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COMPUTATIONAL STUDIES FOR PREDICTION OF ENERGY DEPOSITION IN HUMANS EXPOSED TO RF FIELDS FROM CELLULAR PHONES

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#### **Table of Contents**

#### Computational Studies for Prediction of Energy Deposition in Humans Exposed to RF Fields from Cellular Phones

|      | r i statistica i st                    | age                      |
|------|---|--------------------------|
| Tabl | e of Contents   | . i                      |
| 1.0  | INTRODUCTION  | . 1                      |
| 1.1  | Survey of Computational Electromagnetic Modelling Methods   | . 2                      |
|      | 1.1.1 The Finite-Difference Time-Domain Method1.1.2 Finite-Element Methods1.1.3 Moment Methods1.1.4 The Multiple-Multipole Method | . 2<br>. 2<br>. 2<br>. 2 |
| 1.2  | Applications to Biological Interactions   | . 2                      |
|      | 1.2.1 Antenna Performance in the Presence of an Operator         1.2.2 Exposure to RF Fields         1.2.3 Hyperthermia           | . 2<br>. 3<br>. 3        |
| 1.3  | Electrical Properties Biological Materials  | . 3                      |
| 1.4  | Measurements and Validation   | . 3                      |
| 2.0  | THE FINITE-DIFFERENCE TIME-DOMAIN METHOD  | . 3                      |
| 2.1  | The FDTD Algorithm  | . 3                      |
| 2.2  | FDTD and Radiating Antennas   | . 3                      |
| 2.3  | FDTD with Dispersive Materials  | . 3                      |
| 3.0  | PROGRESS TO DATE AND WORK IN PROGRESS   | . 4                      |
| 3.1  | Modelling of the Cellular Phone   | . 4                      |
| 3.2  | Code Acquisition and Modification   | . 4                      |
| 3.3  | Head Phantoms   | . 4                      |
| Conc | ordia EMC Laboratory  | i                        |

|     |  | rage | 2 |
|-----|--|------|---|
| 4.0 | ACKNOWLEDGEMENTS   | . 4  | ł |
| 5.0 | REFERENCES   | . 5  | 5 |
| 6.0 | APPENDICES   | . 5  | 5 |
|     | Appendix A, Statement of Work  | . 6  | ĵ |
|     | Appendix B, Paper presented at ANTEM '94 Conference, Ottawa,<br>August 2-5, 1994 | . 9  | ) |
| 4   | Appendix C. Presentation Material ANTEM '94                                      | 20   | 1 |

Concordia EMC Laboratory

ii

Computational Studies for the Prediction of Energy Deposition in Humans Exposed to RF Fields from Cellular Phones

#### FINAL REPORT

#### **1.0 INTRODUCTION**

This report is intended to summarize the work that was done in an initial research contract for the study of the methodology which could be applied to the computation of the RF fields associated with the use of cellular telephones. The objectives of the research contract are described in the contractual Statement of Work (SOW) that is enclosed as Appendix A.

Important at the start, was a systematic survey of the state-of-the-art in computational modelling of the transceiver and operator plus an identification of the corresponding measurements which would serve to validate the modelling techniques or which would provide additional quantitative information on the energy deposition or external field distribution aspects.

Vital to any further research, is the acquisition or development of a suitable computer model of the transceiver and the human operator. While probing the availability of existing models, the organization and initiation of the development of a finely-stepped (1 mm), discretized model of the human head was launched. At the same time, we began to plan a series of systematic measurements which could eventually be carried out in Canada at the David Florida Laboratory (DFL) of the Canadian Space Agency (CSA).

The Table of Contents of this report reflects our intentions about its scope. This specific presentation was preempted by the request to participate in a Bio-Session at the ANTEM '94 conference in Ottawa on August 2-5, 1994, and chaired by Dr. Maria Stuchly. This offered us an opportunity to summarize our work preparatory for the technical community at large. Since the contents of the paper duplicate the specific portions of this Final Report, it would have been a wasteful duplication of effort at a time of limited resources to reproduce its contents in a different format. Hence, the published paper is reproduced as Appendix B.

However, to keep the outline of the Table of Contents, the body of this report maintains the individual topic sections and provides brief comments which refer to the appendix material. The oral presentation of the paper included some additional material. Thus hardcopies of the illustrations used for the delivery of the paper are also included as Appendix C. Readers will be able to glean the full scope of what was done by scanning the report in this way.

#### **1.1** Survey of Computational Electromagnetic Modelling Methods

A workable survey of computational modelling methods can be found in the several new books on Computational Electromagnetics. They can be categorized as time-domain or frequency domain methods or distinguished by their integral equation or differential equation formulations. Important to this application is the ability to faithfully represent constituent lossy dielectric materials of the operator as well as the conductive radiating structure of the handset. As can be noted in the Statement of Work, our intention is to concentrate on the methods with which we have developed some prior experience, such as the Finite Difference Time-Domain Method and the Moment Method.

#### 1.1.1 The Finite-Difference Time-Domain Method

A compact description is presented in the ANTEM '94 paper to be found in Appendix B. More detailed descriptions are provided in its references.

#### **1.1.2 Finite-Element Methods**

See current survey text descriptions.

#### 1.1.3 Moment Methods

Our use of moment methods is best described in the recent work on complex RCS models [1] where comparisons of FDTD results and those with the Numerical Electromagnetics Code are featured.

#### **1.1.4** The Multiple-Multipole Method

See current survey text descriptions.

#### **1.2** Applications to Biological Interactions

The applications to biological interactions are reviewed in the ANTEM '94 paper and its references.

#### 1.2.1 Antenna Performance in the Presence of an Operator

The antenna performance in the presence of an operator is currently best outlined (at cellular phone frequencies) in the work of Anderson et al. shown in Appendix C.

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#### **1.2.2** Exposure to RF Fields

The summary comments of the ANTEM '94 paper apply.

#### 1.2.3 Hyperthermia

The summary comments of the ANTEM '94 paper apply as well as the energy deposition curves shown in the oral paper presentation material of Appendix C.

#### **1.3** Electrical Properties Biological Materials

The table of the ANTEM '94 paper apply as well as the dispersion curves of the oral presentation to be found in Appendix C.

#### **1.4** Measurements and Validation

The work of Anderson et al. described in the oral presentation material of Appendix C is representative of available measurements at cellular phone frequencies.

#### 2.0 THE FINITE-DIFFERENCE TIME-DOMAIN METHOD

The summary of the ANTEM '94 paper applies. Its references provide an outline of the necessary adaptations which are required to provide the representation of the transceiver and its radiating antenna and eventually, the dispersive properties of the dielectric materials.

#### 2.1 The FDTD Algorithm

The summary of the ANTEM '94 paper applies.

#### 2.2 FDTD and Radiating Antennas

The summary of the ANTEM '94 paper applies and especially its reference number 7.

#### 2.3 FDTD with Dispersive Materials

The summary of the ANTEM '94 paper plus its references apply as does the additional material of the oral presentation.

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#### 3.0 PROGRESS TO DATE AND WORK IN PROGRESS

The material below will indicate the progress made thus far in the development of the discretized head model. In addition, several planning discussions have been held with DFL and CRC on the extent and timing of a measurement sequence ranging from handset alone, with phantoms, and eventually with a human operator.

#### **3.1** Modelling of the Cellular Phone

Specifications and form factors of available handsets have been sought. Any initial representations of the external field of the handset alone will likely use NEC wire-grid models which can be used for initial measurement checkout and then for comparisons with subsequent FDTD modelling.

#### 3.2 Code Acquisition and Modification

Recent technical conferences have been used to discuss the availability of FDTD and Moment Method codes which have already been adapted to this problem. No firm commitments have been obtained, since each investigator seems to have the codes in a fluid state.

#### 3.3 Head Phantoms

The material of the oral presentation in Appendix C indicates the type of head phantoms that are available.

