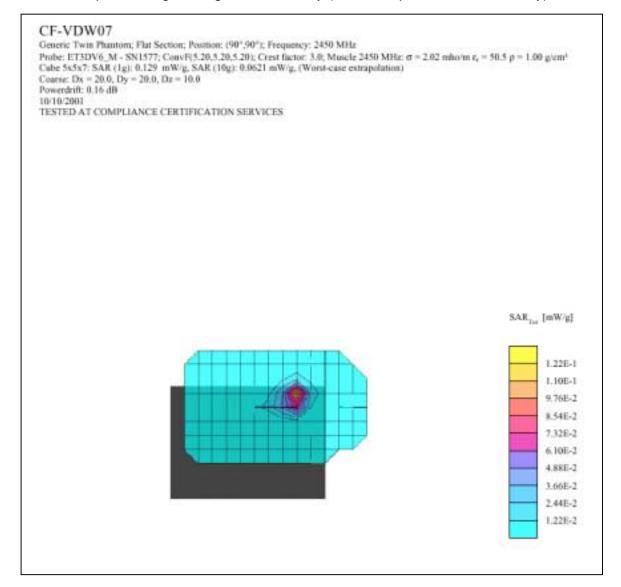
Note: Due to the low SAR (more them 2dB lower them limit), the high and low channels had been optionally skipped.

#### Normal Position Test Plot (EUT Ant-side away from the body)



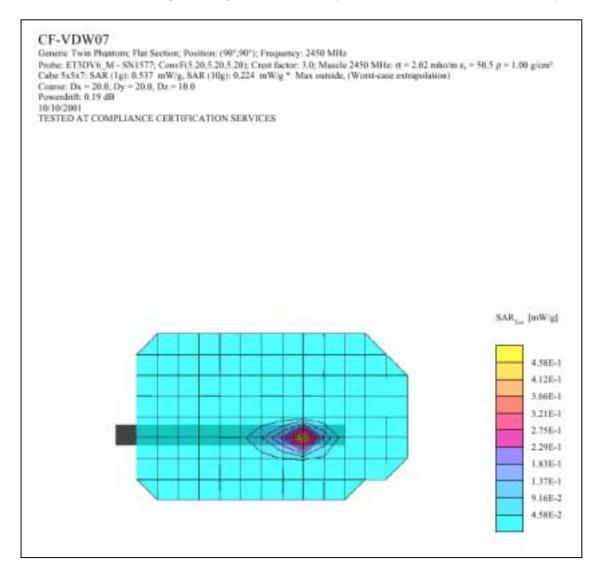
#### **Abnormal Test position**

Reference plot for engineering evaluation only (EUT face/panel toward the body)



#### **Abnormal Test position**

#### Reference plot for engineering evaluation only (EUT Ant side toward the body)



#### 7. REFERENCES

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# 8. APPENDIX

#### 8.1. EUT PHOTOS



# EUT: DSSS Touch Panel Transceiver

# **EQUIPMENTS LIST & CALIBRATION INFO**

Type / Model	Cal. Date	S/N:
DASY3 Professional Dosimetric System	N/A	
Robot RX90BL	N/A	F00/5H31A1/A/01
Robot Controller	N/A	D22134001-1
Teach Pendant	N/A	321
Dell Computer Optiplex GX110	N/A	
Pentium III, Windows NT	N/A	
SPEAG EDC3	N/A	
SPEAG DAE3	4/27/01	421
SPEAG E-Field Probe ET3DV6	4/20/01	1577
SPEAG E-Field Probe ET3DV6	4/20/01	1578
SPEAG Dummy Probe	N/A	
SPEAG Generic Twin Phantom	N/A	
SPEAG Light Alignment Sensor	N/A	261
SPEAG Validation Dipole D1800V2	4/19/01	294
SPEAG Validation Dipole D900V2	4/17/01	108
Brain Equivalent Matter (800MHz)	Daily	
Brain Equivalent Matter (1900MHz)	Daily	
Muscle Equivalent Matter (800MHz)	Daily	
Muscle Equivalent Matter (1900MHz)	Daily	
Robot Table	N/A	
Phone Holder	N/A	
Phantom Cover	N/A	
HP Spectrum Analyzer HP8593GM	6/20/01	3009A00791
Microwave Amp. Model: ZHL-42W	N/A	D072701-5
Power Meter HP436A	4/2/01	2709A29209
Power Sensor HP8482A	4/2/01	2349A08568
Signal Generator HP-83732B	3/21/01	US13449049
Network Analyzer HP-8753ES	7/28/01	MY40001647
Dielectric Probe Kit HP85070A	N/A	

# 8.3. IEEE SCC-34/SC-2 P1528 RECOMMENDED TISSUE DIELECTRIC PARAMETERS

The head tissue dielectric parameters recommended by the IEEE SCC-34/SC-2 in P1528 have been incorporated in the following table. These head parameters are derived from planar layer models simulating the highest expected SAR for the dielectric properties and tissue thickness variations in a human head. Other head and body tissue parameters that have not been specified in P1528 are derived from the tissue dielectric parameters computed from the 4-Cole-Cole equations and extrapolated according to the head parameters specified in P1528.

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

( $\epsilon_r$  = relative permittivity,  $\sigma$  = conductivity and  $\rho$  = 1000 kg/m<sup>3</sup>)

DATE: October 11.2001

# 8.4. EQUIPMENTS CALIBRATION CERTIFICATE

Coughousstrosse 43, 8004 Zurich, Switzerla	nd, Phone +41 1 245 97 00, Fex +41 1 245
Calibratio	n Certificate
Dosimetric	E-Field Probe
Type:	ET3DV6
Serial Number:	1377
Place of Calibration:	Zurich
Date of Calibration:	Apr. 20, 2001
Calibration Interval:	12 months
Schmid & Partner Engineering AG hereby or the date indicated above. The calibration was and procedures of Schmid & Partner Engineer	performed in accordance with specification
Wherever applicable, the standards used in the international standards. In all other cases the dicrowave Electronics at the Swiss Federal liwitzerland have been applied.	standards of the Laboratory for EMF and
Calibrated by:	Mitologia Nervana
Approved by:	21-41

# Schmid & Partner Engineering AG

### DASY - DOSIMETRIC ASSESSMENT SYSTEM

# CALIBRATION REPORT

#### DATA ACQUISITION ELECTRONICS

MODEL:

DAE3 V1

SERIAL NUMBER:

427

This Data Acquisition Unit was calibrated and tested using a FLUKE 702 Process Calibrator. Calibration and verification were performed at an ambient temperature of 23 ± 5 °C and a relative humidity of < 70%.

Measurements were performed using the standard DASY software for converting binary values, offset compensation and noise filtering. Software settings are indicated in the reports.

Results from this calibration relate only to the unit calibrated.

Calibrated by:

E. Meyer

Calibration Date:

April 27, 2001

**DASY Software Version:** 

**DASY3 V3.1c**