

CARLETON UNIVERSITY

MASTER'S THESIS

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## Simulation of an Organic Memristor

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*"Quote"*

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*Abstract*

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Master's of Science in Electrical and Computer Engineering

**Simulation of an Organic Memristor**

by Cem BONFIL

The device that is simulated in this thesis has a conducting PEDOT:PSS sheet with two metal contacts and an electrolyte solution containing lithium and perchlorate ions is placed on top of the conducting sheet. When a potential is applied at both ends of the conducting polymer, lithium ions, which act as reversible counter doping agents, travel into the PEDOT and modify the conductivity of the sheet. Overall this system mainly consists 4 different charge carriers, 3 (holes, lithium and perchlorate ions) mobile and 1 stationary (electrons).

## *Acknowledgements*

The acknowledgements and the people to thank go here, don't forget to include your project advisor...

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# Abbreviations

BTE	Boltzmann Transport Equations
CFL	Courant-Friedrichs-Lowy
$ClO_4^-$	Perchlorate
DD	Drift Diffusion
FD	Finite Difference
$Li^+$	Lithium
PEDOT	Poly(3,4-ethylenedioxythiophene)
PSS	Poly(styrenesulfonate)
SG	Scharfatter-Gummel

# Physical Constants

Boltzmann Constant     $k = 8.6173324 \cdot 10^{-5}$  eV/K

Elementary Charge     $q = 1.6021765 \cdot 10^{-19}$  C

Vacuum Permittivity     $\epsilon_0 = 8.8541878 \cdot 10^{-12}$  F/m

# Symbols

D	Diffusivity	$m^2/s$
E	Electric Field	$V/m$
h	Unit Distance	$m$
I	Current	$A$
J	Current Density	$A/m^2$
$L_D$	Debye Length	$m$
$l_f$	Mean Free Path	$m$
M	Memristance	$\Omega$
n	Electron Density	$m^{-3}$
$N_A$	Acceptor Density	$m^{-3}$
$N_D$	Donor Density	$m^{-3}$
$n_i$	Intrinsic Carrier Density	$m^{-3}$
T	Temperature	$K$
t	Time	$s$
$t_{dr}$	Dielectric Relaxation Time	$s$
U	Generation Recombination Rate	$m^{-3}s^{-1}$
V	Potential	$V$
$V_{th}$	Thermal Voltage	$V$
$\Delta$	Unit Distance	$m$
$\epsilon_r$	Relative Permittivity	unitless
$\mu$	Mobility	$m^2/(Vs)$
$\varphi$	Flux Linkage	$Wb$
$\rho$	Charge Density	$C/m^3$
v	Drift Speed	$m/s$

*For/Dedicated to/To my...*

# **Chapter 1**

## **Introduction**

### **1.1 Thesis Objectives**

The purpose of this thesis is to create a numerical simulation which captures the physics behind the operation of a memristor. The memristor simulation developed in this thesis can be used for researching the physics behind the actual device through further modeling or it could be used as a tool for designing memristors for a specific use.

### **1.2 Thesis Overview**

Initially, discovery of memristor and challenges in memristor research is introduced in the background chapter. It emphasizes the need for better physical modeling and lack of simulation tools required for further analysis and research.

In chapter 2 , Boltzmann Transport Equations (BTE) are shortly introduced. This introduction is followed by explanation of transport mechanisms captured through BTE. Additionally drift diffusion equations (simplified form of BTE) which will be used for the simulation of the memristor are discussed. Finally few analytic solutions to drift diffusion equations, such as charged particles moving over an infinite conductor and PN junction, are generated.

In chapter 3, a numerical solution to drift diffusion equations is formed to calculate the movement of all the charged particles in a memristor. First finite difference method, which is the basis for all the simulations in this thesis, is introduced. Then it is applied to drift diffusion equations. Additionally, a Poisson solver to be used with drift diffusion equations is developed in order to calculate the electric field generated by charged particles and metal contacts. Also, boundary conditions necessary for the solution of

both drift diffusion and Poisson solver are introduced. All the equations and boundary conditions combined generates a set of equations to be solved for the memristor simulation. Various schemes for solving these differential equations as well as the numerical and physical limitations are discussed in detail.

In chapter 4, the numerical solver developed in chapter 3 is tested against analytic solutions as well as a commercially available simulator called COMSOL Multiphysics. A mechanism to limit the maximum density of any particle, which is essential for memristor simulation, is developed and tested.

After the stability and accuracy of the simulation scheme was tested in chapter 4, a memristor model is introduced in chapter 5. The difficulties in the simulation due to physical and numerical restrictions were discussed. A possible approximation 1-D approximation for a 2-D memristor simulation was proposed and simulated. After demonstrating that the simulations produced reasonable results, the validity of the approximations made for 1-D and 2-D simulations are investigated.

A full simulation 2-D is made in chapter 6. A memristor with various PEDOT thicknesses were simulated. The simulation with thinnest PEDOT is compared to 1-D simulations made in the previous chapter. This chapter ends with comparison of simulation results with experimental data and shows how parameter fitting can be used to match experimental data.

The conclusion chapter summarizes the findings of the thesis, discusses advantages and disadvantages of the proposed simulation methods and provides suggestions for improvements on the model and opportunities further research.

### 1.3 Background

The term memristor was first used by Chua in 1971 in his paper called "Memristor The Missing Circuit Element" [1]. He theorized that there is a fourth passive circuit element yet to be discovered in addition to resistor, capacitor and inductor. He said that we already know and use five out of six possible combinations that can be made out of four fundamental circuit variables, current  $I$ , voltage  $V$ , charge  $q$  and flux linkage  $\varphi$ . Chua claimed that we were missing a circuit element which produces a relationship between flux linkage (time integral of the potential) and charge. He introduced a new variable named memristance which has units of resistance and it is a function of charge. The relationship between the current and the potential of a memristor is calculated by replacing the resistance by memristance in ohms law:

$$v(t) = M(q(t))i(t) \quad (1.1)$$

Theoretically memristor retains its resistance in the absence of any power source. When a potential is applied the resistance can be influenced by the direction and the magnitude of the current. If the current flows in one direction resistance increases and if it flows in the other direction resistance decreases. This produces an I-V response which looks like a pinched hysteresis curve. This response is the main characteristic of a memristor.

This new element remained mostly a theory and did not get much academic attention until a group of researchers in Hewlett Packard developed a fully functioning memristor. They successfully fabricated a nano scale memristor using  $TiO_2$  (titanium dioxide). After the discovery in HP labs there was an increase of interest in different types of memristors due to their potential applications for data storage and addition of learning capabilities into passive circuits.

Recently it has been suggested that organic based memristors could be fabricated (ref). This thesis will be concerned with such devices. They are usually constructed on a larger scale than the ones fabricated in HP labs. The memristor studied in this thesis is in millimeter rather than nano meter scale and uses a conductor made out of a polymer called PEDOT:PSS. It is composed of two polymer chains, PEDOT and PSS attached together and conducts electricity via hole transport. Unfortunately the conduction mechanism is not perfectly understood and needs further research due to the complexity of the material(ref).

Unlike a semiconductors like silicon, the structure of a polymer is quite irregular. Polymers have individual molecules with different chain lengths and a variable amount of defects. Additionally, they can be amorphous or partially crystalline and further differences occur through aging. The conduction and electronic properties depend on the orientation of polymer chains which can change in x,y and z directions. These irregularities in the structure makes these conducting polymers resemble amorphous inorganic semiconductors. Following the concept of charge transport in amorphous inorganic semiconductors, the conduction mechanism of conducting polymers is commonly explained by variable range hopping. This mechanism was first introduced by Mott in 1968. He proposed a model for charge transport in systems that are randomly disordered. In variable range hopping charge transport occurs via jumps between available sites. Every charge carrier has a probability of jumping between two sites depending on its energy and the distance to the next available site. Temperature has a big impact on conductivity since it changes the structure of the molecules and increases the energy of charge carriers.

Apart from temperature there are other ways to change the conductivity of PEDOT. It is possible to affect conductivity by either doping or counter doping. Depending on the method used, doping can be reversible or permanent. When reversible doping is employed, resulting device behaves like a memristor since its resistance is now dependent on charge. All these effects make physical modeling, experimentation and simulation very challenging.

There are few approximate analytic solutions for inorganic memristors similar to the one produced in HP labs. These solutions are mostly based on parameter fitting and some basic physics. Studies on organic memristors are even more sparse than studies on inorganic ones. The physics behind the conduction mechanism and the changes in conductivity due to doping requires further research and development of a computer model can help in various ways. First of all it is very difficult to get experimental data on the movement of ions and holes inside PEDOT due its disordered structure. The thickness of PEDOT:PSS (usually in  $\mu\text{m}$  scale) complicates the tracing of the ion in all dimensions. Theories that explain the movement of charged particles and their interactions with each other are difficult to formulate since it is hard to obtain required experimental data. A simulation can be a useful tool in testing various theories without having to set up and perform complicated experiments. Also once a solid understanding of the conduction mechanism has been achieved, simulations can be very useful to test different device structures and optimize them before fabrication.

(need to talk about lack of simulations for memristor)

# Chapter 2

## Theory

### 2.1 Carrier Transport Equations

Drift diffusion equations, which are based on BTE, need to be solved in order to model the complex behaviour of the memristor. Drift-diffusion equations can be obtained by simplifying BTE equation via approximations. These simplifications dictate the limits of the drift-diffusion model and serve as guidelines for where this model can be used, therefore they need to be well understood.

Derivation of Boltzmann transport equation starts by stating that any charged particles can be defined its position in space  $\mathbf{r}$  and momentum  $\mathbf{k}$  in time  $t$  using a probability distribution function  $f(k,r,t)$ . This results in the most general form of Boltzmann transport equation.

$$\frac{d}{dt}f(k,r,t) = 0 \quad (2.1)$$

This general form of BTE needs to be expanded and relevant physical equations need to be placed in to get an appropriate equation describing a specific problem or a device. Many different device simulators use some sort of approximation to BTE. In a semiconductor simulations they are commonly used for the movement and the density distribution of charge carriers, holes (p) and electrons (n).

After this brief introduction to BTE, rather than going through the tedious mathematical derivation of the drift diffusion model, the approximations that are made along the process of derivation can be discussed in order to get a better insight on the model. As a particle travels in a solid state device it collides with other particles as well as the atoms in the device. For drift-diffusion equations individual lattice scattering events or

collisions are averaged and the particles have an average constant velocity under the effects of an electric field. This means that all the particles respond instantaneously to the changes in the electric field. The movement of the particles due to electric field is called the drift current. The relationship between drift velocity and the electric field is given by the following equation:

$$v = \mu E \quad (2.2)$$

$\mu$  is called mobility constant and it determines the speed at which the particles are going to move when subject to an electric field. Drift current density can be derived based on drift velocity.

$$J_E = qn\mu E = qnv \quad (2.3)$$

Where  $q$  is the elementary charge and  $n$  is the electron density. In addition to the previous assumptions, it is assumed that the lattice is perfectly uniform, has a uniform temperature distribution and all the particles are close to the temperature of the lattice. Based on this assumption we can say that all the particles that move due to thermal effects has the same thermal velocity ( $v_{th}$ ) and mean free path ( $l_f$ ). These quantities can be combined into one single coefficient called diffusion constant.

$$D = v_{th} l_f \quad (2.4)$$

Drift and diffusion coefficients are related to each other via Einstein relationship:

$$D = \frac{\mu k T}{q} \quad (2.5)$$

Where  $k$  is the Boltzmann constant and  $T$  is the lattice temperature. This results in a second term which contributes to carrier movement and it is called the diffusion current density.

$$J_D = qD \frac{dn}{dx} \quad (2.6)$$

Unlike the drift current density, which is directly related to the carrier density, diffusion current density is related to the carrier density's first order derivative in space.

Combining these two terms we get the current density equation for electrons in one dimension.

$$\vec{J}_n^x = q\mu_n n \vec{E}_x + qD_n \frac{dn}{dx} \quad (2.7)$$

This equation can be easily extended to higher dimensions by simply using the appropriate terms.

$$\vec{J}_n^y = q\mu_n n \vec{E}_x + qD_n \frac{dn}{dy} \quad (2.8)$$

Current density equations can be used for both positive and negatively charged particles by changing the sign of the diffusion current density.

$$\vec{J}_p^x = q\mu_p p \vec{E}_x - qD_p \frac{dp}{dx} \quad (2.9)$$

Anisotropic drift and diffusion coefficients can be handled with ease by using different coefficients for different directions. Also Mobility and diffusivity can be a function of any variable such as position or temperature. This is a useful property for a memristor simulation since hole mobility depends on lithium density in the PEDOT:PSS at any given position.

For the memristor simulation the movement of lithium and perchlorate ions inside the electrolyte is modeled as if they were moving in a solid state device. So all the derivations that were made for positive and negatively charged particles include holes and electrons as well as lithium and perchlorate ions. Viscosity of the liquid and all possible deformations due to movement of ions are simply ignored. This approximation was made based on the assumption that the amount of ions in the electrolyte are very large compared what PEDOT:PSS can absorb. Additionally the solution is very dilute. So the distribution of ions inside the electrolyte remain mostly unchanged and the exchange of ions do not disturb the shape or the physical characteristics of the solution on top of PEDOT:PSS.

Current density equations by themselves are not enough to solve a time dependent problem. It is necessary to account for the movement of charge over time which is captured in the continuity equation. It is basically a statement of conservation of particle density over time. The change in the amount of carriers over time in a particular area must be equal to the difference in current density over the same area. Additionally the amount of charge can change due to generation-recombination of charged particles.

$$\frac{\partial n}{\partial t} = \frac{1}{q} \nabla \cdot \vec{J}_n + U_n \quad (2.10)$$

$$\frac{\partial p}{\partial t} = -\frac{1}{q} \nabla \cdot \vec{J}_p + U_p \quad (2.11)$$

$U_n$  and  $U_p$  are net generation recombination rates. These terms were not used for the modeling of the memristor. (**Not sure how to justify this. Maybe I can say I have not used it because there is no equation/model for it ?**)

Electric field can be generated in two different ways. One is through the distribution of net charge over the area and the other one is an externally applied potential. It is possible calculate the potential distribution over an area by using Poisson's equation. Once the potential is known the electric field can be obtained by just calculating the negative gradient of the electric potential.

$$\nabla \cdot (\varepsilon \nabla V) = -\rho = -q(p - n + N_D^+ - N_A^-) \quad (2.12)$$

$$\vec{E} = -\nabla V \quad (2.13)$$

Electric field can be split into one, two or three components depending on the dimensions of the problem. In a 2-D case they are  $\vec{E}_x$  and  $\vec{E}_y$ .

$$\vec{E}_x = -\frac{\partial V}{\partial x} \quad (2.14)$$

$$\vec{E}_y = -\frac{\partial V}{\partial y} \quad (2.15)$$

Once the electric field and current density is known, the amount of current in a particular place in the device can be easily calculated by using the following integral.

$$I = \int_s \vec{J}_{tot} \cdot ds = \int_s (J_n + J_p + \varepsilon \frac{\partial E}{\partial t}) ds \quad (2.16)$$

Through the electric field and net charge distribution, Poisson's equation and drift-diffusion equations are coupled and non linear. The strength of the non linearity depends on the size of the device and charge density which determines the total amount of charge over an area. In a strongly coupled system of differential equations, small changes in either the charge density or electric field can easily cause instabilities in the simulation.