The colour illustrations of the head models shown in Appendix C also indicate the model development progress which has been made. It is important to derive these models to launch the computations, either to trade or compare with others and to have a complete and thorough understanding of their composition.

It is also important to remember that eventually discretized models of the phantoms will be required in order to match the corresponding measurement results.

#### 4.0 ACKNOWLEDGEMENTS

The collaboration of Dr. Wilfred Lauber and his patient direction are gratefully acknowledged. Dr. Maria Stuchly was an invaluable source of information and guide on work that is now being done at other laboratories. Discussions with Dr. Shantnu Mishra of DFL have been giving us the encouragement that meaningful measurements can soon be incorporated into this project.

The dedicated and meticulous work of our research assistant, Mina Danesh, gave us an encouraging start with the head model development.

#### 5.0 REFERENCES

 Trueman, C.W. and Kubina, S.J., "HF Ground Wave Radar Studies," Final Report, TN-EMC-94-02, DSS/DREO Contract No. W7714-3-9707/01-SV, EMC Laboratory, Electrical & Computer Engineering, Concordia University, August 1994.

#### 6.0 **APPENDICES**

Appendix A - Statement of Work, Computational Studies for Prediction of Energy Deposition in Humans Exposed to RF Fields from Cellular Phones, Contract Number MIST/CRC 36001-3-3603

Appendix B - Paper Presented at the ANTEM '94 Conference, Ottawa, August 2-5, 1994

Appendix C - Illustrations Used for Oral Presentation of Paper, ANTEM '94 Conference, Ottawa, August 2-5, 1994

## **APPENDIX** A

Statement of Work, Computational Studies for Prediction of Energy Deposition in Humans Exposed to RF Fields from Cellular Phones Contract Number MIST/CRC 36001-3-3603

**EMC Laboratory** 

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6

30 August 1993

#### Statement of Work

#### Computational Studies for Prediction of Energy Deposition in Humans Exposed to RF Fields from Cellular Phones

#### **Aims and Objectives**

The aim of this project is to study the most promising methods of computational modelling which can be applied to predict the energy deposition levels in the human head exposed to the RF fields of modern hand-held cellular telephones. A validated computational methodology would form an essential part of the research aimed at assessing the RF safety of these devices.

This study will concentrate on the frequency band (~900 MHz) presently in use and will serve as a base of evaluation of the methodology for its extension to the higher planned frequencies for personal communications.

#### **Technical Approach**

The exploratory work on this project shall be directed at the survey of the state-of-art of the modelling and measurements in this field, the acquisition of all the physical parameters associated with human body composition and the RF cellular devices, the analysis of the external field distribution (and correlation with known reliable measurement data), in preparation for the volumetric modelling of sources and human body (head, shoulders, arms..) by an adaptation of the FDTD computer programs that are currently in use at the EMC Laboratory.

#### 1. State-of-Art Review

All available literature sources will be examined, with the assistance of DOC, to establish the extent of available modelling information on this problem at frequencies in the cellular radio bands (initially focusing at 900 MHz), corresponding measurement data and the relevant physical parameters of the sources and irradiated subject.

#### 2. Definition of Physical Parameters

From the literature survey, data at the Department of Health and Welfare and any subsequent measurements or estimates, the e.m. properties of the human subject will be catalogued in relation to an appropriate geometric description to be used in the computational modelling at the frequencies specified for this initial study.

Particular note shall be taken of the frequency dispersion of these parameters.

The physical and RF properties of the sources (cellular devices and test dipoles) shall be catalogued for use in the source modelling aspect of the external and internal problem with the most likely computer codes.

#### 3. Analysis of the External Problem

One or two of the available EFIE, or Hybrid integral equation-based codes shall be used to model the external field distribution produced by the sources used in measurements. This is necessary for the realistic definition of source fields to be used with the FDTD method. Comparisons with modelling data of other investigators will be made at this stage.

Close collaboration and contact will be maintained with DOC and the measurement program which is planned to take place as a part of this program.

#### 4. Volumetric Modelling - FDTD

The volumetric modelling of the upper portions of the subject and sources will be carried out in a methodical manner starting with exploratory testing of crude models to simulate published results to the progressive refinement of the discretization of the subject and source to simulate actual measurements that will be undertaken with a phantom and later, human subjects.

Testing will be done with select Gaussian excitations in the available FDTD formulations to concentrate on the applicable frequency bands and also to gain some experience with the frequency dispersion aspects of the modelling problem.

#### **Reporting Requirements**

Brief Summary Quarterly Reports will be prepared to coincide with the invoicing periods as usually required under contract. Yearly Summary Status Reports will be provided and a formal Final Report will be provided.

As work progresses, conference and journal papers will be prepared and cleared with the Technical Authority.

Dr. S. J. Kubina, Eng. 30th August 1993

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## **APPENDIX B**

Paper Presented at the ANTEM '94 Conference Ottawa, Ontario August 2-5, 1994

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#### CELLULAR PHONES AND HUMANS-A NEOPHYTE OUTLOOK

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Abstract-A cellular phone is a portable radio that radiates 0.6 watt from an antenna located very close to the operator's head. Concern has been expressed as to whether a cellular phone is capable of inducing damaging field strengths inside the user's head. This paper reviews the application of electromagnetic modelling techniques using the finite-difference time-domain(FDTD) method to this problem. The construction of an FDTD model of the cellular phone and of the head is discussed. The relative permittivity and conductivity of biological tissue varies with frequency, and this paper reviews modifications of the FDTD method to handle such dispersive media. Some specific applications of measurement methods and of the FDTD method to the calculation of the amount of electromagnetic power delivered to the body are discussed.

#### Introduction

Recently concern has been expressed by the general public and the news media over whether cellular telephones pose a health risk[1]. Cellular phones operate between 824 and 850 MHz, and radiate 0.6 watt of power. Cellular phone sets meet the existing standard for portable radio sets[2], which limit the power that the radio can cause the body to absorb. Tissue damage largely depends on heating due to power delivered by the electromagnetic field[1], given by the "specific absorption rate" (SAR) in watts per kilogram. Standards limit the power absorption to 80 mW/kg average SAR over the entire body, and 1.6 W/kg peak SAR to any 1 gram of tissue for 30 minutes or more. If a device radiates less than 0.74 W at the frequency of cellular phones, it is considered absolutely safe because it cannot deliver the limiting power levels outlined above.

This paper discusses computational modelling of the head in three dimensions, for the purpose of determining the local SAR levels throughout its volume when a cellular phone is in use. The emphasis is on the "finite-difference timedomain" (FDTD) method, although a variety of other techniques are also feasible[3,4]. We will review the basics of FDTD and how an FDTD model of a cellular phone and a head is constructed. Techniques for modelling the dispersive nature of biological tissue will be discussed, and then we will touch on some results that have been published that are pertinent to the assessment of cellular phone hazards.

#### The Finite-Difference Time-Domain Method

In Yee's finite-difference time-domain method[5], space is subdivided into cells of size  $\Delta x$  by  $\Delta y$  by  $\Delta z$  and time into steps of size  $\Delta t$ . The partial derivatives in Maxwell's curl equations are replaced by central-difference formulas involving the components of the electric field along the edges of the cells, and the components of the magnetic field normal to the faces of the cells. Initially all the fields are zero. The time function for the generator voltage is known and is used to obtain a known electric field applied at the base of the antenna. Time is advanced in steps of  $\Delta t$ , and at each time step, the value of the electric field component along each edge of each cell is calculated and then the value of the magnetic field component perpendicular to the center of each cell face is found, using Yee's "leap frog" update scheme. The electric field at time step (n + 1/2) is found from its previous value and the adjacent magnetic field values at step n. Then the magnetic field at step (n + 1/2). In this way the generator voltage is used to determine the electric and magnetic fields throughout the cell space as a function of time.

The FDTD cell space size of  $N_x$  by  $N_y$  by  $N_z$  cells is usually limited by computer resources to  $N_x N_y N_z = 500,000$  or perhaps 1,000,000 cells, although more might be used if a supercomputer is available. The cell space contains the cellular phone and head models at its center, separated from the outer boundary by a layer of free-space cells. Field components within the cell space are computed with the Yee FDTD algorithm. Field components on the outer surfaces of the cell space are updated using an "absorbing boundary condition" (ABC), which is intended to cause the outer surfaces of the cell space to absorb all the energy incident upon them. The most common ABC is the second-order Mur boundary[6]. Because the ABC is imperfect, there can be a significant amount of reflection from the outer boundary. A sufficiently large number of free-space cells must separate the surfaces of the FDTD model from the outer boundary, such that interactions between the model and the outer boundary are reduced to insignificant levels.

The bandwidth of the FDTD cell model extends up to the frequency where the cell size is one-tenth of the wavelength in the dielectric material that has the shortest wavelength. The largest time step  $\Delta t$  that can be used in FDTD is called the "Courant Limit", and for cubical cells is given by  $\Delta t = \Delta x/(\sqrt{3}c)$ , where  $\Delta x$  is the cell size and c is the free-space speed of light. The speed-of-light in the FDTD grid is somewhat dependent on the direction of travel of the wave; the error is larger as the frequency approaches the grid's limiting frequency.

#### Modelling a Person Operating a Cellular Phone

To model a cellular phone operating near a person's head, cells at appropriate locations must be given material properties to represent the cellular phone itself, its antenna, and the head. The cellular phone is modelled by specifying a parallelepiped block of cells to be perfectly-conducting, representing the electrical circuit board and internal components[7]. The plastic case can be modelled with a layer of cells set to a typical relative permittivity for plastic, such as  $\varepsilon_r \approx 2[7]$ . The phone's wire antenna is modelled by setting the electric field to zero along the edges of the cells which lie in a straight line along the path of the wire; currently available techniques require the antenna to be parallel to the x -, or y -, or z -axis. Special care is taken in updating the magnetic field components at the centers of the cell faces adjacent to the antenna[8]. The antenna is excited by specifying the voltage time function applied to the base of the antenna, which is used to specify the electric field on the cell edge modelling the base of the antenna[7] at each time step.

From the electromagnetic point of view, a head is made up of regions of lossy dielectric with various permittivities and conductivities. To build an FDTD model of the head, each cell in the model is assigned electrical parameters appropriate to the tissue at the cell's location: skin, bone, eye, muscle, brain, and so forth. The FDTD update equations for the components of the electric field in each cell use the permittivity and conductivity for that cell. In updating electric field components that lie on the boundaries between cells having differing material types, a weighted average of

the permittivity and conductivity of each cell should be used[9,10]. This is included as a volume-averaged permittivity in Ref. [11], accounting for the fraction of a cell occupied by free-space compared to that composed of, for example, muscle.

#### Antenna Source Voltage Functions

The source voltage is applied as an electric field across the gap at the base of the antenna. The source time function must not contain any frequency components above the bandwidth for the FDTD cell space. If the source voltage time function is to be a sine wave, then it must turn on gradually, as an abrupt start would violate the grid's bandwidth restriction. Then the FDTD algorithm must be run for a sufficient number of time steps that the turn-on transients die away, and sinusoidal steady-state to be reached. For high-permittivity resonant structures, a great many time steps may be required[9], as very long time constants are involved. Under certain circumstances, resonance is encountered in electromagnetic models of human beings[12]. When steady-state is reached each field component in the grid will be sinusoidal, and the amplitude and phase can be determined from the time history for the last period of the input function.

To obtain the frequency response over a wide bandwidth, the FDTD code is often run with a sine wave input many times at individual frequencies. Alternately, we can take advantage of FDTD's solution of Maxwell's Equations in the time domain to obtain information over a wide bandwidth in a single run. The source voltage can be a Gaussian pulse or other time-limited function whose spectral content is restricted to bandwidth for which the FDTD grid is valid. Then the FDTD algorithm finds the time response of each field component in the cell space due to the source voltage pulse. Again, the FDTD algorithm must be run for a sufficient number of time steps to trace the transient response of each field component to zero value. The Fourier transform is used to find the response as a function of frequency. Yee's FDTD assumes that neither the conductivity nor the permittivity vary with frequency, and must be modified to model biological tissues over wide bandwidths, as discussed below.

#### **Computing the Radiated Fields**

The FDTD method obtains the time functions giving the value of the electric fields on each of the cell edges, and the corresponding magnetic fields at the centers of the cell faces, at each time step. These near field time functions are used to find the far fields or radiated fields of the cellular phone operating near the head, with a near-zone to far-zone transformation[13]. Thus at each time step, the far field in any direction can be found from the values of the near fields on a closed integration surface, usually taken to be a few cells inside the absorbing boundary of the cell space.

#### **Electrical Parameters of Biological Materials**

To construct an electromagnetic model of a human being, detailed anatomical information is required. Ref. [14] gives cross-sections of the head and body at approximately 2.5 cm spacing, obtained by dissection. By superimposing the grid of FDTD cells on each cross-section, it is possible to assign a tissue type to each cell. Ref. [15] constructed a cell model of a person in this way, assigning one of 14 different material types to each cell. For computation the cell size was 1.31 cm, with a total of about 145,000 cells in the model. At each frequency the relative permittivity of each of the tissue types are found from the literature[3,16-21] and are listed in Tables 1 and 2 for tissues relevant to modelling the head. Note that the permittivity of some types of tissue is quite high, and that both permittivity and conductivity change very significantly in the frequency range covered by the tables.

| Tissue | Ref             | Free      | quency ( | MHz)                  |      |      |       |     |      |       |      |
|--------|-----------------|-----------|----------|-----------------------|------|------|-------|-----|------|-------|------|
|        |                 | 10        | 27.12    | 100                   | 350  | 400  | 500   | 750 | 900  | 1000  | 1500 |
| Brain, | 3               | 163-352   |          | 57-90                 |      |      |       |     |      | 37-55 |      |
| nerve  | 3               |           | 155      | 52                    | 60   |      |       |     |      |       |      |
|        | 16              |           |          |                       |      |      |       | 49  |      |       | 46   |
| •      | . 17            |           |          | 63                    |      | 50.3 |       |     | 41.2 |       |      |
| . •    | 20              |           |          |                       | 74   |      |       |     |      |       | •    |
| Bone,  | 3               | 37        |          | _23                   |      |      |       |     |      |       |      |
| Fat    | 3               |           | 29       | 7.5                   | 5.7  |      |       |     |      |       |      |
|        | 16              |           | •        |                       |      |      |       | 5.6 |      |       | 5.6  |
| , .    | 17              |           | •        | 12.2                  |      | 9.2  | ,     |     | 7.3  |       |      |
|        | 20              | 1.00.00.4 |          | <i>C</i> <b>A D D</b> | 5.7  |      |       |     |      | 57 50 |      |
| Muscle | 3               | 162-204   | 100      | 04-90                 | 50   |      |       |     |      | 57-59 |      |
|        | 3               |           | 100      | /4                    | 23   |      |       | 52  |      |       | 40   |
|        | 10              |           |          | 70 5                  |      | 67 5 |       | 52  | 517  |       | 47   |
|        | 10              |           |          | 71 76                 |      | 02.5 | 52 51 | •   | 54.7 | 10-52 |      |
|        | 10              |           |          | 63                    |      | 526  | 52-54 |     | 52   | 49-34 |      |
|        | 19              |           |          | 05                    |      | 52.0 | 52.4  |     | 50 5 |       |      |
|        | $\frac{21}{20}$ |           |          |                       | 54   |      |       |     | 50.5 |       |      |
| Slain  | 20              |           | 106      | 25                    | 176  |      |       |     |      |       |      |
| SKIII  | 16              | · ·       |          | 20                    | 17.0 |      |       | 52  |      |       | 49   |
| `      | 18              |           |          | 57                    |      |      | 46 5  | 54  |      | 43-46 |      |
|        | 20              |           |          | . 51                  |      | 17.4 | 10.0  |     |      |       |      |
| Eve    | 3               |           | 155      | 85                    | 80   |      |       |     |      |       |      |
| 2)0    | 16              |           |          |                       |      |      |       | 80  |      | •     | 80   |
| Blood  | Ĩ               |           | 102      | 74                    | 65   |      |       |     |      |       |      |
|        | 18              |           |          | 69-81                 |      |      | 67-70 |     |      | 60.5  |      |
|        | 20              |           |          | . –                   | 65   |      |       |     |      |       |      |

 Table 1

 The permittivity of biological materials at various frequencies.