This is further discussed in the next chapter while developing a numerical scheme to solve the system of equations described in this chapter.

The main equations for the drift-diffusion model that are used for memristor simulation can be compactly written as:

$$\nabla \cdot (\varepsilon \nabla V) = -q(p - n + N_D^+ - N_A^-) \quad (2.17)$$

$$\vec{J}_p = q\mu_p p \vec{E} - qD_p \nabla p \quad (2.18)$$

$$\vec{J}_{N_A^-} = q\mu_{N_A^-} N_A^- \vec{E} + qD_{N_A^-} \nabla N_A^- \quad (2.19)$$

$$\vec{J}_{N_D^+} = q\mu_{N_D^+} N_D^+ \vec{E} - qD_{N_D^+} \nabla N_D^+ \quad (2.20)$$

$$\frac{\partial p}{\partial t} = -\frac{1}{q} \nabla \cdot \vec{J}_p \quad (2.21)$$

$$\frac{\partial N_A^-}{\partial t} = \frac{1}{q} \nabla \cdot \vec{J}_{N_A^-} \quad (2.22)$$

$$\frac{\partial N_D^+}{\partial t} = -\frac{1}{q} \nabla \cdot \vec{J}_{N_D^+} \quad (2.23)$$

This system of equations consist of 4 different charge carriers. There are 3 mobile species lithium ( $N_D^+$ ) and perchlorate ( $N_A^-$ ) and holes (p) and one fixed charge, electrons (n). The particles are coupled to each other through Poisson's equation since they all carry electric charge.

Unfortunately even after so many approximations and simplifications to BTE, drift diffusion equations have analytical solutions for only few isolated cases. Analytical solutions developed in the rest of this chapter will be used in testing of the numerical method developed for simulating a memristor.

## 2.2 Analytic Solutions

Three different analytic solutions to drift diffusion equations will be developed in this section. It starts with a simple steady state problem with uniform electric field. Second example, which is more complex than the first one, is a transient solution for a charge distribution moving under uniform electric field. Finally PN junction problem incorporates drift diffusion equation with Poisson's equation therefore provides a steady state solution for electron/hole density distributions, potential and electric field.

### 2.2.1 Steady State Solution Over a Finite Domain

The problem that will be solved in this section consists of a finite amount of charge and a uniform electric field pushing this charge against a wall ( $J=0$ ). It is assumed that the charge density is low enough not to affect the uniform electric field, therefore Poisson's equation is not needed. To simplify the problem even further, transient solution is ignored. Also it is important to note that the initial distribution of the charge density does not matter in this problem since everything will be redistributed in steady state. The only relevant information is the total amount of charge subject to the electric field. At steady state, drift and diffusion currents are in equilibrium and net current density is zero. Following equation shows hole density at equilibrium:

$$J_p(x) = q\mu_p pE - qD_p \frac{dp}{dx} = 0 \quad (2.24)$$

Total current density will only be zero when the drift current density generated by the electric field is completely balanced by the diffusion current density.

$$q\mu_p pE = qD_p \frac{dp}{dx} \quad (2.25)$$

$$\frac{\mu_p E}{D_p} p = \frac{dp}{dx} \quad (2.26)$$

This is a simple differential equation that can be solved by assuming that hole density has the following form:

$$p(x) = Ce^{ax} \quad (2.27)$$

$C$  and  $a$  are arbitrary constants. Now we can insert equation (2.27) into (2.26)

$$\frac{\mu_p E}{D_p} C e^{ax} = a C e^{ax} \quad (2.28)$$

$$a = \frac{\mu_p E}{D_p} \quad (2.29)$$

A general form for the solution is generated by placing  $\mathbf{a}$  back to equation (2.27).

$$p(x) = C e^{\frac{\mu_p E}{D_p} x} \quad (2.30)$$

It is possible to solve for  $\mathbf{C}$  by observing that the total number of charges at steady state must be equal to the initial number of charges. So the integral of the initial hole density distribution must be equal to the integral of charge density at steady state.

$$\int_0^L C e^{\frac{\mu_p E}{D_p} x} dx = \int_0^L p(t=0, x) dx$$

$$C = \frac{\int_0^L p(t=0, x) dx}{\int_0^L e^{\frac{\mu_p E}{D_p} x} dx}$$

$$C = \frac{\int_0^L p(t=0, x) dx}{\frac{D_p}{\mu_p E} [e^{\frac{\mu_p E}{D_p} L} - 1]} \quad (2.31)$$

So the full solution takes the following form:

$$p(x) = \frac{\int_0^L p(t=0, x) dx}{\frac{D_p}{\mu_p E} [e^{\frac{\mu_p E}{D_p} L} - 1]} e^{\frac{\mu_p E}{D_p} x} \quad (2.32)$$

The solution shows that increasing electric field will concentrate the charge density at the edge of the area. Physically this behavior is reasonable, since the force that is pushing the particles against a wall is getting greater and it is making the charges accumulate further. Following plot demonstrates the effect of increasing electric field on hole distribution.

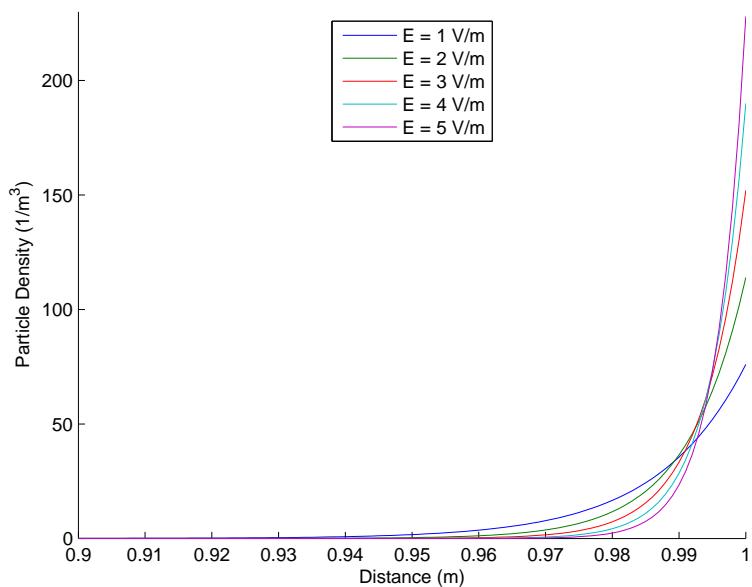


FIGURE 2.1: Increased accumulation of particles due to increased electric field

### 2.2.2 Transient Solution Over an Infinite Domain

The problem that will be solved in this section consists of an initial particle density distribution subject to a uniform electric field over an infinitely long conductor. The charge distribution will drift and diffuse over time. This requires a transient solution and the continuity equation (2.10) needs to be employed. Again, charged particles do not affect the electric field therefore solution of Poisson's equation is not necessary.

The first step of the solution process is the insertion of (2.7) into (2.10).

$$\frac{\partial n}{\partial t} = \frac{1}{q} \nabla \cdot (\vec{J}_n) \quad (2.33)$$

$$\frac{\partial n}{\partial t} = \frac{1}{q} \nabla \cdot (q\mu_n n \vec{E} + qD_n \frac{dn}{dx}) \quad (2.34)$$

For 1-D above equation can be simplified to:

$$\frac{\partial n}{\partial t} = \mu_n E \frac{dn}{dx} + D_n \frac{d^2 n}{dx^2} \quad (2.35)$$

The separation of variables can be used which assumes that the solution can be separated into a time and space dependent functions.

$$n(t, x) = n(t)n(x) = n_t n_x \quad (2.36)$$

Placing equation (2.36) into (2.35) and dividing by  $n(t,x)$ ,

$$\frac{1}{n_t} \frac{dn_t}{dt} = \mu_n E \frac{1}{n_x} \frac{dn_x}{dx} + D \frac{1}{n_x} \frac{d^2 n_x}{dx^2} \quad (2.37)$$

Assuming both sides of the equation are equal to a constant  $-k$ , time dependent part of the problem becomes a simple first order differential equation.

$$\frac{1}{n_t} \frac{dn_t}{dt} = -k$$

$$\frac{dn_t}{dt} = -kn_t$$

Based on the above differential equation  $n_t$  can take the following form:

$$n_t = C_1 e^{-kt} \quad (2.38)$$

Now for  $n_x$  assuming a solution of the form below,

$$n_x = C_2 e^{-j\omega x} \quad (2.39)$$

Placing (2.39) into (2.37)

$$\omega^2 D C_2 e^{-j\omega x} - j\omega \mu_n E C_2 e^{-j\omega x} + k C_2 e^{-j\omega x} = 0 \quad (2.40)$$

Simplifying equation (2.40) and solving for  $k$  gives,

$$k = \omega^2 D + j\omega \mu_n E \quad (2.41)$$

Combining equation (2.38), (2.39) and (2.41) to get the initial form of the solution.  
( $C = C_1 C_2$ )

$$n = n_t n_x = C e^{(-\omega^2 D + j\omega \mu_n E)t} e^{-j\omega x} \quad (2.42)$$

The application of superposition principle leads to:

$$n = n_t n_x = \int_{-\infty}^{\infty} C(\omega) e^{(-\omega^2 D + j\omega \mu_n E)t} e^{-j\omega x} d\omega \quad (2.43)$$

The distribution of  $n_x$  is known at  $t=0$ .

$$n(x, t=0) = \int_{-\infty}^{\infty} C(\omega) e^{-j\omega x} d\omega \quad (2.44)$$

$C(\omega)$  is just the inverse fourier transform of  $n(x, 0)$ .

$$C(\omega) = \int_{-\infty}^{\infty} n(x, 0) e^{-j\omega x} dx \quad (2.45)$$

The final form of the solution is generated by placing equation (2.45) into (2.43):

$$n = \int_{-\infty}^{\infty} \int_{-\infty}^{\infty} n(z, 0) e^{j\omega z} dz e^{(\omega^2 D - j\omega \mu_n E)t} e^{-j\omega x} d\omega \quad (2.46)$$

Rearranging equation (2.46),

$$n = \int_{-\infty}^{\infty} \int_{-\infty}^{\infty} n(z, 0) e^{(\omega^2 D - j\omega \mu_n E)t} e^{-j\omega x} e^{j\omega z} dz d\omega \quad (2.47)$$

Using a gaussian initial distribution results into:

$$n(x, 0) = e^{-(\frac{x-x_0}{\sigma})^2} \quad (2.48)$$

$$n(x, t) = \frac{1}{\sqrt{4D_n \sigma^{-2} t + 1}} e^{-\frac{(t\mu_n E - x + x_0)^2}{4D_n t + \sigma^2}} \quad (2.49)$$

If the initial distribution is a rectangular then the solution takes the following form:

$$n(x, 0) = \prod (w(x)) \quad (2.50)$$

$$n(x, t) = \frac{1}{2} \operatorname{erf}\left(\frac{w + 2t\mu_n E - 2x}{4\sqrt{D_n t}}\right) - \frac{1}{2} \operatorname{erf}\left(\frac{-w + 2t\mu_n E - 2x}{4\sqrt{D_n t}}\right) \quad (2.51)$$

Following plots show the evolution of two particle densities with different initial distributions described above.

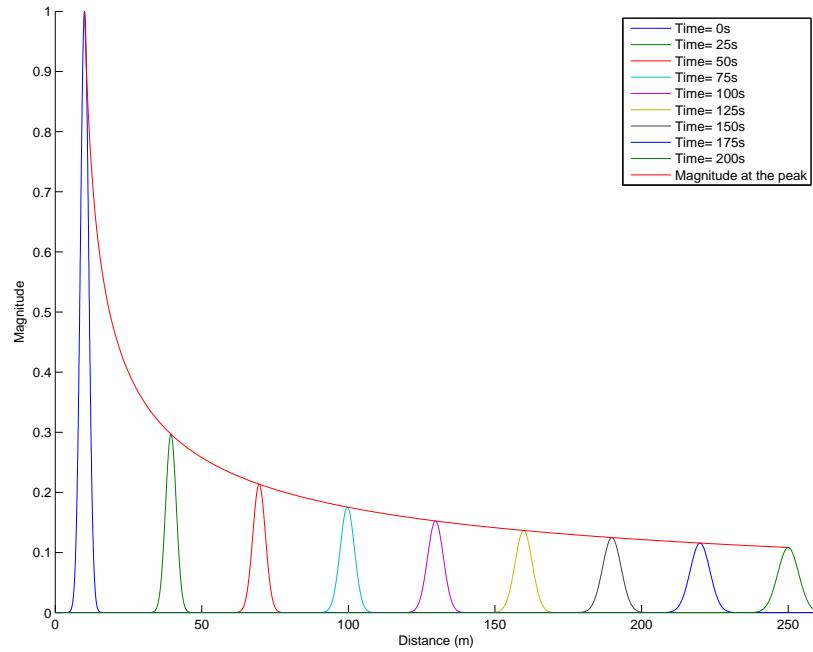


FIGURE 2.2: A gaussian particle density distribution drifting and diffusion over time

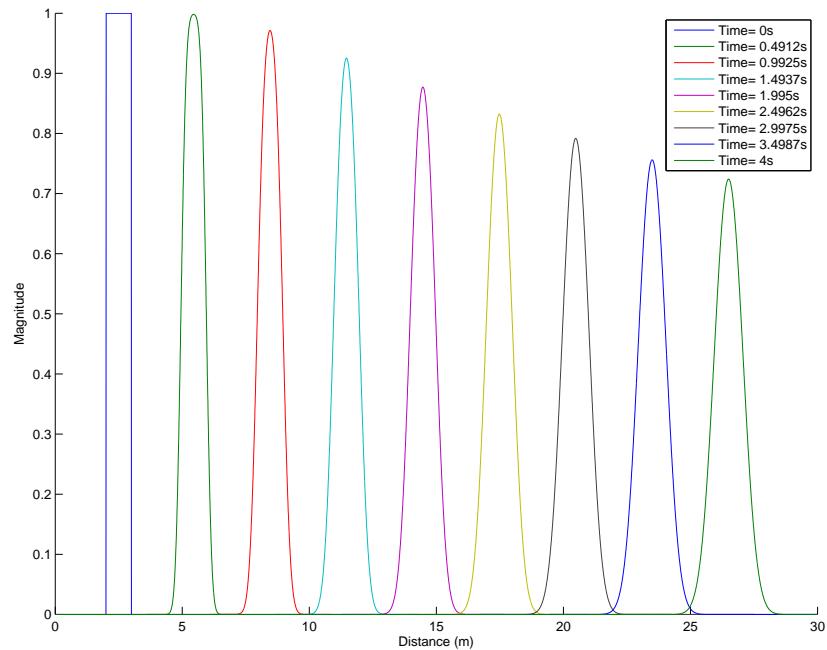


FIGURE 2.3: A gaussian particle density distribution drifting and diffusion over time

### 2.2.3 PN Junction

Previous analytical solutions involved the solution of Poisson's equation and continuity equation which were not coupled. Now we can look at an example where these equations are tightly coupled. There are usually no direct analytical solution for coupled equation but it is possible to get a closed form solution by making use of certain approximations. One simple example of this situation is an abrupt p-n junction. An abrupt p-n junction is created when two materials of uniform opposite doping, p type and n type, are brought together. A p type material has an excess number of acceptors  $N_A$  and an n type material has an excess number of donors  $N_D$ . The junction is defined at the interface where  $N_A = N_D$ .