#### Modelling Dispersive Materials with FDTD

Tables 1 and 2 illustrate that tissue is quite "dispersive", that is, the permittivity and conductivity of tissue varies quite rapidly with frequency. But in Yee's FDTD the relative permittivity  $\varepsilon$ , and conductivity  $\sigma$  cannot change with frequency. To obtain the fields inside a model of a head or body over a wide frequency band, the FDTD code must be run with a sine wave excitation at many individual frequencies, with the appropriate permittivity and conductivity at each frequency. But we lose the inherent advantage of a time domain method, in which the response to a pulse input function obtains wideband information in a single run. To regain this capability, the FDTD algorithm must be modified to directly incorporate the dispersive nature of biological materials.

| Tissue | Re | f F       | requen | cy (MHz)  |       |      |      |      |      |           |      |
|--------|----|-----------|--------|-----------|-------|------|------|------|------|-----------|------|
|        |    | 10        | 27.12  | 100       | 350   | 400  | 500  | 750  | 900  | 1000      | 1500 |
| Brain. | 3  | 0.21-0.63 |        | 0.48-0.95 |       |      | •    |      |      | 0.81-1.2  |      |
| nerve  | 3  |           | 0.45   | 0.53      | 0.65  |      |      |      |      |           |      |
|        | 16 |           | •      |           |       |      |      | 1.2  |      |           | 1.4  |
|        | 17 |           |        | 4.7       |       | 7.5  |      |      | 12.2 |           |      |
|        | 20 |           |        |           | 0.62  |      |      |      |      |           |      |
| Bone.  | 3. | 0.024     |        | 0.057     |       |      |      |      |      |           |      |
| Fat    | 3  | · ·       | 0.04   | 0.07      | 0.072 |      |      |      |      |           |      |
|        | 16 | 1         |        | • .       |       |      |      | 0.09 |      |           | 0.12 |
|        | 17 |           |        | 0.215     |       | 0.88 |      |      | 1.4  |           |      |
|        | 20 |           |        |           | 0.07  |      |      |      |      |           |      |
| Muscle | 3  | 0.69-0.96 | •      | 0.75-1.05 |       | •    |      |      |      | 1.38-1.45 |      |
|        | 3  |           | 0.74   | 1.0       | 1.33  |      |      |      |      |           |      |
|        | 16 |           |        |           |       |      |      | 1.54 |      |           | 1.77 |
|        | 17 |           |        | 6.8       |       | 9    |      |      | 13.8 |           |      |
|        | 19 |           |        | 0.62      |       | 0.68 | 0.72 |      | 0.92 |           |      |
|        | 21 | х<br>э    | •      |           |       |      |      |      | 1.2  |           |      |
|        | 20 |           |        |           | 1.3   |      |      |      |      |           | •    |
| Skin   | 3  |           | 0.74   | 0.55      | 0.44  |      |      |      |      |           | -    |
|        | 16 |           |        |           |       |      |      | 1.54 |      |           | 1.77 |
|        | 17 |           |        |           | 0.42  |      |      |      |      |           |      |
| Eve    | 3  | ,         | 0.45   | 1.9       | 1.9   |      |      |      |      |           |      |
|        | 16 |           |        |           |       |      |      | 1.9  |      |           | 1.9  |
| Blood  | 3  |           | 0.28   | 1.1       | 1.2   |      |      |      |      |           |      |
|        | 20 |           |        |           | 1.22  |      |      |      |      | •         |      |

 Table 2

 The conductivity of biological materials at various frequencies.

The simplest model of materials with frequency-dependent permittivity and conductivity is given by the Debye equation[3]. The frequency dependence of the complex relative permittivity is described with a single relaxation time, according to

$$\varepsilon_r(\omega) = \varepsilon_{r \cdot \cdot \cdot} + \frac{\varepsilon_{rs} - \varepsilon_{r \cdot \cdot}}{1 + j \omega \tau}$$

where  $\omega$  is the operating frequency in rad/sec,  $\varepsilon_{r_{er}}$  is the relative permittivity at high frequencies,  $\varepsilon_{r_{er}}$  is the "static" relative permittivity at zero frequency, and  $\tau$  is the relaxation time. Ref. [22] describes a frequency-dependent finite-difference time-domain or (FD)<sup>2</sup>TD method. The relation between the electric flux density, the frequency-dependent permittivity and the electric field in the frequency domain

$$D(\omega) = \varepsilon_r(\omega)\varepsilon_0 E(\omega)$$

becomes a convolution in the time domain. Ref. [22] modifies the Yee FDTD algorithm to compute the convolution as a running sum, which must be maintained for each location in the cell space. Water, which has a slowly-varying permittivity in the range 0 to 80 GHz, is modelled with  $\varepsilon_{rs} = 81$ ,  $\varepsilon_{rs} = 1.8$ , and relaxation time  $\tau = 9.4$  picoseconds with good agreement with the exact solution in a one-dimensional geometry.

Ref. [23] applies the (FD)<sup>2</sup>TD method to model muscle from 40 to 433 MHz. The relative permittivity of muscle varies with frequency much more rapidly than that of water. At 40.68 MHz, it is 97.3, declining to 53 at 433 MHz, while the conductivity rises from 0.693 S/m at 40.68 MHz to 1.43 at 433 MHz[24]. Ref. [23] choses  $\varepsilon_{rs} = 15$ ,  $\varepsilon_{rs} = 120$ ,  $\sigma = 0.64$  S/m and relaxation time  $\tau = 6.67$  nanoseconds to obtain a reasonable representation of the frequency dependence of the relative permittivity and conductivity of muscle from 40 to 400 MHz. The paper models a two-dimensional cylindrical structure with (FD)<sup>2</sup>TD and compares the result to the exact solution in the form of a Bessel function series evaluated at individual frequencies, with excellent agreement.

Ref. [25] extends the  $(FD)^2TD$  method to handle multiple second-order Lorentz poles. The permittivity is represented as

$$\varepsilon_r = \varepsilon_{r\infty} + (\varepsilon_{rs} - \varepsilon_{r\infty}) \sum_{p=1}^{p} \frac{G_p \omega_p^2}{\omega_p^2 + 2j \omega \delta_p - \omega^2}$$

where  $\sum_{p=1}^{r} G_p = 1$ , P is the number of poles, and  $\omega_p$  and  $\delta_p$  are the resonant frequency

and damping coefficient, respectively, of the p-th pole. The method is an extension of the convolution technique used in Ref. [22] for a simple Debye pole. Ref. [26] extends the method to include multiple simple poles of the Debye type as well as complex poles.

Ref. [11] offers an alternate formulation for  $(FD)^2TD$  based on the solution of a differential equation relationship between the electric flux density D and the electric field E at each time step. The method is applied to model material properties in terms of two simple Debye poles, that is, using two relaxation times. Ref. [27] gives the permittivity of muscle in terms of five relaxation times, valid over a wide frequency range. In Ref. [26] the representation is simplified to use only two time constants, to obtain a reasonable representation of frequency variation of muscle's permittivity from 20 MHz to 20 GHz. The  $(FD)^2TD$  method is applied to the parts of a whole-body model composed of muscle to calculate induced currents in the body at 40, 150 and 350 MHz.

Ref. [28] provides estimates of the parameters required to represent 12 types of biological tissue with the two relaxation time model, valid up to 3 GHz. Each tissue has its individual values of the two "static" permittivities, and the "high frequency" permittivity required for the dispersion model. But all the tissue types are modelled as having the same relaxation times  $\tau_1$  and  $\tau_2$ . This permits the actual permittivity of each cell of the FDTD model to be chosen by volume averaging according to the fraction of the cell occupied by each type of tissue. This improves the resolution of models having a somewhat coarse cell size. Then a single run of the (FD)<sup>2</sup>TD code obtains the field strengths in each cell up to 915 MHz, which would require many runs of the Yee FDTD code with a single-frequency sine wave excitation.

#### SAR Studies using the FDTD Method

It is thought that the principal cause for concern for exposure to electromagnetic fields such as those radiated by cellular phones is due to the heating of tissue. The "specific absorption rate" (SAR) gives the rate at which energy is supplied to tissue, or the power supplied per unit mass, and is given for one cell of an FDTD model by

$$SAR = \frac{\sigma}{2\rho}(E_x^2 + E_y^2 + E_z^2)$$

where  $\sigma$  is the conductivity,  $E_x$ ,  $E_y$ , and  $E_z$  are amplitudes of the x, y, and z components of the sinusoidally-varying electric fields, and  $\rho$  is the mass density of the tissue in the cell. Most studies seek to evaluate the SAR for comparison with safety standards.

Before computational methods that are able to handle dielectric materials became available, measurements were the only means to determine the radiation patterns of portable radios operated near a human head or body. Indeed measurements will continue to provide a quick and accurate means of determining whether cellular phone antenna provides adequate coverage. A typical measurement uses a "head phantom" filled with a material designed to simulate the average electrical properties of biological tissue[29]. Also, embedded field strength probes or temperature probes in a head phantom may provide a useful means of confirming the accuracy of field strength or SAR values obtained by computation. Ref. [30] used a whole-body phantom to measure the power absorbed by a person near a cellular phone antenna on a car. The car phone system is designed to radiate between 3 and 10 watts. The study showed that, for a child standing 15 cm from the antenna, which is very close, the antenna would have to radiate 35 watts to exceed the safety standard.

Ref. [29] considered the energy deposited in the head by a portable radio at 840 MHz, with a whip antenna and a sleeve dipole. The radio differs from a cellular phone in that it is normally held in front of the face with the antenna extending vertically near the operator's eyes and forehead. In Ref. [20] the temperature rise in a head phantom was measured. A "hot spot" was found near the antenna, but the associated temperature rise was so small that there is no potential for tissue damage.

Ref. [20] studied the energy absorbed by a human in the field of a distant antenna, represented by an incident plane wave. Ref. [20] used an FDTD model of the torso having 1.27 cm cells, derived from the cross-sectional anatomy of Ref. [14]. The material properties of each cell were assigned depending on the volume fraction of the cell occupied by each tissue type. The torso model had 16628 cells, and was centered in a space of 36 by 24 by 44 cells for a total of 38016 cells. A sinusoidal excitation was used at both 100 MHz and 433 MHz, with time stepping for three cycles to reach steady state. The SAR was determined for cross-sections of the torso. The study was extended in Ref. [15]. The torso model was extended to become a complete model of a "standard man" 175 cm tall, weighing 70 kg. At 100 MHz, 2.62 cm cells were used with an FDTD space size of 23x12x68 or 18768 cells. At 350 MHz, 1.31 cm cells were used for better resolution, and the overall space used 45x24x135 or 145800 cells. As before, a sinusoidal source was used, and timestepping was carried out for three cycles to reach "steady state". It was found that at 100 MHz there is a considerable difference between the SAR in the body isolated in space compared to the body standing on a ground plane, but at 350 MHz, there is not much difference.

Ref. [21] studied the changes in the performance of a handheld telephone set at 900 MHz and 1900 MHz due to the presence of the operator's hand and head. The telephone set was modelled as a perfectly-conducting box with a quarter-wave monopole on top. The hand and the head were represented as pure muscle tissue. The head was modelled as a sphere of radius 9 cm; the hand was a block 10 cm wide and 2 cm thick wrapped around the lower part of the telephone set. The excitation was a Gaussian pulse modulated with a cosine wave, permitting the impedance and other parameters to be found over a small bandwidth around the cosine's frequency. The total number of cells in the model was 141,680; time stepping was carried out for 1500 steps. The input impedance of the antenna and the azimuth radiation pattern of the FDTD model were compared with the measured impedance and pattern of a real person holding the telephone set. The changes in the impedance and azimuth pattern of the telephone set due to the operator that were expected from the FDTD model were found in the measurement. However, the model was too simple to predict these quantities precisely.

FDTD is often used to study the problem of hyperthermia, in which an applicator is applied to the body to deliver electromagnetic energy to a tumor, which is then destroyed by electromagnetic heating. Ref. [31] studied the use of dipoles to apply energy to tumors at 70 and 95 MHz. In this application, CAT scans taken of an individual patient's body were used to construct an FDTD model with 34751 cells. The FDTD model predicts the internal SAR values when energy is applied with a standard array of dipoles, excited with a sinusoid at a single frequency. The objective is to design the excitation of the array to achieve high SAR in the tumor, but low absorption rates elsewhere to prevent damage to healthy tissue.

Ref. [28] used (FD)<sup>2</sup>TD to study the SARs in the whole body when exposed to a plane wave. As discussed above, the electrical parameters of each tissue type were described with a dispersion model using two relaxation times, valid to 3 GHz. The method was used to find the SAR averaged over layers of the body at frequencies from 20 to 915 MHz, due to plane wave exposure. (FD)<sup>2</sup>TD allows the computation of the fields in the body as a function of time due to a narrow pulse, which contains energy over a wide range of frequencies. Ref. [28] graphs the net current induced at various cross-sections of the body as a function of time, including the dispersive effects of the tissues.

#### Conclusion

Because of the concerns of the general public that cellular phone operation might have an associated health risk, there is tremendous interest and activity in predicting field strengths and specific absorption rates inside the head. But at present there is little published work dealing specifically with the SAR in the head at the frequency of cellular telephones. Existing studies of hazards associated with radiofrequency fields were used to develop the standard maximum field strengths and power levels[2] such that, to the best of available knowledge, cellular telephones pose no health risk. As discussed in this paper, the analytic and numerical tools for detailed accurate computation of field strength levels inside the head have greatly improved in the past five years. We can expect a landslide of information as these methods are applied to assess the SAR levels to be expected in the head when using a cellular phone.

#### Acknowledgement

Dr. M.A. Stuchly's assistance in supplying reference material is gratefully acknowledged.

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**Concordia University** 

### **APPENDIX C**

Illustrations Used for Oral Presentation of Paper Presented at ANTEM '94 Conference Ottawa, Ontario August 2-5, 1994

• • •

EMC Laboratory

Laboratory

CELLULAR PHONES AND HUMANS - A NEOPHYTE OUTLOOK

# S.J. Kubina\*, C. W. Trueman\*, W. Lauber\*\* and S. R. Mishra

\*Concordia University \*\* CRC Ottawa DFL/SPA



Antem 94

# Purpose of Paper

Our Goal: Internal/External Flds-Modelling/Measure ments

- Bio-Materials
- Models and Phantoms
- Modelling

- Measurements
- Target Methods: Computational/Meas urement

Antem 94

# CONSIDERATIONS

Antem 94

## Physical

- Set Form Factors
- Humans m/f/c
- Electromagnetic
  - Frequency
  - Antenna/Feed
  - Polarization
  - Power
  - .....next......