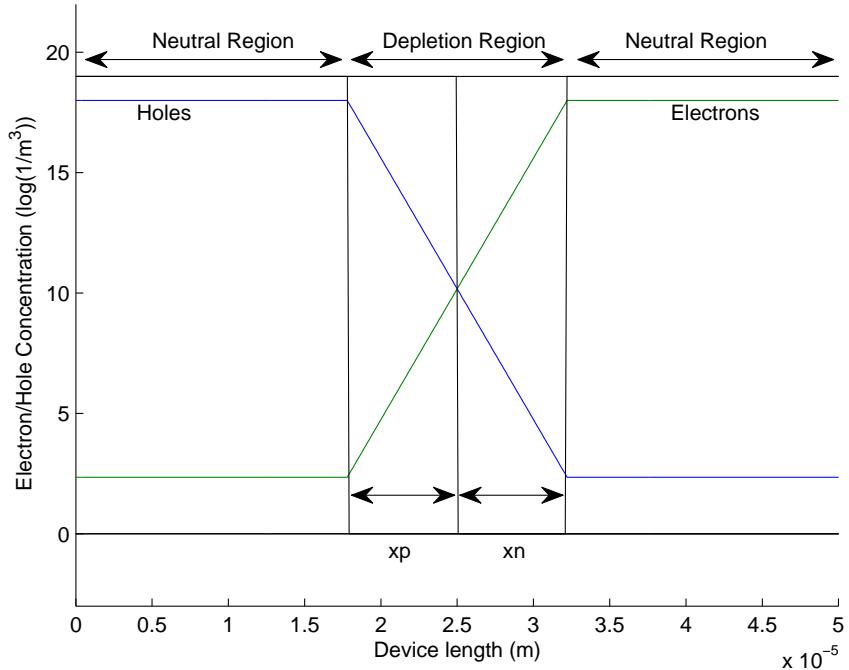


FIGURE 2.4: PN junction electron/hole density

In this example an analytical solution for an abrupt p-n junction will be derived. To get a solution for this problem depletion region approximation we will be used. This approximation starts by assuming that the charges are fully depleted around the junction. All the electric field is confined in the depletion region and regions far away from the junction are neutral. Based on this the net charge density over the entire region is:

$$\frac{d\vec{E}}{dx} = \frac{\rho}{\varepsilon} = \frac{q}{\varepsilon}(-N_A + N_D) \quad (2.52)$$

$$\rho = \begin{cases} 0 & \text{for } x < -x_p \\ -qN_A & \text{for } -x_p \leq x \leq 0 \\ qN_D & \text{for } 0 \leq x \leq x_n \\ 0 & \text{for } x > x_n \end{cases} \quad (2.53)$$

The electric field over the entire region can be calculated by integrating  $\rho$ .

$$E = \begin{cases} \int \frac{-qN_A}{\varepsilon} dx + C_1 & \text{for } -x_p \leq x \leq 0 \\ \int \frac{qN_D}{\varepsilon} dx + C_2 & \text{for } 0 \leq x \leq x_n \end{cases} \quad (2.54)$$

It is possible to solve for  $C_1$  and  $C_2$  since electric field must go to zero at  $x_p$  and  $x_n$ .

$$E(x = -x_p) = 0 \Rightarrow C_1 = \frac{-qN_A}{\varepsilon} x_p \quad (2.55)$$

$$E(x = x_n) = 0 \Rightarrow C_2 = \frac{qN_D}{\varepsilon} x_n \quad (2.56)$$

Then  $E(x)$  becomes:

$$E(x) = \begin{cases} \frac{-qN_A}{\varepsilon}(x + x_p) & \text{for } -x_p \leq x \leq 0 \\ \frac{qN_D}{\varepsilon}(x - x_n) & \text{for } 0 \leq x \leq x_n \end{cases} \quad (2.57)$$

Additionally, the electric field must be continuous across the interface therefore the electric field in the p-type side and the n-type side must equal each other at the interface or when  $x = 0$ .

$$\frac{-qN_A}{\varepsilon}(x_p) = \frac{qN_D}{\varepsilon}(-x_n) \quad (2.58)$$

$$N_A x_p = N_D x_n \quad (2.59)$$

This equation makes physical sense since it states that the total charge on one side of the junction must be the same as the total charge on the other. In other words, the net charge on each side keeps the electric field confined to the depletion region.

To find the voltage as a function of distance, equation 2.13 can be integrated.

$$V(x) = \begin{cases} \int -E(x)dx = \int \frac{qN_A}{\varepsilon}(x + x_p)dx = \frac{qN_A}{\varepsilon}(\frac{x}{2} + x_p) + C_3 & \text{for } -x_p \leq x \leq 0 \\ \int -E(x)dx = \int \frac{qN_D}{\varepsilon}(x - x_n)dx = \frac{qN_D}{\varepsilon}(-\frac{x}{2} + x_n) + C_4 & \text{for } 0 \leq x \leq x_n \end{cases} \quad (2.60)$$

The potential at one side of the junction can be set to zero. Defining the voltage on the p type side as zero, such that at  $x = x_p$ ,  $V=0$ . This gives the constant  $C_3$  as:

$$C_3 = \frac{qN_A}{2\varepsilon}x_p^2 \quad (2.61)$$

$$V(x) = \frac{qN_A}{2\varepsilon}(x + x_p)^2 \quad \text{for } -x_p \leq x \leq 0 \quad (2.62)$$

$C_4$  can be found by using the fact that the potential on the n-type side and p-type side are equal at the interface, such that:

$$V_p(x = 0) = \frac{qN_A}{2\varepsilon}x_p^2 = V_n(x = 0) = \frac{qN_A}{2\varepsilon}(x_n - \frac{x}{2})x + C_4 \quad (2.63)$$

$$C_4 = \frac{qN_A}{2\varepsilon}x_p^2 \quad (2.64)$$

Now an overall expression for  $V(x)$  can be obtained.

$$V(x) = \begin{cases} \frac{qN_A}{\varepsilon}(x + x_p)^2 & \text{for } -x_p \leq x \leq 0 \\ \frac{qN_D}{\varepsilon}(-\frac{x}{2} + x_n)x & \text{for } 0 \leq x \leq x_n \end{cases} \quad (2.65)$$

The maximum voltage across the junction is at  $x = x_n$ , which is:

$$V_{bi} = \frac{q}{2\varepsilon}(N_Dx_n^2 + N_Ax_p^2) \quad (2.66)$$

Using (2.59) in the above equation and rearranging allows  $x_p$  and  $x_n$  to be determined. They are:

$$x_n = \sqrt{\frac{2\varepsilon V_{bi}}{q} \frac{N_A}{N_D(N_D + N_A)}} \quad (2.67)$$

$$x_p = \sqrt{\frac{2\varepsilon V_{bi}}{q} \frac{N_D}{N_A(N_D + N_A)}} \quad (2.68)$$

The value of the built in potential can also be calculated using fermi levels of p and n doped materials.

$$E_{FN} - E_i = kT \ln\left(\frac{N_D}{n_i}\right) \quad (2.69)$$

$$E_i - E_{FP} = kT \ln\left(\frac{N_A}{n_i}\right) \quad (2.70)$$

$E_{FN}$  and  $E_{FP}$  are fermi energy levels of electrons and holes respectively. The difference between the fermi levels divided by the single electron charge gives us the built in potential of the pn junction.

$$E_{FN} - E_{FP} = qV_{bi} = kT \ln\left(\frac{N_D}{n_i}\right) + kT \ln\left(\frac{N_A}{n_i}\right) = kT \ln\left(\frac{N_A N_D}{n_i^2}\right) \quad (2.71)$$

$$V_{bi} = \frac{kT}{q} \ln\left(\frac{N_A N_D}{n_i^2}\right) \quad (2.72)$$

The calculation of the built in potential completes all the necessary equations for the analytical solution of the pn junction without any external bias. Following graphs shows the plots of approximate solutions for net charge, electric field and the junction potential.

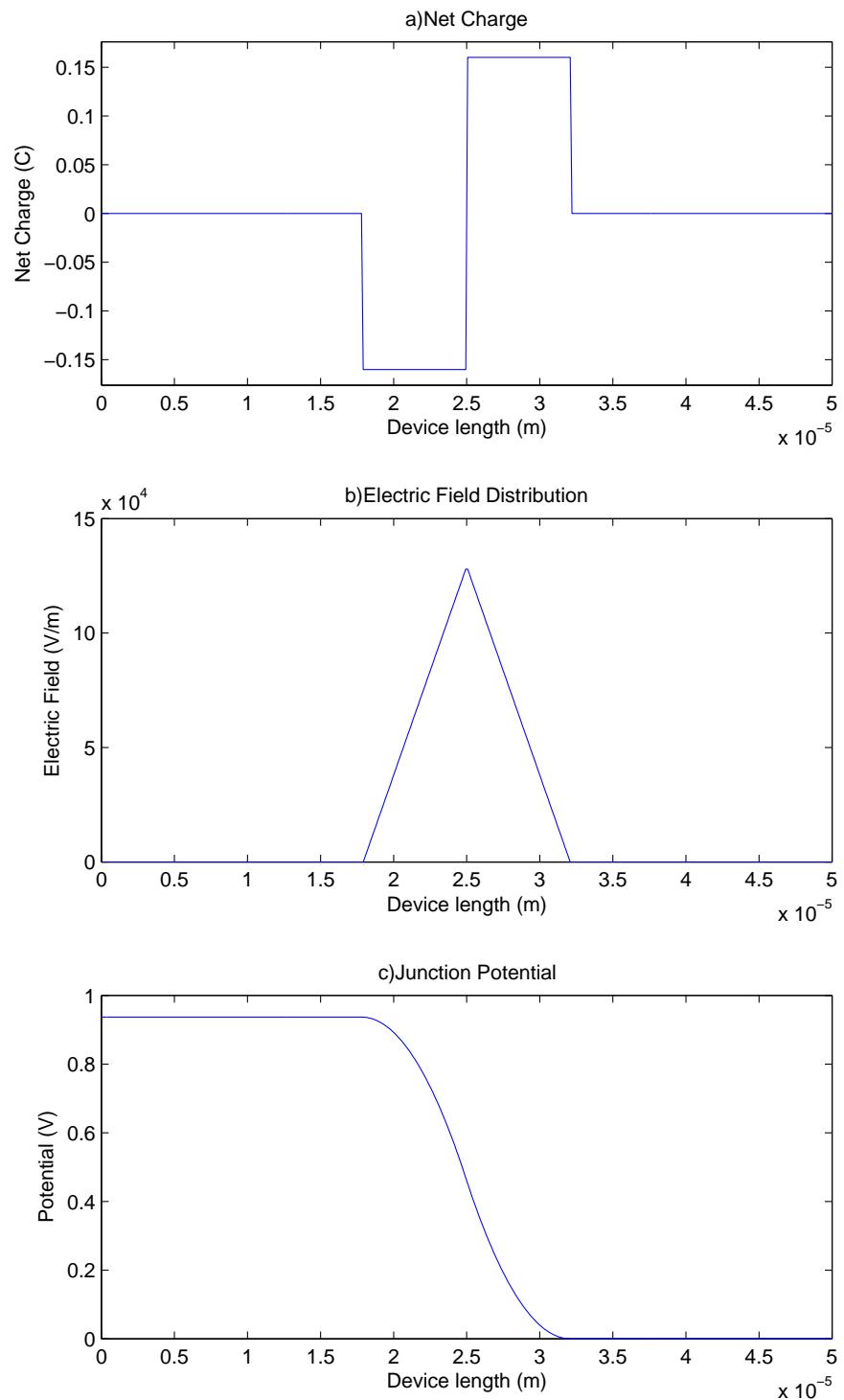


FIGURE 2.5: Approximate Solution of a PN Junction

## Chapter 3

# Numerical Solution of Drift-Diffusion Equation

As discussed in the introduction, physical memristor simulations are almost non existent. Memristor simulation developed in this thesis requires certain properties ,such as limitation of particle density or combination of 1-D and 2-D simulations, which may not available in most commercial simulators. It is also possible that other unusual properties that are not studied in this thesis will be needed for further research. For this reason, in this chapter a method for solving drift diffusion equations as well as Poisson's equation will be developed based on finite difference. This fully functioning simulator will have needed flexibility for current and future research opportunities.