36 C-5

# l technology

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Components for cellular handsets, PCN and private radio networks.

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# CONSIDERATIONS (Cont'd)

- Electromagnetic ... - Radiation Pattern
  - Volumetric
  - Near Field Structure
    - External
  - Internal Fields
    - Deposition Rates



# ANATOMICAL MODELS & PHANTOMS

# Primary Source

- Eycleshymer & Schoemaker 1911
- Utah Prof. Om Ghandi et al.
- University of Ottawa
- Commercial
   Other...



Antem 94





FREQUENCY IN MHz

Figure 8-3 Whole-body-averaged specific absorption rate of prolate spheroidal models of average man, rhesus monkey, medium rat, and medium mouse for E polarization at incident plane-wave power density of 1 mW/cm<sup>2</sup>. (From Durney, C. H., Johnson, C. C., Barber, P. W., et al. [1978]. Radiofrequency radiation dosimetry handbook, 2nd ed. Pub. No. SAM-TR-78-22, USAF School of Aerospace Medical Division (AFSC), Brooks Air Force Base, Tex.)

from Gandhi, Om P., Ed., "Biological Effects and Medical Applications of Electromagnetic Energy," Prentice Hall, 1990, ISBN 0-13-082728-2.

Sulcus hypothalamicus [Monroi] Corpus fornicis Massa intermedia Foramen interventriculare [Monroi], Lobulus paracentralis Corpus mamillare, Os parietale Sulcus corporis callosi. Thalamus Tela chorioidea ventriculi tertii · Os frontale Ventriculus tertius Sulcus cinguli Praecuneus Commissura posterior [cerebri] Gyrus frontalis superior Recessus posterior fossae interpedun: Genu corporis callos Corpus pineale Commissura anterior [cerebri] Corpora quadrigemina Aquaeductus cerebri [Sylvii] Lamina terminalis Velum medullare anterius Os nasale obulus centralis Recessus opticus, Cuneus Chiasma opticum... Monticulus Concha nasalis superior ... minae medullares Cartilago septi nasi et septum nasi. īī Concha nasalis media occipitale Hypophysis. Gyrus lingualis Concha nasalis inferior Maxilla-Tuber [vermis] >... Pyramis [vermis] Palatum durumpus medullare M. orbicularis orisvula [vermis] Lingu Ostium pharyngeum tubae auditivae et os occipitale Tela chorioidea ventriculi tertir Pons [Varoli] Mandibula M. transversus meinti Foramen caecum N. geniogloscus / M. geniohyoideus M. mylohyoideus M. semispinalis cervicis Usulaz Os hyoideum/ Lig. nuchae Pharynx [pars oralis] / Cartilago epiglottica et epiglottis/ Medulla spinalis Vestibulum laryngis/ Pharynx [pars laryngea]/ Mm. constrictores pharyngis

C-9

#### KEY-FIGURE II

9(b) C-10 Cartilago septi nasi se Cavum nasi Naris et vomer se M. nasalis [pars transversa] M. nasalis 27 s Pharynx [pars nasalis] Concha nasalis inferior of M. caninus M. quadratus labii superioris aș Meatus nasi interior # Sinus maxillaris [Highmori] N. infraorbitalis of Ostium pharyngeum tubae auditivae Torus tubarius # N. infraorbitalis et v. angularis V. angularis et sinus maxillaris [Highmori] 🕫 M. quadratus labii superioris A. angularis 91 no Maxilla A. palatina descendens #g 11 M. pterygoidaus internus M. constrictor pharyngis superior as, ts M. orbicularis ocull A. maxillaris interna 😝 D Tuba auditiva (Eustachii) Lig. longitudinale anterius #2 A Os zygomaticum M. tensor veli palatini & #Recessus pharyngeus [Rosenmuelleri] et m. sygomat: M. temporalis as ris interna et mandibula Mandibula [processus coronoideus] 84 [processus coronoideus] M. pterygoidous internus as, r veli palatini et v. pharyngea N. mandibularis 82 massatar M. longus capitis 81 longus capitis et m. rectus capitis anterior M. pterygoideus externus & adibularis M, pterygoideus externus M. levator veli palatini 78 es M. Jevator vali palatini N. messeter 75\_ A meningea media et glandula perotis Glandula parotis 77\_ N. auriculotemporalis et v. facialiis posterio Lig. temporomandibulare 78\_ Lig. sphenomandibulare et a. temporalis A. meningea media et lig. sphenomandibulare 75... st A. carotis interna superficialis es Truncus sympathicus et glandula parotis facialis posterior et a. temporalis superficialis 74 ... N. auriculotemporalis et a. carotis interna 73 ... e Cartilago auriculae Cartilago auriculae et glandula parotis 73\_... N. accessorius et n. glossopharyngeus 71.--V. jugularis externa et emissarium condyloideum 70.--s: Ganglion nodosum et n. accessorius JI N. facialis 🗢 V. jugularis interna N. facialis #. St Cellulae mastoideae Ganglion nodosum er Emissarium condyloideum Cellulae mastoideae #7-M. rectus capitis lateralia M. sternocleidomastoideus ar N. hypoglosous N. hypoglossus es. Sinus transversus Sinus transversus #4 Sutura occipitomastoidea Os temporale #3-M. obliquus capitis superior ## A. et v.occipitalis Tonsilla cerebelli #1/ 41 M. splenius capitis A. vertebralis 60/ 44 A.vertebralis M. splenius capitis se." 43 N. hypoglossus et n. accessorius 44 Os occipitale [pars basilaris] N. hypoglossus 58/ 45 A. et v. occipitalis [ramus] Lobulus biventer si Pyramis medullae oblongatae se 44 N. occipitalis major 47 Oliva et nucleus funiculi gracilis A. et v. occipitalis [ramus] 55 48 M. trapezius N. occipitalis major su 49 Decussatio lemniscorum Lobulus semilunaris inferior 33 M. semispinalis capitis se 50 Os occipitale et lig. nuchae Nucleus funiculi gracilis 31

#### SECTION II





from Gandhi, Om P., Ed., "Biological Effects and Medical Applications of Electromagnetic Energy," Prentice Hall, 1990, ISBN 0-13-082728-2.





from Gandhi, Om P., Ed., "Biological Effects and Medical Applications of Electromagnetic Energy," Prentice Hall, 1990, ISBN 0-13-082728-2.

C-12





from Gandhi, Om P., Ed., "Biological Effects and Medical Applications of Electromagnetic Energy," Prentice Hall, 1990, ISBN 0-13-082728-2.

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C-13

| Lu         | ung Material. (C) Castable Bon<br>quid Bone Material Componen   | ne Material Composits. | onents. (D) |
|------------|---|------------------------|-------------|
| <b>.A.</b> | Muscle and brain material   | Percentag              | e by weight |
| Ma         | aterial   | Muscle                 | Brain       |
|            | Water   | 52.4                   | 40.4        |
|            | Salt (NaCl)   | 1.4                    | 2.5         |
|            | Sugar   | 45.0                   | 56.0        |
|            | HEC   | 1.0                    | 1.0         |
|            | Bacteriacide  | 0.1                    | 0.1         |
| В.         | Lung material   |                        |             |
| Ma         | Salt (NaCl)<br>Sugar<br>HEC<br>Bacteriacide<br>B. Lung material<br>Material<br>Muscle material (above)<br>Microspheres<br>C. Bone material (castable)<br>Material<br>Two ton epoxy<br>Epoxy<br>Hardener<br>KCl Solution | Percentag              | e by volume |
|            | Muscle material (above)<br>Microspheres   |                        | 47<br>53    |
| <b>C</b> . | Bone material (castable)  |                        |             |
| Ma         | terial  | Percentag              | e by weight |
| •          | Two ton epoxy   |                        |             |
|            | Ероху   | . 3                    | 5.0         |
|            | Hardener  | . 3                    | 5.0         |
|            | KCl Solution  | 2                      | 8.0         |
| D.         | Bone material (liquid)  | -                      |             |
| Ma         | terial  | Percentag              | e by weight |
|            | TWEEN   | 5                      | 7.0         |
|            | n-Amyl alcohol  | 2                      | 8.5         |
|            | Paraffin oil  |                        | 9.5         |
|            | Water   |                        | 4.5 ·       |
|            | Salt (NaCl)   |                        | 05          |

TABLE 1. (A) Composition by Weight of Muscle and Brain Equivalent Material. (B) Percentage by Volume of Filler Used in

from Hartsgrove, G., Kraszewski, A., and Surowiec, A., "Simulated Biological Materials for Electromagnetic Radiation Absorption Studies," Bioelectromagnetics 8:29-36 (1987), © Alan R. Liss, Inc.

C-14

5.4

 TABLE 2. Dielectric Constant and Conductivity of Tissue Equivalent Materials at Selected

 Frequencies

|             | Frequency (MHz) |      |            |      |             |      |  |  |  |  |  |
|-------------|-----------------|------|------------|------|-------------|------|--|--|--|--|--|
|             | 1               | 00   | . 4        | 00   | 900         |      |  |  |  |  |  |
| Material    | e'              | σ    | £'         | σ    | ε'          | e    |  |  |  |  |  |
| Muscle      | 70.5            | 6.8  | 62.5       | 9.0  | 54.7        | 13.8 |  |  |  |  |  |
| Brain       | 63.0            | 4.7  | 50.3       | 7.5  | 41.2        | 12.2 |  |  |  |  |  |
| Lung        | 37.0            | 3.4  | 32.6       | 4.3  | <b>28.0</b> | -6.6 |  |  |  |  |  |
| Bone cast   | 13.6            | 0.08 | 9.3        | 1.1  | 7.4         | 1.6  |  |  |  |  |  |
| Bone liquid | 10.8            | 0.35 | <b>9.1</b> | 0.66 | 7.2         | 1.2  |  |  |  |  |  |

from Hartsgrove, G., Kraszewski, A., and Surowiec, A., "Simulated Biological Materials for Electromagnetic Radiation Absorption Studies," Bioelectromagnetics 8:29-36 (1987), © Alan R. Liss, Inc.

C-15

PROPERTIES OF BIOMATERIALS

 Categorization
 Typical Values
 Dispersion Modelling - Hurt
 Recent Results - Furse, Chen , Ghandi, May 94



| TABLE 5-3 Dielectric Constant at 37° C of Various Tissues at Radio and Microwave Frequencies |   |  |   |  |   |                                  |  |  |  |  |
|--|---|--|---|--|---|----------------------------------|--|--|--|--|
|  | Frequency   |  |   |  |   |                                  |  |  |  |  |
| Tissue   | 100 kHz   | 1 MHz  | 10 MHz  | 100 MHz  | 1 GHz                                     | 10 GHz                           |  |  |  |  |
| Skeletal muscle<br>Liver<br>Spleen<br>Kidney<br>Brain<br>Bone                                | $\begin{array}{c} (14.4-24.8)\times10^3\\ (9.8-13.7)\times10^3\\ 3.3\times10^3\\ (10.9-12.5)\times10^3\\ (1.96-3.8)\times10^3\\ 280\end{array}$ | $\begin{array}{c} (1.9-2.5)\times10^{3}\\ 1.97\times10^{3}\\ 1.45\times10^{3}\\ (2.39-2.69)\times10^{3}\\ (0.54-1.25)\times10^{3}\\ 87\end{array}$ | 162–204<br>251–338<br>321–410<br>190–204<br>163–352<br>37 | 64–90<br>65–82<br>69–101<br>66–95<br>57–90<br>23 | 57–59<br>47–49<br>50–55<br>42–50<br>37–55 | 43-45<br>35<br>41<br>40<br>38-44 |  |  |  |  |

| TABLE 5-4 | Conductivity at 37° C of Various Tissues at Radio and Microwave |
|-----------|---|
| · · · ·   | Frequencies in Siemens per Meter                                |

| •               | Frequency |           |                |           |             |        |  |  |  |  |
|-----------------|-----------|-----------|----------------|-----------|-------------|--------|--|--|--|--|
| Tissue          | 100 kHz   | 1 MHz     | : 10 MHz 100 ] |           | 0 MHz 1 GHz |        |  |  |  |  |
| Skeletal muscle | 0.38-0.59 | 0.58-0.85 | 0.69-0.96      | 0.75-1.05 | 1.38-1.45   | 11.5   |  |  |  |  |
| Liver           | 0.15-0.16 | 0.27-0.3  | 0.42-0.47      | 0.6-0.72  | 0.95-1.1    | 8.9    |  |  |  |  |
| Spleen          | 0.62      | 0.63      | 0.5-0.84       | 0.73-1.05 | 1.09-11.3   | 10.1   |  |  |  |  |
| Kidney          | 0.24-0.25 | 0.360.37  | 0.500.68       | 0.66-1.05 | 0.95-1.0    | 9.7    |  |  |  |  |
| Brain           | 0.12-0.17 | 0.140.21  | 0.21-0.63      | 0.48-0.95 | 0.81-1.2    | 8-10.8 |  |  |  |  |
| Bone            | 0.014     | 0.017     | 0.024          | 0.057     |             |        |  |  |  |  |

from Gandhi, Om P., Ed., "Biological Effects and Medical Applications of Electromagnetic Energy," Prentice Hall, 1990, ISBN 0-13-082728-2.

C-18

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## The Permittivity of Biological Materials at Various Frequencies

|    | Tissue   | Freque  | ency (M       | Hz)   |      |      |       |      | •    |       |      |
|----|----------|---------|---------------|-------|------|------|-------|------|------|-------|------|
|    | Drain    | 10      | 27.12         | 100   | 350  | 400  | 500   | 750  | 900  | 1000  | 1500 |
|    | nerve    | 105-552 | 155           | 52    | 60   |      |       |      |      | 51-55 |      |
|    |          | •       | • •           | 62    |      | 50.2 |       | 49   | 41.0 |       | 46   |
|    | •        | • •     | <del></del> . | . 03  | 74   | 50.5 |       |      | 41.2 |       |      |
|    | Bone,    | 37      | • •           | 23    |      |      |       |      | •    | • .   |      |
|    | Fat      |         | 29            | 7.5   | 5.7  |      |       | 56   |      |       | 56   |
|    |          |         | •             | 12.2  |      | 9.2  |       | 2.0  | 7.3  |       | 5.0  |
|    | Musele   | 162-204 |               | 64-00 | 5.7  |      |       |      |      | 57.50 |      |
|    | 141USCIC | 102-20+ | 106           | 74    | 53   |      |       |      |      | 57-55 |      |
|    |          |         | •             | 70.5  |      | 675  | ~     | 52   | 517  |       | 49   |
|    | x        |         |               | 71-76 |      | 02.5 | 52-54 | ·    | 54.7 | 49-52 |      |
|    |          |         |               | 63    |      | 52.6 | 52.4  |      | 52   |       |      |
|    | •<br>•   |         |               |       | 54   |      |       | ٠    | 20.2 |       |      |
|    | Skin     |         | 106           | 25    | 17.6 |      |       |      |      |       |      |
|    |          |         |               | 57    |      |      | 46.5  | 52   |      | 43-46 | 49   |
|    |          |         | •             |       |      | 17.4 | -10.2 |      |      | .5 .0 |      |
| •. | Eye      |         | 155           | 85    | 80   |      |       | . 80 |      |       | 80   |
|    | Blood    |         | 102           | 74    | 65   |      |       | 00   |      |       | 00   |
|    |          | •••     |               | 69-81 | 65   |      | 67-70 |      |      | 60.5  |      |
|    |          |         |               |       | 05   |      |       |      |      |       |      |