### 3.1 Finite Difference Method

There are many different methods that can be used to solve drift diffusion equations such as finite elements, finite difference or meshless methods. Finite difference was chosen as an appropriate method for this thesis due to its simplicity which will allow straightforward implementation of unusual physical properties. Finite difference uses an approximation for the derivative of a function based on the mathematical definition of the derivative.

$$\frac{df}{dx} = \lim_{h \rightarrow 0} \frac{f(x + h) - f(x)}{h} \quad (3.1)$$

It is possible to obtain a numerical approximation for the first derivative by dropping the limit and assuming that  $h$  is small enough to get a value for the derivative reasonably

close to the actual value. As  $h$  gets smaller the approximation becomes more and more accurate. The difference between the calculated value and the real value is called the truncation error and it is captured using  $O(h^n)$  notation.  $n$  signifies the order of  $h$  which determines how fast the approximation is approaching the real solution as  $h$  decreases.

$$\frac{df}{dx} = \frac{f(x+h) - f(x)}{h} + O(h) \quad (3.2)$$

It is possible to uniformly discretise the entire region over which a function is defined in order to calculate its derivative. The first step is the division of the region over which the function is defined into  $n-1$  segments therefore there will be  $n$  number of points. Then the length of each segment is defined using the following relationship:

$$h = \frac{L}{n} \quad (3.3)$$

A function is defined at the edge of every segment. All the points can be labeled consecutively,  $x_0, x_1, x_2 \dots x_{n-1}$  where  $x_i = ih$ . The function is discretely defined on  $f_i = f(x_i)$  where  $i=0,1,2..n-1$ . It is possible to use equation 3.2 to discretely calculate the first derivative of the function with respect to  $x$ .

$$\frac{df(x_i)}{dx} = \frac{f(x_{i+1}) - f(x_i)}{h} + O(h) \quad (3.4)$$

Above equation is called forward difference because the derivative for point  $x_i$  was calculated using the point that is coming right after it,  $x_{i+1}$ .

$$f'_i = \frac{f_{i+1} - f_i}{h} + O(h) \quad (3.5)$$

Forward difference is not the only way to calculate a discrete derivative. Here are a few other ways calculate the same derivative by making use of different points.

$$f'_i = \frac{f_i - f_{i-1}}{h} + O(h) \quad (3.6)$$

$$f'_{i+\frac{1}{2}} = \frac{f_{i+1} - f_i}{h} + O(h^2) \quad (3.7)$$

Equation 3.6 is called backward difference and equation 3.7 is called central difference. One important thing to note here is that in the central difference formula the derivative falls exactly in the middle of two points. It also gives more accurate results using the same number of points as forward and backward difference.

Using finite difference formulas it is possible to construct higher derivatives. A formula for a second order derivative at point  $x_i$  using central difference can be calculated using first order derivative on  $x_{i-\frac{1}{2}}$ ,  $x_i$  and  $x_{i+\frac{1}{2}}$

$$f'_{i+\frac{1}{2}} = \frac{f_{i+1} - f_i}{h} \quad (3.8)$$

$$f'_{i-\frac{1}{2}} = \frac{f_i - f_{i-1}}{h} \quad (3.9)$$

$$f'_i = \frac{f_{i+\frac{1}{2}} - f_{i-\frac{1}{2}}}{h} \quad (3.10)$$

Second order derivative is constructed by taking the second derivative of the last function. Then equations 3.8 and 3.9 is placed into 3.10.

$$f''_i = \frac{f'_{i+\frac{1}{2}} - f'_{i-\frac{1}{2}}}{h}$$

$$f''_i = \frac{\frac{f_{i+1}-f_i}{h} - \frac{f_i-f_{i-1}}{h}}{h}$$

$$f''_i = \frac{f_{i+1} - f_i - f_i + f_{i-1}}{h^2}$$

Second order derivative takes the following form:

$$f''_i = \frac{f_{i+1} - 2f_i + f_{i-1}}{h^2} + O(h^2) \quad (3.11)$$

Overall these finite difference equations are enough to solve drift diffusion equations. Even though all the derivations were done in 1-D it is trivial to extend them to higher dimensions. This method can be used to solve Poisson's equation and drift diffusion equations.

## 3.2 Poisson Solver

Poisson's equation needs to be solved before drift diffusion equations in order find the potential distribution as well as the electric field inside the device. In order to solve for electric field and potential Poisson's equation will be simplified through assumptions and then finite difference will be used to solve this simplified equation. The first step of simplification is assuming that the permittivity is isotropic.

$$\nabla \cdot (\varepsilon \nabla V) = -\rho \quad (3.12)$$

$$\nabla \cdot (\varepsilon \nabla V) = \varepsilon \nabla^2 V \quad (3.13)$$

Dividing both sides by permittivity and expanding the left hand side,,

$$\nabla^2 V = -\frac{\rho}{\varepsilon} \quad (3.14)$$

$$\nabla^2 V = \frac{\partial^2 V}{\partial x^2} + \frac{\partial^2 V}{\partial y^2} \quad (3.15)$$

After discretizing electric potential over a 2-D uniform grid and using the second order central finite difference formula (3.11) laplacian of the electric potential can be calculated using:

$$\nabla^2 V_{i,j} = \frac{V_{i+1,j} - 2V_{i,j} + V_{i-1,j}}{\Delta x^2} + \frac{V_{i,j+1} - 2V_{i,j} + V_{i,j-1}}{\Delta y^2} \quad (3.16)$$

Since the grid is uniform the distance between two nodes in x and y directions are equal only one variable is need for distance between two points.

$$\Delta = \Delta x = \Delta y \quad (3.17)$$

Net charge density and the permittivity is also discretized over the same uniform mesh. Discretized form of Poisson's equation is generated by combining equations 3.14 ,3.16 and 3.17.

$$\nabla^2 V_{i,j} = \frac{V_{i-1,j} + V_{i,j-1} - 4V_{i,j} + V_{i+1,j} + V_{i,j+1}}{\Delta^2} = -\frac{\rho_{i,j}}{\varepsilon_{i,j}} \quad (3.18)$$

This equation can be rearranged into the form below:

$$\varepsilon_{i,j}(V_{i-1,j} + V_{i,j-1} - 4V_{i,j} + V_{i+1,j} + V_{i,j+1}) = -\Delta^2 \rho_{i,j} \quad (3.19)$$

Equation 3.19 is valid for almost all the nodes in the system except two cases, boundary nodes and interface nodes. There are two different types of boundary conditions. The first one is Dirichlet boundary condition which forces a particular value for the potential at the boundary.

$$V_{i,j} = V_b \quad (3.20)$$

Where  $V_b$  is the value of the potential at the boundary. The other possible boundary condition is called Neumann boundary condition which states that the derivative of the potential at the boundary is zero. This gives the following equation:

$$\frac{\partial V}{\partial x} = \frac{V_{i+1,j} - V_{i,j}}{\Delta} = 0 \quad (3.21)$$

So for a boundary over y axis:

$$V_{i+1,j} = V_{i,j} \quad (3.22)$$

Neumann boundary condition over x axis is obtained using the same procedure.

$$V_{i,j+1} = V_{i,j} \quad (3.23)$$

Combining the equations above (3.19, 3.20, 3.21 and 3.22) it is possible to turn Poisson's equation, which is a second order differential equation, into a linear equation.

$$D_2 \vec{V} = -\Delta^2 \vec{\rho} - \vec{V}_b \quad (3.24)$$

One can easily get the potential distribution by simply solving the matrix equation obtained. Due to the nature of the problem resulting matrix is quite sparse and using a sparse LU decomposition dramatically increases the computational efficiency.

$$\vec{V} = D_2^{-1}(-\Delta^2 \vec{\rho}_{i,j} - \vec{V}_b) \quad (3.25)$$

After solving for the potential distribution it is easy to calculate the electric field distribution discretely using the relationship between electric field and electric potential (2.13) and central difference equation (3.7).

$$\vec{E}_{i,j} = -\frac{V_{i+1,j} - V_{i-1,j}}{2\Delta} \quad (3.26)$$

$$\vec{E}_{i,j} = -\frac{V_{i,j+1} - V_{i,j-1}}{2\Delta} \quad (3.27)$$

### 3.3 Current Density Equations

Both drift and diffusion currents can be calculated over the entire grid. Drift current does not involve any differentials but it is a function of electric field and the diffusion current can be calculated using first order central difference. The current density is calculated in such a way that it falls between two points which simplifies the application of the boundary conditions.

$$J_{i+\frac{1}{2},j,k}^x = q\mu_n n_{i+\frac{1}{2},j,k} E_{i+\frac{1}{2},j,k}^x + D_n \frac{n_{i+1,j,k} - n_{i,j,k}}{\Delta} \quad (3.28)$$

The electric field was calculated exactly on the nodes and linear interpolation was used in order to get a value between the nodes. Same argument is also valid for particle densities  $p$  and  $n$ . They were defined on the nodes but can be linearly interpolated.

$$\begin{aligned} n_{i+\frac{1}{2},j,k} &= \frac{n_{i+1,j,k} + n_{i,j,k}}{2} \\ E_{i+\frac{1}{2},j,k}^x &= \frac{E_{i+1,j,k}^y + E_{i,j,k}^y}{2} \end{aligned}$$

Current density in y direction is calculated following the same method:

$$\begin{aligned} J_{i,j+\frac{1}{2},k}^y &= q\mu_n n_{i,j+\frac{1}{2},k} E_{i,j+\frac{1}{2},k}^y + D_n \frac{n_{i,j+1,k} - n_{i,j,k}}{\Delta} \quad (3.29) \\ n_{i,j+\frac{1}{2},k} &= \frac{n_{i,j+1,k} + n_{i,j,k}}{2} \\ E_{i,j+\frac{1}{2},k}^y &= \frac{E_{i,j+1,k}^y + E_{i,j,k}^y}{2} \end{aligned}$$

#### 3.3.1 Boundary Conditions

For a drift diffusion problem there are few different possibilities for boundary conditions. The first one is when the particles cannot go past through a certain boundary, no flow boundary condition. This can be achieved by using Dirichlet boundary condition and setting the flow at the boundary to zero.

$$J = 0 \quad (3.30)$$

For memristor simulation this boundary condition is used to conditionally limit the lithium density. The implementation of the limiting process is discussed in detail and tested thoroughly in section 4 of chapter 4. Dirichlet boundary condition is also used

for metal contacts. It is assumed that metal contacts have infinite amount of charge and the boundary is always charge neutral. For example for holes, electrons, positive and negative doping it is assumed that at the boundary positive charge concentration will be equal to the negative charge concentration.

$$N_D + p = N_A + n \quad (3.31)$$

For a semiconductor holes and electrons has to obey mass action law.

$$np = n_i^2 \quad (3.32)$$

$n_i$  here is the concentration of the semiconductor at equilibrium before getting doped. Solving (3.32) and (3.31) together results in the equation below:

$$n = \frac{1}{2}(N_D - N_A + \sqrt{(N_D - N_A)^2 + 4n_i^2}) \quad (3.33)$$

Once electron concentration is obtained it is possible to calculate the hole concentration using mass action law.

$$p = \frac{n_i^2}{n} \quad (3.34)$$

During simulation, the application of this boundary condition differs from the no flow boundary which was applied implicitly. All the boundaries are simulated using a no flow condition but their values are set to the appropriate value at the end of every time step. Lack of charge is compensated and excess charge is taken off by the metal contact. The difference between the boundary value of the charge density and its actual value is used to calculate the derivative of the current density with respect to time.

$$\frac{dn}{dt} = \frac{n_{cr} - n_{ct}}{\Delta t} \quad (3.35)$$

$n_{cr}$  is the excess carrier density and  $n_{ct}$  is the equilibrium carrier density at the contact. Using these boundary conditions for holes and electrons and the following two equations it is possible to calculate incoming and outgoing currents.

$$Q = An \quad (3.36)$$

$$I = \frac{dQ}{dt} \quad (3.37)$$

Equation 3.36 is an approximate relationship between charge and charge density where Q is total charge, A is the area holding that charge. Equation 3.37 is simply the general definition of current. By combining these two equations it is possible to derive a formula for calculating current at any metal contact.

$$I = A \frac{dn}{dt} \quad (3.38)$$

### 3.3.2 Continuity Equation

Continuity equation is needed to calculate a transient solution for drift diffusion equations. The equation is simple to discretize using finite difference method. There are two terms that need to be discretized, a first order derivative in time and space. First the divergence term in equation (2.10) needs to be evaluated.

$$\nabla \cdot J = \frac{\partial J}{\partial x} + \frac{\partial J}{\partial y} = \frac{dJ_x}{dx} + \frac{dJ_y}{dy} \quad (3.39)$$

It is possible to replace the derivative with central finite difference terms.

$$\frac{dJ_x}{dx} = \frac{J_{i+\frac{1}{2},j,k}^x - J_{i-\frac{1}{2},j,k}^x}{h} \quad (3.40)$$

$$\frac{dJ_y}{dy} = \frac{J_{i,j+\frac{1}{2},k}^y - J_{i,j-\frac{1}{2},k}^y}{h} \quad (3.41)$$

$$\nabla \cdot J_{i,j,k} = \frac{J_{i+\frac{1}{2},j,k}^x - J_{i-\frac{1}{2},j,k}^x}{h} + \frac{J_{i,j+\frac{1}{2},k}^y - J_{i,j-\frac{1}{2},k}^y}{h} \quad (3.42)$$

This is the general form of the divergence of the current density. Placing current density formulas in equation 3.42 and collecting all the terms in a matrix gives rise to the following equation:

$$\nabla \cdot J_k = B \quad (3.43)$$

The time derivative can also be replaced by a forward or backward finite difference terms respectively.

$$\frac{\partial \vec{n}_k}{\partial t} = \frac{\vec{n}_{k+1} - \vec{n}_k}{\Delta t} \quad (3.44)$$

$$\frac{\partial \vec{n}_k}{\partial t} = \frac{\vec{n}_k - \vec{n}_{k-1}}{\Delta t} \quad (3.45)$$

It is possible to find a numerical transient solution for the drift-diffusion problem by combining finite difference form of the time derivative ((3.44) or (3.45)) and the divergence of the current density equations (3.43).

Forward difference approximation can be used to get an explicit solution for continuity equation.

$$\frac{\vec{n}_{k+1} - \vec{n}_k}{\Delta t} = B$$

$$\vec{n}_{k+1} = \vec{n}_k + \Delta t B \quad (3.46)$$

Both forward and backward difference formulas work sequentially in order to generate a transient solution. Solution from the previous time step is needed to calculate the next time step. Forward difference gives an explicit solution which has few advantages. This solution can be implemented, without forming any matrices, by directly calculating the divergence of the current density for each node and then marching through time using equation 3.46. Additionally, unlike backward difference, there are no equations to be solved for every time step. These two properties ease the computational load of the problem and speed up the solution process. Unfortunately this scheme has very strict stability conditions which has to be met in order to get a solution.