## The Conductivity of Biological Materials at Various Frequencies

| Tissue | Frec      | uency | (MHz)     |       |      |      |       |      |           |      |
|--------|-----------|-------|-----------|-------|------|------|-------|------|-----------|------|
| Drain  | 10        | 27.12 | 100       | 350   | 400  | 500  | 750   | 900  | 1000      | 1500 |
| nerve  | 0.21-0.05 | 0.45  | 0.48-0.93 | 0.65  |      |      |       |      | 0.01-1.2  |      |
|        |           |       | • • •     |       |      |      | 1.2   |      | •         | 1.4  |
|        |           |       | 4.7       | 0.00  | 7.5  |      |       | 12.2 | •         |      |
| Rone   | 0.024     | · •   | 0.057     | 0.62  |      |      |       |      |           |      |
| Fat    | 0.024     | 0.04  | 0.07      | 0.072 |      |      |       |      |           |      |
|        |           |       |           | 0.072 |      |      | 0.09  |      |           | 0.12 |
|        | ·         |       | 0.215     |       | 0.88 |      |       | 1.4  |           |      |
| 76 1   | 0.00.000  |       | 0.75 1.05 | 0.07  |      |      |       |      | 1 00 1 45 | • •  |
| Muscle | 0.69-0.96 | Ö 74  | 0.75-1.05 | 1 22  |      |      |       |      | 1.38-1.45 |      |
|        |           | 0.74  | 1.0       | 1.55  |      |      | 1 54  |      |           | 177  |
|        |           |       | 6.8       |       | 9    |      | 1.01  | 13.8 |           | //   |
|        |           |       | 0.62      |       | 0.68 | 0.72 |       | 0.92 |           |      |
| · · ·  |           |       |           |       |      |      |       | 1.2  |           | •    |
| Slein  |           | 074   | A 55      | 1.3   |      |      |       | -    | ,         |      |
| SKIII  |           | 0.74  | 0.55      | 0.44  |      |      | 1 54  |      |           | 1 77 |
|        |           |       |           | 0.42  |      |      | 1.57  |      |           | 1.// |
| Eye    |           | 0.45  | 1.9       | 1.9   |      |      |       |      |           |      |
|        |           |       |           |       |      |      | - 1.9 |      |           | 1.9  |
| Blood  |           | 0.28  | . 1.1     | 1.2   |      |      |       |      |           |      |
|        | •         |       | •         | 1.22  |      |      |       |      |           |      |

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# DISPERSION

DEBYE EQUATION WITH 2 RELAXATION CONTHINTS

 $e^{\dagger}(\omega) = e_{o}\left[\frac{\varepsilon_{oo} + \frac{\varepsilon_{s_{1}} - \varepsilon_{oo}}{1 + j\omega r_{1}} + \frac{\varepsilon_{s_{2}} - \varepsilon_{oo}}{1 + j\omega r_{2}}\right] - \dots + (+)$ 

## FURSE et al.

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Antem 94

64

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| TABLE 1<br>DEBYE CONSTANTS FOR TISSUES $\tau_1 = 46.25$ ns, $\tau_2 = 0.0907$ ns<br>(Average of optimum for fat and muscle.) |       |       |       |
|--|-------|-------|-------|
|  |       |       |       |
| Muscle   | 40.0  | 3948. | 59.09 |
| Bone / Cartilage   | 3.4 - | 312.8 | 7.11  |
| Blood  | 35.0  | 3563. | 66.43 |
| Intestine  | 39.0  | 4724. | 66.09 |
| Liver  | 36.3  | 2864. | 57.12 |
| Kidney   | 35.0  | 3332. | 67.21 |
| Pancreas / Spleen  | 10.0  | 3793. | 73.91 |
| 1/3 Lung   | 10.0  | 1224. | 13.06 |
| Heart  | 38.5  | 4309. | 54.58 |
| Brain/Nerve  | 32.5  | 2064. | 56.86 |
| Skin   | 23.0  | 3399. | 55.59 |
| Eye  | 40.0  | 2191. | 56.99 |

from Furse, Cynthia M., Chen, Jin-Yuan, and Gandhi, Om P., "The Use of the

Frequency-Dependent Finite-Difference Time-Domain Method for Induced Current and SAR Calculations for a Heterogeneous Model of the Human Body," IEEE Transactions on EMC, Vol. 36, No. 2, May 1994, pp. 128-133.



Fig. 1. Relative dielectric permittivity versus frequency: curve is best fit for five-term Debye expression to data (x's) found in Table I.

from Hurt, William D., "Multiterm Debye Dispersion Relations for Permittivity of Muscle," IEEE Transactions on Biomedical Engineering, Vol. BME-32, No. 1, January 1985, pp. 60-64. 64





from Hurt, William D., "Multiterm Debye Dispersion Relations for Permittivity of Muscle," IEEE Transactions on Biomedical Engineering, Vol. BME-32, No. 1, January 1985, pp. 60-64.

63)

C-23

•MEASUREMENTS & MODELLING Toftgard, Hornsleth, Bach Anderson, June 93 Furse et al..., May 94 DFL Measurements Deslile et al... Other

Antem 94



Fig. 1. The telephone-operator model used for the simulations. Both hand and head are modeled as pure muscle.

from Toftgård, Jørn, Hornsleth, Sten N., and Andersen, Jørgen Bach, "Effects on Portable Antennas of the Presence of a Person," IEEE Transactions on Antennas and Propagation, Vol. 41, No. 6, June 1993, pp. 739-746.



Simulated and measured results for a  $\lambda/4$  monopole mounted on the telephone with the operator included for f=914 MHz (a) Radiation pattern in the XY plane. (b) Radiation pattern in the XZ plane. (c) Input impedance.

from Toftgård, Jørn, Hornsleth, Sten N., and Andersen, Jørgen Bach, "Effects on Portable Antennas of the Presence of a Person," IEEE Transactions on Antennas and Propagation, Vol. 41, No. 6, June 1993, pp. 739-746.



from Toftgård, Jørn, Hornsleth, Sten N., and Andersen, Jørgen Bach, "Effects on Portable Antennas of the Presence of a Person," IEEE Transactions on Antennas and Propagation, Vol. 41, No. 6, June 1993, pp. 739-746.

900

(c) Fig. 2. Simulated and measured results for a  $\lambda/4$  monopole mounted on the telephone without the operator included for f =914 MHz. (a) Radiation pattern in the XY plane. (b) Radiation pattern in the XZ plane. (c) Input impedance.

950

Freq. [MHz]

1000

1050 1100

850

-100 -

7





c) Far field horizontal diagram (maximum value corresponds to 51.2 mV/m at a distance of 1000 m)

## Fig. 6: Electric field of a hand-held transmitter with inclined antenna (27.5°) at 430 MHz

from Brüns, H.-D., Singer, H., and Mader, T., "Field Distributions of a Hand-Held Transmitter Due to the Influence of the Human Body," 10th International Zurich Symposium and Technical Exhibition on EMC, March 9-11, 1993, Session on Biological Effects, pp. (3A3) 9-14.

# FDTD MODEL PARAMETERS

Antem 94

Paper list: Cell Size (~ 3. (mm) White Space ~ 30 - 2<sup>nd</sup> order Source - Lucbbers --Dispersion - Later Output

# NEXT STEPS - CONCLUSION

Antem 94

Search for Models, Codes & Data Codes - External Problem Model Development FDTD Code Refinement Concurrent Measurement **Program/Validation** Contact with Community



C-31









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INDUSTRY CANADA / INDUSTRIE CANADA