Using backward difference can be used instead of forward difference to get an implicit solution.

$$\begin{aligned}\frac{\vec{n}_k - \vec{n}_{k-1}}{\Delta t} &= \frac{1}{q}(qB) \\ \vec{n}_k - \Delta t B &= \vec{n}_{k-1}\end{aligned}$$

Since all the equations in B matrix are linear  $n$  can be taken and out of the parenthesis ( $B = Cn$ ).

$$\begin{aligned}\vec{n}_k - \Delta t C \vec{n}_{k-1} &= \vec{n}_{k-1} \\ \vec{n}_k &= (I - \Delta t C)^{-1} \vec{n}_{k-1}\end{aligned} \quad (3.47)$$

This solution needs a matrix inversion every time step but it is unconditionally stable if not used together with Poisson's equation. The decision to use implicit or explicit solution is not very simple and it will be discussed in detail the next section.

## 3.4 Stability and Computational Efficiency

Before discussing numerical limitations of solving drift diffusion equation via finite difference it is important to look into physical limitations of the problem. These limitations persist no matter what kind of numerical scheme is employed to solve drift diffusion equations.

### 3.4.1 Physical Limitations

Debye length is the length over which mobile charge carriers screen out an external electric field. It determines how steeply charges will accumulate over a certain distance when subject to an electric field.

$$L_D = \sqrt{\frac{\epsilon V_{th}}{qn}} \quad (3.48)$$

Debye length limits how coarse the grid can be since we need to be able to accurately capture the distribution of charge density. As it can be seen from the formula above the higher the charge density is the steeper the charge will accumulate. This can become a major problem for device sizes in millimetre range or higher and high charge concentrations since the mesh density needs to be extremely high.

The amount of time it takes for charge fluctuations to disappear is called Dielectric relaxation time. It limits the maximum time step of a simulation since the fluctuations that are not properly resolved over time will make the simulation unstable.

$$t_{dr} = \frac{\epsilon}{qn\mu} \quad (3.49)$$

Dielectric relaxation time is only important when electric potential is highly affected by redistribution of charge over time. Otherwise it has minimal impact on the stability of the problem.

### 3.4.2 Numerical Limitations

There are also numerical limits which can affect convergence and stability of a solution when using an explicit scheme. These are called Courant-Friedrichs-Lowy (CFL) conditions. We can look at CFL conditions for pure diffusion and pure drift.

$$\frac{\Delta^2}{2D_n} > \Delta t \quad (3.50)$$

Above condition is for pure diffusion and it restricts the maximum time step. Following condition is for drift dominated systems:

$$\frac{2\Delta}{\mu E} > \Delta t \quad (3.51)$$

This is the second numerical restriction on our simulation. Interestingly the condition for drift depends on the electric field therefore it needs to be satisfied as the electric field changes over time during simulation.

Both physical and numerical constraints have to be evaluated and mesh density and time step need to be selected in order to satisfy all these conditions discussed above. Particularly mesh density have a very strong effect on the accuracy, stability and computational efficiency of the simulation. Increasing mesh density increases the computational time needed to calculate every time step since there are more points. Additionally because of the CFL condition for diffusion time step is related to the square of the mesh size. This means that maximum allowed step size decreases much quicker than the mesh density. Also, increasing charge density can decrease the maximum mesh size to a very small value. This can be fixed by using a non uniform mesh which can dramatically decrease the amount points needed for the simulation. Unfortunately this is usually not very straightforward to implement in a finite difference scheme. Also small mesh size requires small time steps. This cannot be avoided through non uniform meshing. Numerical and physical constraints in memristor simulation are further discussed in chapter 5.

### 3.4.3 Explicit vs. Implicit Solution

Overall explicit and implicit solutions have their advantages and disadvantages. Choosing one over the other requires careful analysis of the problem. Implicit solution by itself is unconditionally stable therefore it can support very large time steps without any stability issues. However with increased time step, the accuracy of the transient solution decreases but the steady state solution does not get affected. So for steady state solutions it is better to use an implicit method which can reach steady state very quickly. This advantage disappears when particle densities are high enough to affect the electric field and Poisson's equation needs to be solved for every time step. In this scenario the maximum step size is determined by dielectric relaxation time which tends

to be around the same order as CFL conditions. Now since the time step is going to be around the same order for both implicit and explicit methods it makes sense to use the explicit one because it is computationally less expensive.

Usually implicit solution is preferable when there is no coupling between Poisson's equation and drift diffusion equations and the transient response is not very important. Explicit solution have an edge over the implicit solution due to its lower computational resource requirement when the equations are coupled and the time steps for both schemes are restricted to fairly small values. For memristor simulation, drift diffusion equations are strongly coupled with Poisson's equation. For this reason all memristor simulations use explicit time stepping.

### 3.5 Equations Used for Memristor Simulation

### 3.6 Simulation Procedure

Different equations and schemes that are used to solve drift diffusion and Poisson's equation was shown over the past few sections. Using all this information it is possible create a general approach to solve a drift diffusion problem.

The geometry and physical properties of the problem as well as all initial and boundary conditions needs to be defined at the beginning of the solution process. Initialization sets up the first time step of the problem at  $t = 0$ . Once this first step is done it is possible generate required vectors and matrices and solve the problem for the next time steps,  $t = t_i$ .

The solution process starts by solving Poisson's equation using the charge distribution at current time step. Once it is solved, the electric field distribution is calculated and used in drift diffusion equations to calculate current density distribution. Next explicit time stepping is used to determine carrier density in the next time step. Finally once the carrier distribution for the next step is calculated it is possible check for a stopping criterion. If this criterion is not met then the whole process will start all over again with a small difference. If the charge concentration is so small that the equations are decoupled then it is possible to skip solving Poisson's equation for the rest of the simulation which speeds up the solution process.

There are two different criteria that can be used to decide weather to finalize the simulation process or not. The simulation can stop if it reaches a certain time point. This is quite simple since the current time can be checked and if it is equal or greater than the required simulation time then the simulation can be stopped. It is also possible to simulate until the simulation reaches steady state. This can be determined by comparing the current carrier distributions with distribution at the previous time step. If the difference is very small then the time derivatives of the carrier densities are very close to zero and the simulation has reached steady state therefore the simulation process can be stopped. The flowchart in figure 3.1 summarizes the solution procedure.

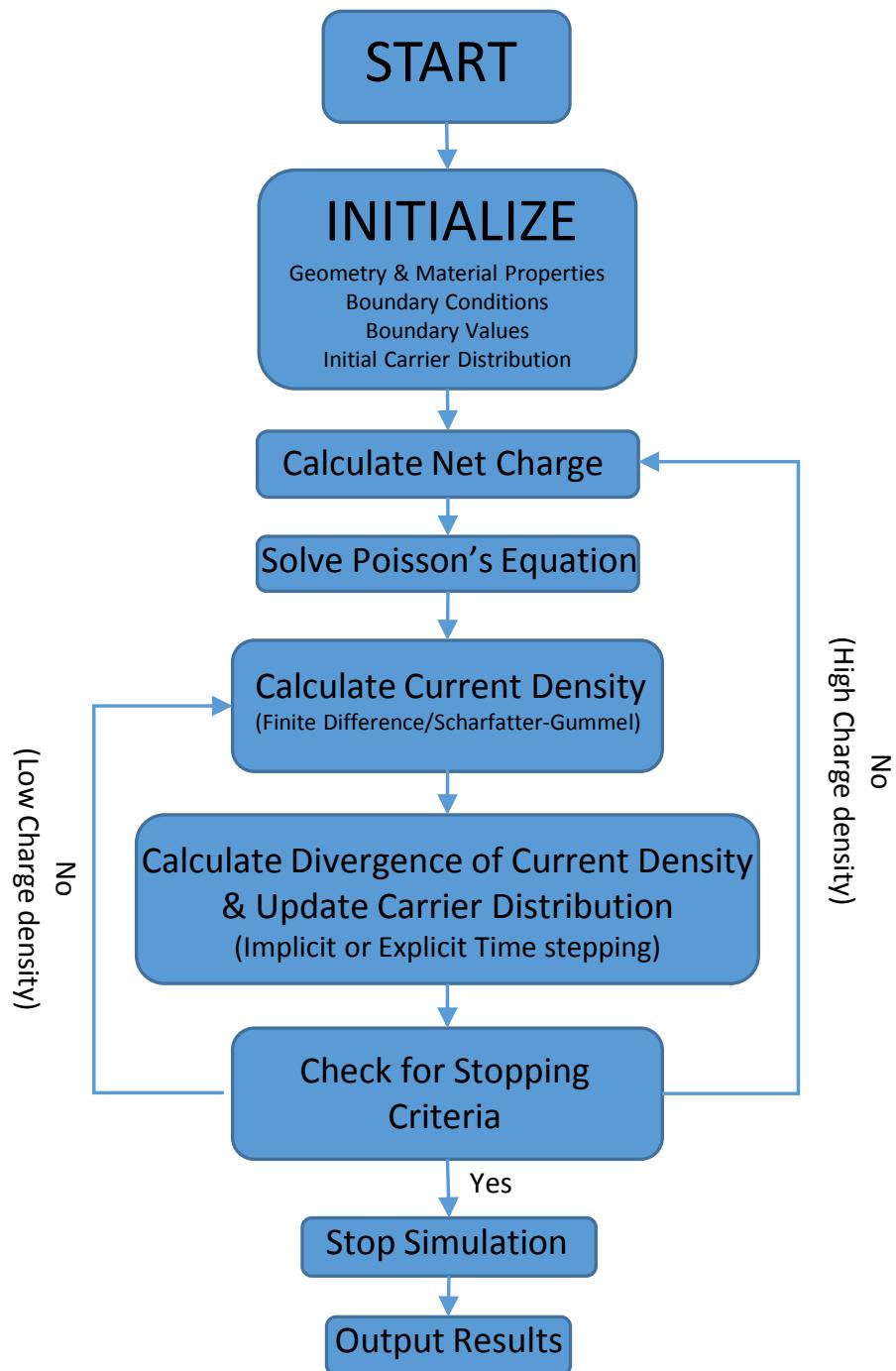


FIGURE 3.1: Finite Difference Drift-Diffusion Scheme Flowchart

## Chapter 4

# Testing Drift Diffusion Solver

In the past chapter went through the details of how to solve drift-diffusion and Poisson's equation using finite difference method. In this chapter steady state and transient analysis for drift diffusion equations using finite differnce is compared to analytical solutions as well as a commercially available simulator called 'COMSOL Multiphysics' which uses finite element method instead of finite difference. Following test cases were made to ensure that the key parts of the finite difference scheme runs properly and does not produce unexpected results.

### 4.1 Solution for Closed Boundary

In this section the accuracy of the finite difference solution in steady state will be tested. In order to do this simple 1-D problem where with a finite number of negatively charged particles over a certain distance subject to constant electric field is used. This is the same problem that was solved analytically in section 3.3.1. It is also assumed that the charge density is very low and does not affect the electric field. Both ends of the simulation domain have no flow boundary conditions for charged particles. The solution process requires an initial distribution for charge density over the area. For this problem the density of the negative particles was initialized to be uniform over the entire area.

Since the differential equations are uncoupled solving Poisson's equation only once is sufficient to determine the electric field over the course of the entire simulation. Figures 4.2 and 4.1 show the potential and the electric field distribution over the entire simulation area calculated from Poisson's equation using finite difference.

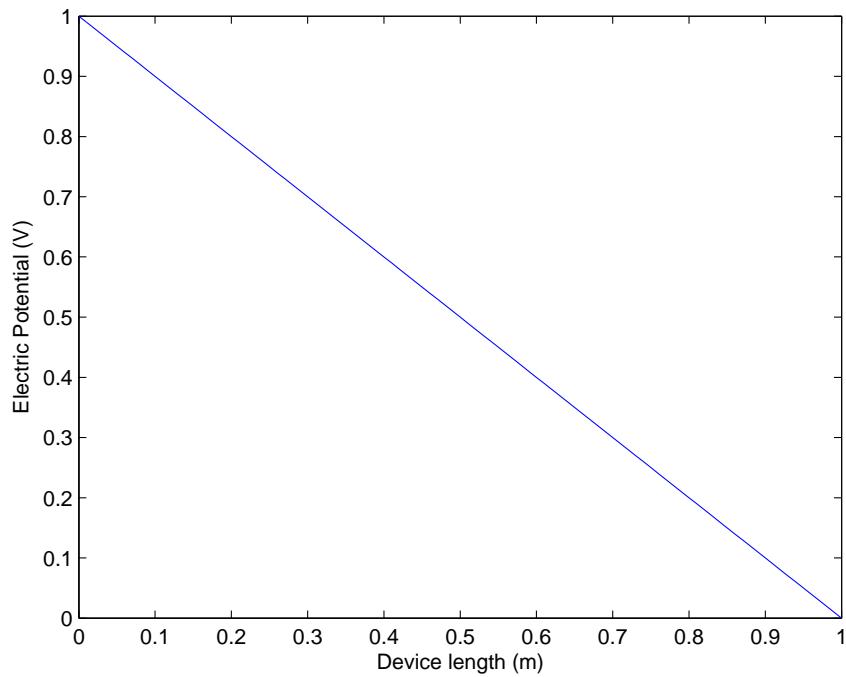


FIGURE 4.1: Potential Distribution

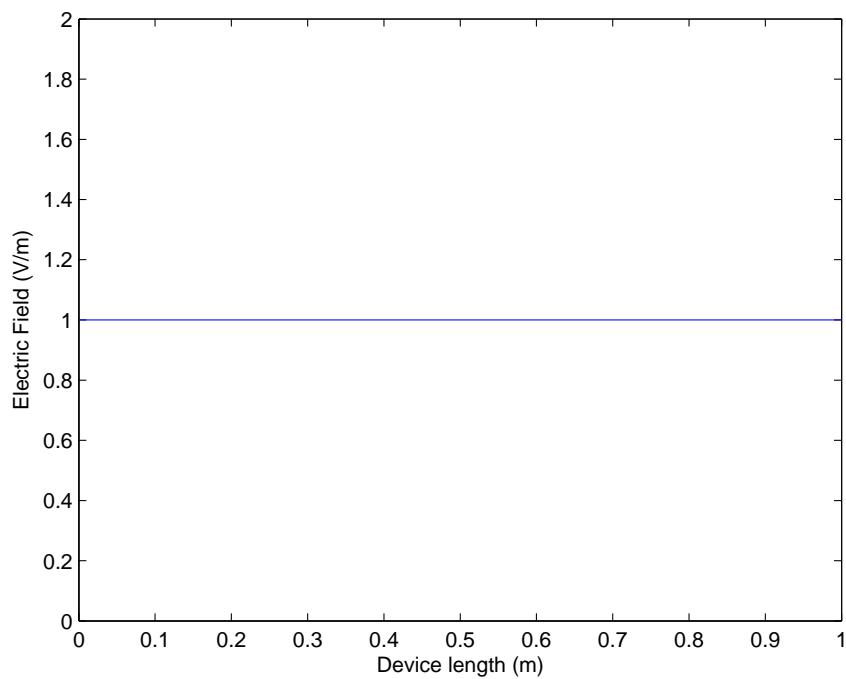


FIGURE 4.2: Electric Field Distribution

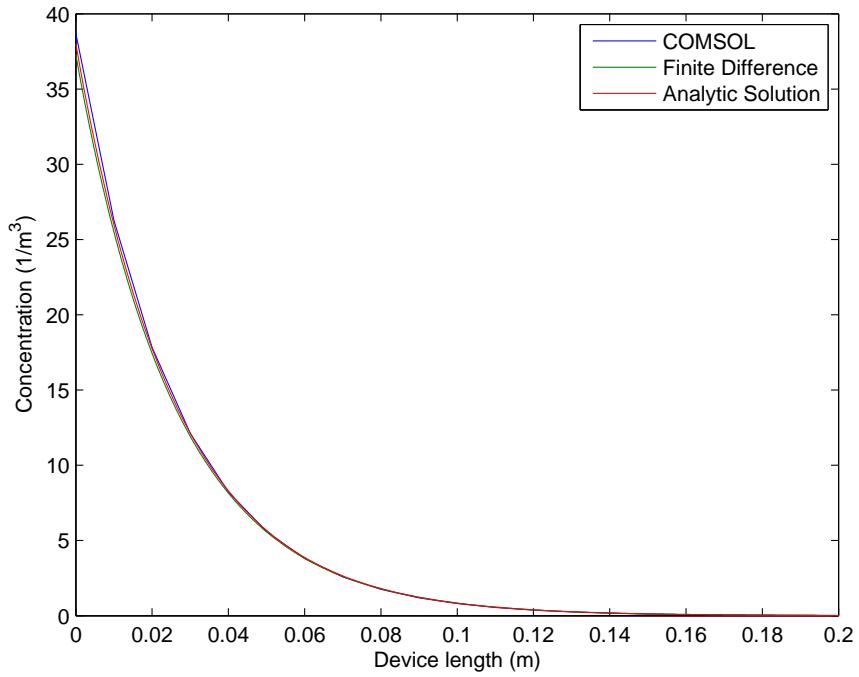


FIGURE 4.3: Steady State Negative Charge Density

Figure 4.3 has two simulation results as well as the exact solution of this problem. The green line represents the result given by the finite difference method at steady state. It can be seen from the graph that the steady state solution generated by both COMSOL and finite difference matched the analytical solution.

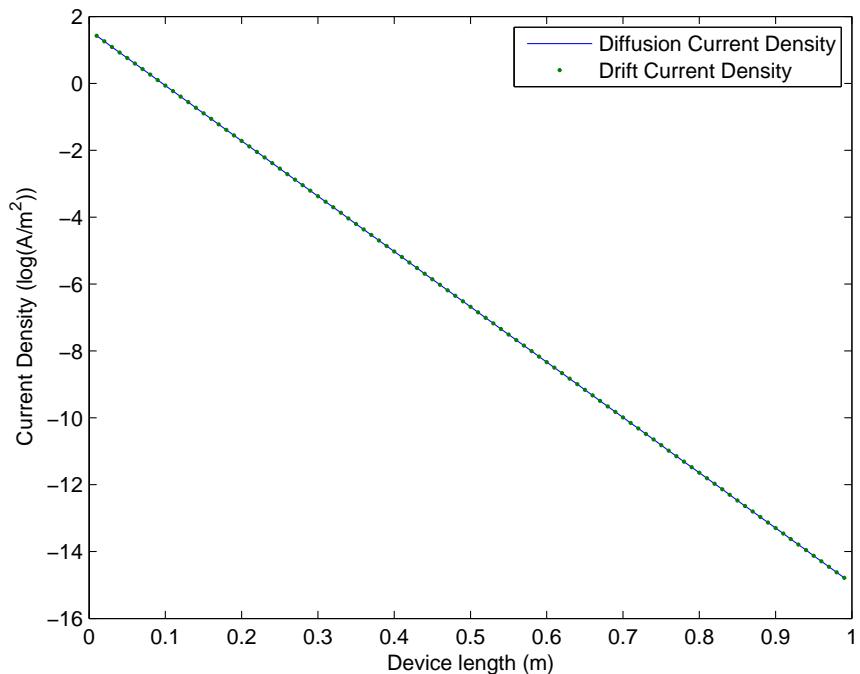


FIGURE 4.4: Finite Difference Drift and Diffusion Current Densities

While deriving an analytical solution for this problem we have assumed that at steady state the drift current density must be equal to the diffusion current density. In figure 4.4 we see drift and diffusion current densities in log scale. Overall both currents match quite tightly.

(Add problems with high electric fields and debye length)

(This example tested the simulation for, steady state  $dn/dt=0$  etc.. Generates good results for low E field but problems with high E fields)

## 4.2 Solutions for Open Boundary

Another crucial aspect of drift-diffusion simulation is its transient response. Like the previous test case, the analytic solution can be used to test the accuracy of the transient response. Two different analytic solutions for a similar drift diffusion problems which involved infinite boundaries and a uniform electric field were generated in section 3.3.2. The only difference between two cases are their initial carrier distributions. One has a rectangular and the other one has a gaussian initial carrier distribution. Finite difference scheme does not allow simulation over an infinitely long conductor. For this simulation, a solution will be generated until the carrier distribution gets close to a wall.

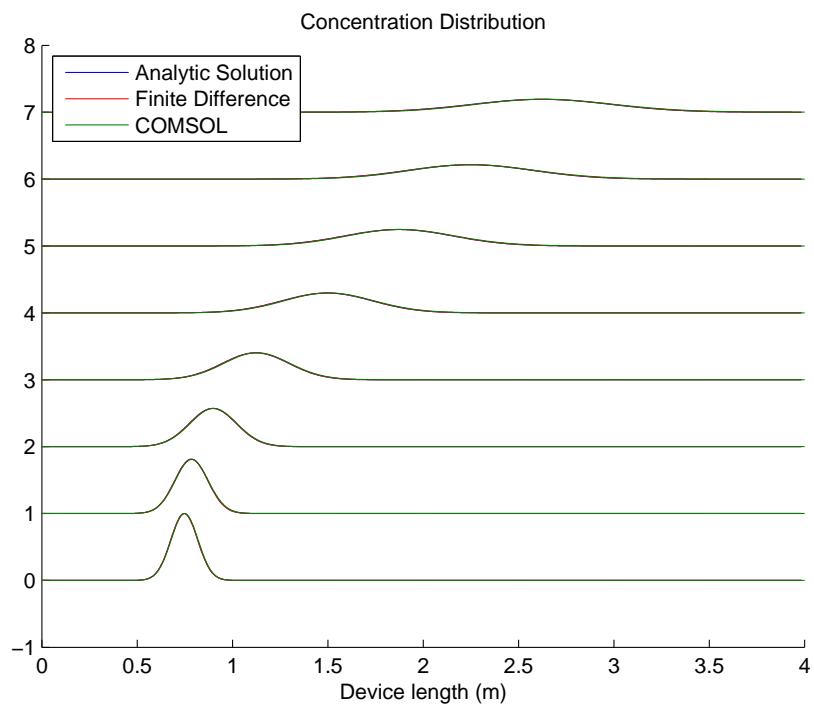


FIGURE 4.5: Gaussian Carrier Distribution Evolving Over time

Figure 4.5 has the transient response from COMSOL, finite difference and analytical solution. Snapshots of the carrier distributions were taken for each method at different time steps and they were superposed on top of each other. Increasing levels on y axis represent a carrier distributions at a different time starting from the bottom and moving forward in time towards the top. Figure 4.5 has a gaussian initial carrier distribution. It can be seen from that the transient solution generated by both COMSOL and finite difference are quite close to the analytical solution.

(Needs conclusion, What this example showed ?)

### 4.3 PN Junction

In previous test cases, Poisson's equation was not coupled with drift diffusion equations. In this example both drift diffusion and Poisson's equation will be tested to investigate how well they work when they are coupled together. A simple pn junction is quite adequate for this task since it has analytical solutions (under certain assumptions) for electric potential, electric field and carrier distribution.

For the simulation ,the initial hole and electron distributions were determined using mass action law and they were assumed to be constant at the boundaries. Keeping carrier concentrations constant at the boundaries creates a mechanism in which the charge can move in and out of the simulation domain. If the charge density at any time step is higher than the fixed density then the difference will move out of the system. If there is a lack of charge at the boundary then carriers will move in to fill in the gap. Following figure (4.6) shows the final result of bringing p and n type materials together.

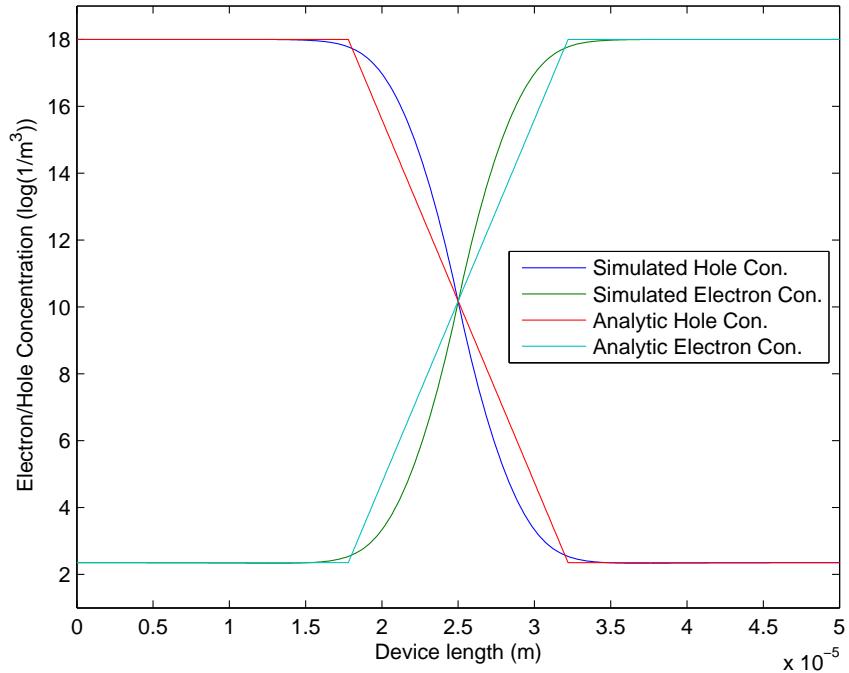


FIGURE 4.6: Electron/hole Concentration of a PN Junction

There is a little mismatch between simulated and analytic concentration densities. Analytic solution have sharp edges and simulated solution does not. This is due to all the assumptions made in order to find an analytic solution. This mismatch can also be seen for electric field, electric potential and net charge .

Figure 4.7 shows the potential distribution for simulated and analytic solution. Close match in electric potential distribution shows that coupled equations can generate fairly accurate solutions.

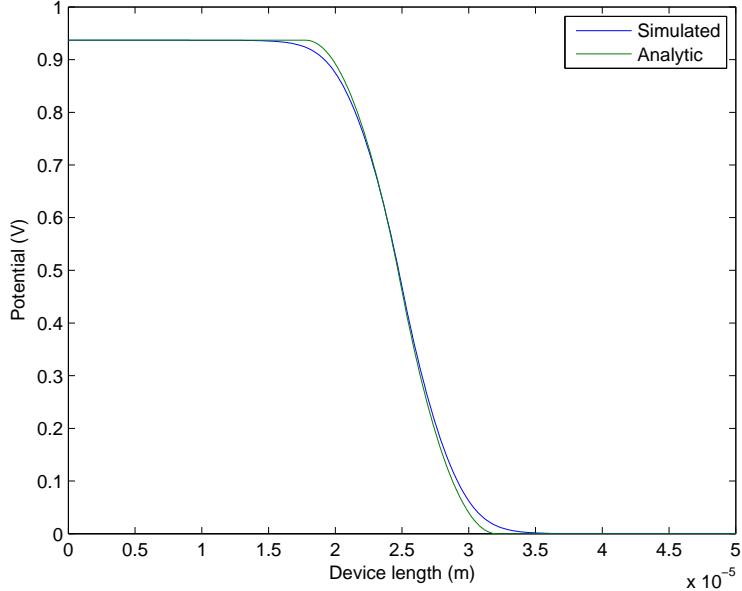


FIGURE 4.7: Potential Distribution of a PN Junction

Calculation of the electric field involves one basic derivative. Since simulated potential is matching the analytic solution quite nicely the electric field should follow a similar pattern. Figure 4.8 shows that this is indeed the case, simulated electric field matches the calculated electric field.

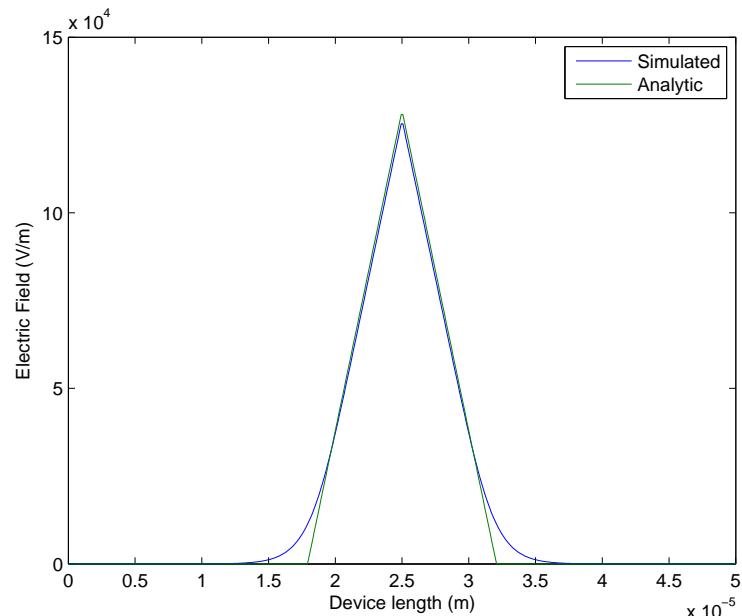


FIGURE 4.8: Electric Field Distribution of a PN Junction

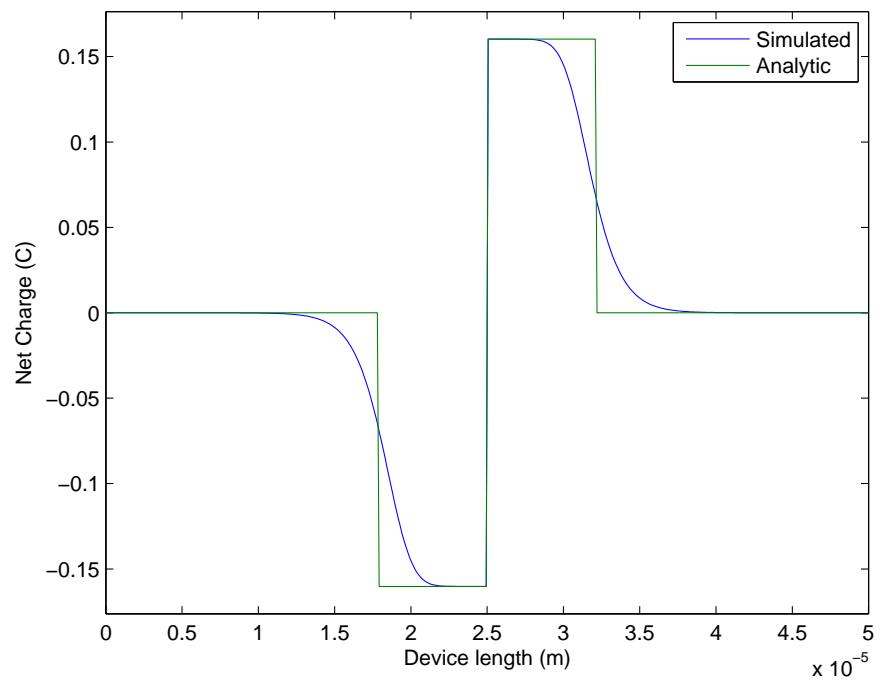


FIGURE 4.9: Total Charge Distribution of a PN Junction

Figure 4.9 shows the total charge distribution at steady state. Simulated net charge density follows the analytic one except the abrupt changes at two ends.

(Needs conclusion, What this example showed ?)

## 4.4 Region Specific Particle Density Limit

The last property of the drift diffusion simulation scheme to be tested concerns the movement of lithium ions. As lithium ions move into the PEDOT:PSS they bond with PSS polymers and replace holes. PEDOT:PSS can only absorb lithium as long as there are available PSS polymers to bond with therefore PEDOT:PSS has limited capacity to accept lithium ions. This behavior can be captured during simulation by using a Dirichlet boundary condition ( $J=0$ ) to stop the particle flow into a node if the density will go over a preset limit. This can be achieved in two different ways.

A soft limit can be set by making particle mobility a function of its density. This function can be defined such a way that it switches from high to a low value when the particle density approaches its limit. This implementation is straightforward and can be used in commercial simulators but it has a drawback. Once the particle density is reached the mobility and diffusivity of the lithium particles are stuck at very low values. If there is an outflux of particles from that particular node then the density is stuck at the limit until the mobility function goes back to a value which allows particle flow. This can introduce a considerable lag in the response of lithium current density.

An alternative approach is precalculating the particle density of a density limited node at the next time step, setting any influx to zero if the density is going to go over the limit and finally recalculating the particle densities at the next time step using the updated current densities. This mechanism sets a hard limit on the density since there is a sudden cut instead of a gradual cut in the current flow. This density limiting mechanism is more responsive than the previous one but it requires the calculation of the next time step twice for the species that has a limit. Also large currents caused by either a big time step or high electric field can force the algorithm to cut the current flow into a node before the density reaches its limit.

To avoid adding any lag into the system latter method was chosen for memristor simulations in this thesis. This method was implemented in finite difference drift diffusion solver and it is tested in this chapter. Also a soft limit was implemented in COMSOL for comparison since it did not support the hard limit method.

In order to test density limiting method implemented for memristor simulation a simple example was set up. Two positively charged particles were initialized like the figure below (4.10). One set of particles has a concentration limit of  $2 \cdot 10^{10} m^{-3}$  on the right side of the simulation area and no limit on the left side. The other set has no restrictions.

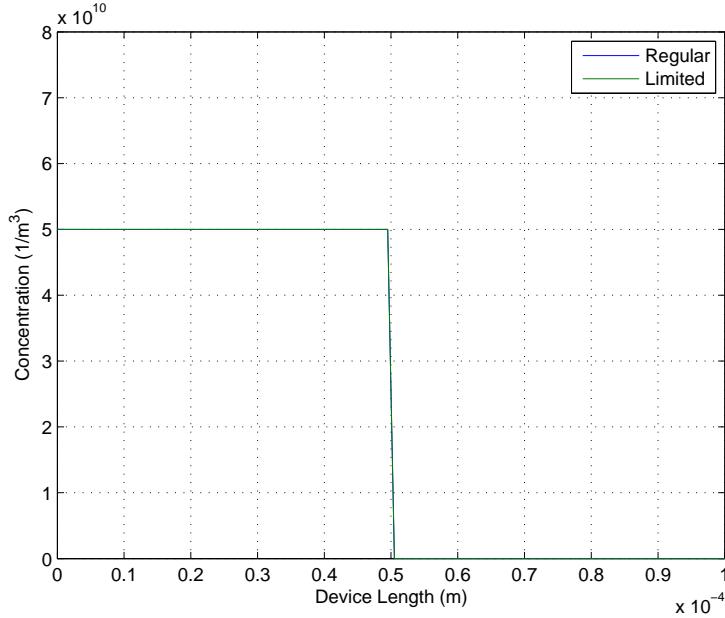


FIGURE 4.10: Initial Particle Density

This transient simulation was done using a potential train. The potential is applied to the left side of the device and the right side is always grounded. For the first half of the simulation applied potential was positive and for the second half it was negative. Figure 4.10 shows how two simulations differ when the particles are pushed towards the right wall due to the electric field created by the positive potential. Particles with no limit on the right side move freely and accumulate on the right side. The limit for the other particles is effectively stopping them from going in and accumulating freely. Once the limit is reached at a certain node the density cannot increase any further and that node turns into a no flow wall.

When the potential is switched all the particles accumulate freely to the left side of the simulation domain since there are no restrictions on this side (figure 4.12).

This last figure (4.13) shows the transient response over time of a single node on the right side. The potential is positive for the first 1.5 seconds and it is switched to negative for the last 1.5 seconds. The node without limit keeps accepting charge until steady state has been reached but the node with a limit on stops accepting charged particles once the limit is reached. After the potential is switched density limited node has no problem releasing the particles.

Additional to the test that was run for finite difference, COMSOL was used to test the soft limiting mechanism. COMSOL does not have a built in option that allows limiting particle density. One possible solution to this is making particle mobility and diffusivity a function particle density. It is possible to use a sigmoid function which switches from

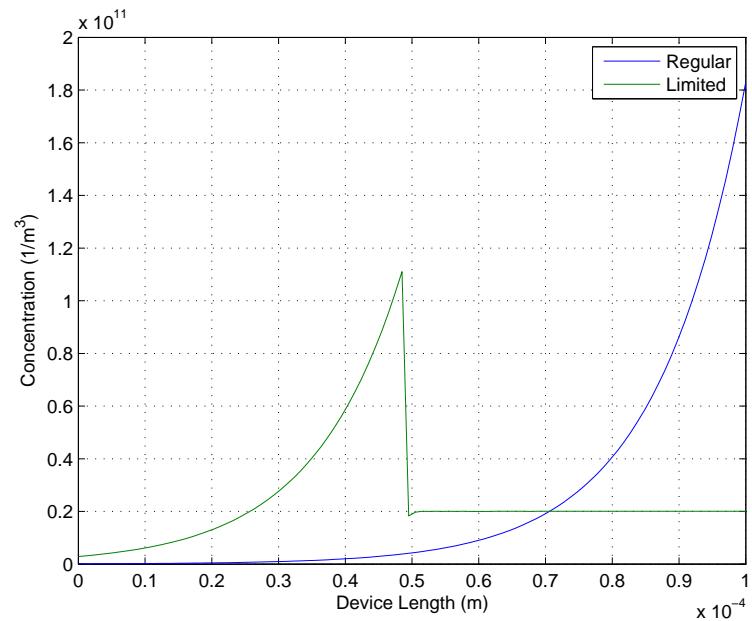


FIGURE 4.11: Limited Concentration Accumulation on the Right Side

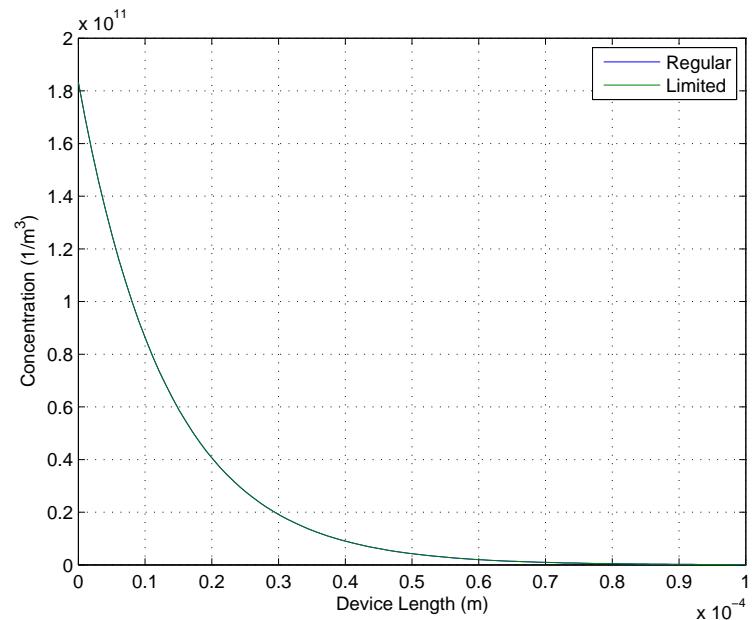


FIGURE 4.12: Limited Concentration Accumulation on the Left Side

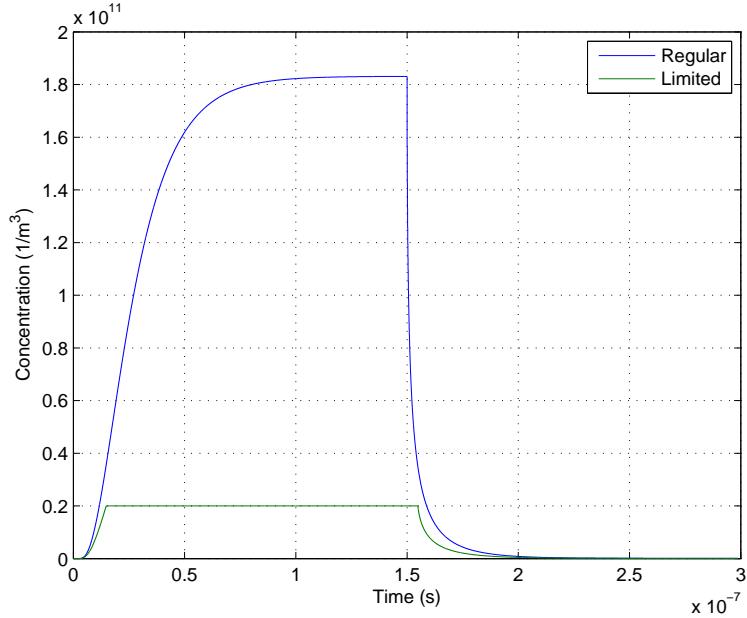


FIGURE 4.13: Accumulation at the right wall over time

1 to 0 very quickly when particle density is close to its limit. Here is the equation of the sigmoid function used to limit the particle flow:

$$\mu = \frac{\mu_0}{1 + e^{\sigma(n-n')}} \quad (4.1)$$

$\mu_0$  is the original mobility of the charge carrier.  $\sigma$  controls the sharpness of the switch and  $n'$  determines the density at which the switch will be.

Another problem with COMSOL was the definition of mobility and diffusion coefficients over different areas. If these constants are defined as one single value per area COMSOL simulates without any errors but if they are defined as a function of concentration it causes convergence issues. This can be overcome by using two more sigmoid functions to distinguish between two areas with different mobilities and diffusion constants. Left sigmoid in figure 4.14 was multiplied by the mobility/diffusivity of the left side of the area and the right sigmoid was multiplied by the mobility/diffusivity of the right side of the area. Both functions were summed to obtain a function which describes the characteristics of the entire area.

$$\mu = \frac{\mu_l}{1 + e^{\sigma_x(x-x')}} + \frac{\mu_r}{1 + e^{-\sigma_x(x-x')}} \quad (4.2)$$

In this problem  $\mu_r$  is the same as equation 4.1 since the region on the right side has a particle density limit. A sharp switch between right and left side mobility using the

function above would be ideal for this simulation but unfortunately it causes convergence issues in COMSOL therefore there is a gradual change between two mobilities.

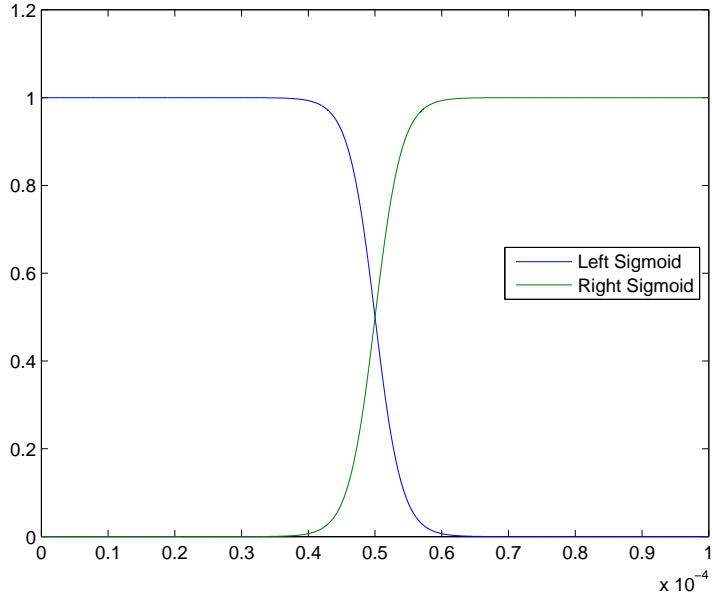


FIGURE 4.14: Mobility change from left to right side

The initial carrier distribution for COMSOL was set to be exactly the same as finite difference simulation. Figures 4.15 and 4.16 show results for both COMSOL and finite difference simulations at steady state before the potential switch. The plot on the left side gives insight on how COMSOL simulation behaves for limited and limitless accumulation on the right wall. Due to the gradual change of mobility and diffusion constants between two areas we end up with concentration on the right side higher than the limit which is  $2 \cdot 10^{10}$ . Additionally, the particle density goes over its limit near the right wall. In figure 4.16 the difference between COMSOL and finite difference becomes more visible. In FD simulation the accumulation goes much higher due to higher electric field and unlike COMSOL it does not penetrate the right half of the simulation domain.

In figure 4.17 it is possible to see the accumulation of charge near the middle after the potential is switched. This is due to mobility being a function of distance and concentration. As the ions move from left to right they go from a low mobility region to a high mobility region and they slowly accumulate around the area where the change in mobility occurs. Figure comparing both COMSOL and finite difference shows that the accumulation does not happen in the case of finite difference due to the way concentration limiting mechanism was implemented.

With the decrease of ion concentration on the limited region the difference between low and high mobility regions diminish. Once the concentration on the limited side is low enough the whole system behaves as if there was no limit and ion mobility becomes

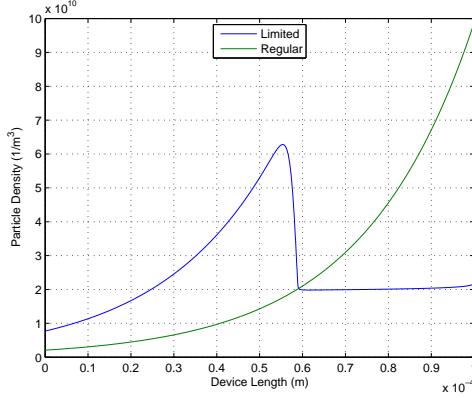


FIGURE 4.15: COMSOL Simulation for Particle Density Limit

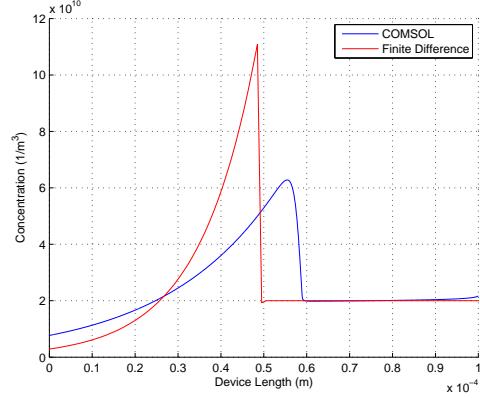


FIGURE 4.16: COMSOL and Finite Difference Simulation

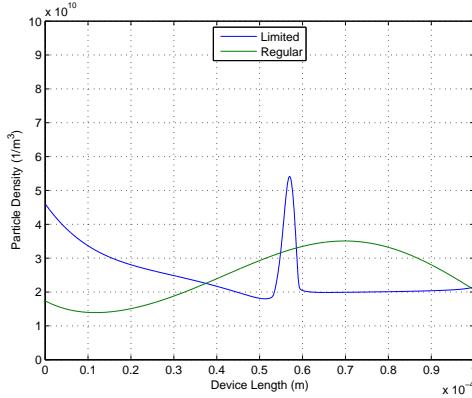


FIGURE 4.17: COMSOL Simulation for Particle Density Limit

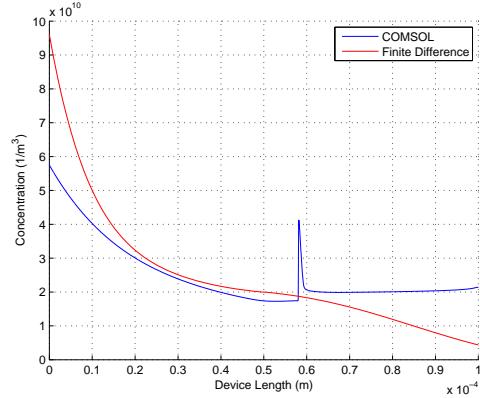


FIGURE 4.18: COMSOL and Finite Difference Simulation

equal for all regions and the ions freely accumulate on the left wall (figure 4.19). Aside from the difference in electric field strength both simulations behave the same way as they approach steady state. Figure 4.20 shows concentration densities at steady state. COMSOL simulation has a lower electric field since it has convergence issues with high electric fields.

This comparison can be finalized by looking the particle density transient response of the rightmost node. For the first half of the simulation everything is the same as the finite difference case except COMSOL goes a little bit over the limit. When the potential is switched, node without the limit has no noticeable difference in behaviour. The node with concentration limit has a lag when it comes to releasing the particles. This is due to the sigmoid function used to achieve a limiting behaviour. Once the limit is reached mobility and diffusivity are stuck at a very low value until the particle density starts to go lower than the limit.

The simulations above showed that is possible to impose a density limit over any area

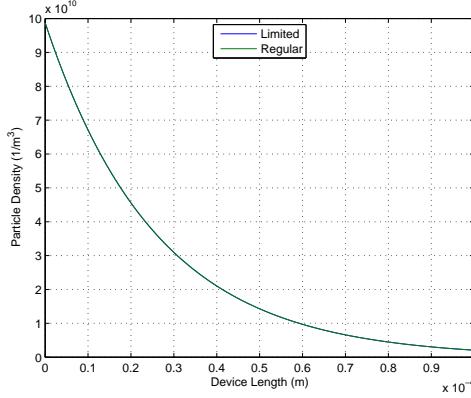


FIGURE 4.19: COMSOL Simulation for Particle Density Limit

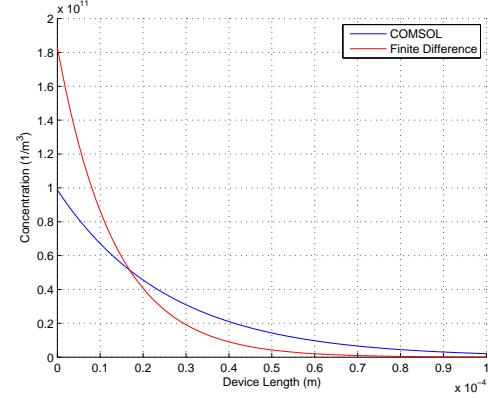


FIGURE 4.20: COMSOL and Finite Difference Simulation

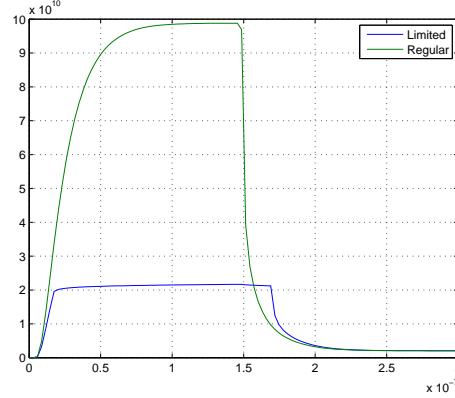


FIGURE 4.21: Density on the right wall over time using COMSOL

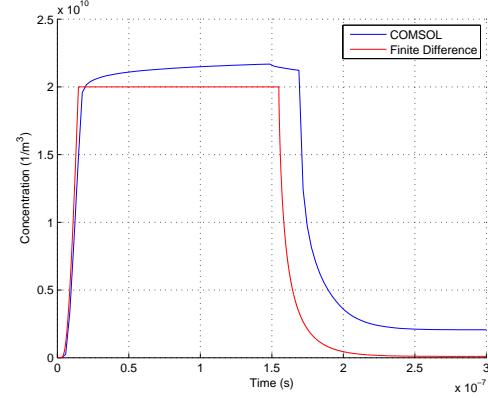


FIGURE 4.22: Density on the right wall over time, COMSOL vs. Finite Difference

using a simple no flow boundary condition in finite difference. For COMSOL some workarounds had to be implemented in order to simulate the same behavior which caused convergence issues.

**(Need a better conclusion)**

# Chapter 5

## Memristor Simulation

A numerical method for a memristor simulation was developed and tested in previous chapters based on drift-diffusion equations and finite difference. This chapter introduces the memristor's structure and physical parameters used for the simulation. It continues with a preliminary problem analysis to determine required mesh density and maximum possible time step. This preliminary analysis is followed by 1-D simulations of a memristor under various conditions.

### 5.1 Memristor Structure

Following figure (5.1) shows the structure of a simple memristor which will be taken as a basis for all the memristor simulations presented in this thesis. It consists of 2 metal contacts, a polymer conductor (PEDOT:PSS) and an electrolyte solution which has lithium and perchlorate ions (perchlorate/lithium density  $\approx 6.02 \cdot 10^{23} m^{-3}$ ). The memristor is about 1 cm long and 1 mm wide. The thickness of the conductive layer is around 1  $\mu m$ . During experimentation the electrolyte is deposited on PEDOT via a syringe so its thickness can vary drastically but as long as the amount of ions in the electrolyte solution is enough to saturate PEDOT this does not make a significant difference in the operation of the memristor. For simulation it was assumed that there were always more than enough ions to saturate the PEDOT so the electrolyte was modeled as an infinite source/sink of ions. The top boundary of the electrolyte was assumed to be charge neutral at all times which provides a mechanism for moving ions in and out of the system. This way the movement of ions near the surface of the PEDOT can still be captured without having to simulate the ion movement for the entire electrolyte solution which is variable in size.

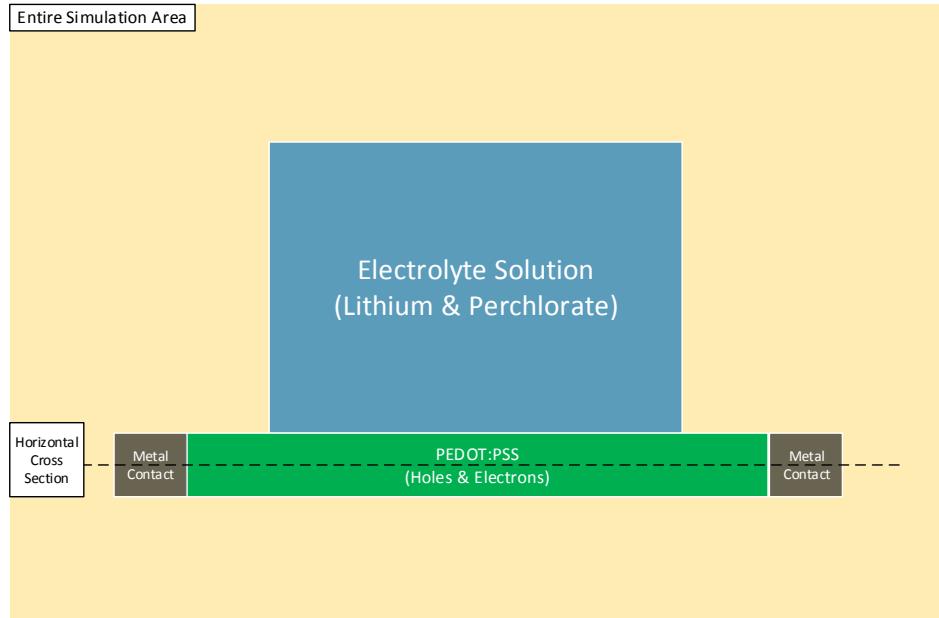


FIGURE 5.1: The structure of the memristor used for simulation (not to scale)

The initial conditions for all the charge carriers are the same. All charges are balanced and uniformly distributed but the carrier density in the electrolyte solution is higher than the carrier density in PEDOT. Perchlorate ions are not allowed to move outside the electrolyte solution so a no flow boundary condition was used around the electrolyte. Lithium ions are free to move between PEDOT and the electrolyte solution but their maximum concentration is limited inside the PEDOT. The mobility of lithium ions has further restrictions inside PEDOT. Lithium has higher mobility in PEDOT right under the electrolyte solution, wet PEDOT, than the region without any contact with electrolyte, dry PEDOT. In fact due to this difference very little amount of lithium reaches the metal contacts. This decrease in mobility was modeled by making the mobility of lithium a function of position. The mobility of lithium ions were assumed to be 100 times slower than the mobility of holes in the wet PEDOT and it is zero in the dry PEDOT ( $\mu_{hole} \approx 10^{-3} m^2/Vs$ ).

PEDOT:PSS is a regular conductor with fixed negative charge and mobile holes. Holes can move in and out of the PEDOT through the metal contacts which hold the charge neutrality of the initial condition throughout the simulation. The interface between PEDOT and electrolyte only allows the exchange of lithium ions. During simulation, the movement of lithium ions changes the conductivity of the PEDOT by increasing or decreasing the amount of available holes through coulomb forces. In the actual device lithium ions change the conductivity via various physical effects like changing the mobility of holes through modifying their hopping distance. Even though the mobility of

the holes can simply be made a function of time the physical details of these additional affects are beyond the scope of this thesis.

## 5.2 Simulation Requirements

It is important to analyze computational requirements of a simulation in order to asses the feasibility of the computation scheme. In this case, it is possible to determine spacial and temporal requirements using the equations 3.48 to 3.51 which describe physical and numerical limitations of the simulation. Following graph 5.2 shows the requirements for a memristor of the scale discussed above and a typical semiconductor device size around  $1 \mu\text{m}$ . The mesh density has to be high enough in order to capture the exponential charge accumulation for charge shielding so the minimum step size was set to be 5 times smaller than the Debye length. Plots 5.2.a and 5.2.c show the amount of points required to simulate a semiconductor and a memristor based on minimum step size. It is important to note that these values are for 1-D simulation and they can be converted to 2-D and 3-D by squaring or cubing y axis values respectively. Plots 5.2.b and 5.2.d were created using CFL conditions for drift and diffusion and dielectric relaxation time. A typical simulation time was estimated using mobility and electric field. Based on the estimated simulation time the number of time steps were calculated using the minimum time step obtained from CFL conditions and dielectric relaxation time.

It can be seen from graphs 5.2.a and 5.2.c that memristor simulations require much higher mesh densities compared to a typical semiconductor simulation such as  $1 \mu\text{m}$  long PN diode. This is due to the larger size and higher charge density of the memristor. Graph 5.2.a 5.2.b show that a memristor with  $10^{26} \text{ m}^{-3}$  charge density of the electrolyte would require close to  $10^9$  points and  $10^{14}$  time steps to simulate in 1-D. These requirements make the simulation of the memristor extremely challenging. In order investigate and find possible solutions for this issue, first a memristor low charge density ( $\approx 10^{15}$ ) was simulated in order to ensure that the simulation functions as designed. Then memristors with different charge densities were simulated and compared with each other to asses whether the behavior at low charge densities will be comparable to behavior at high charge densities.

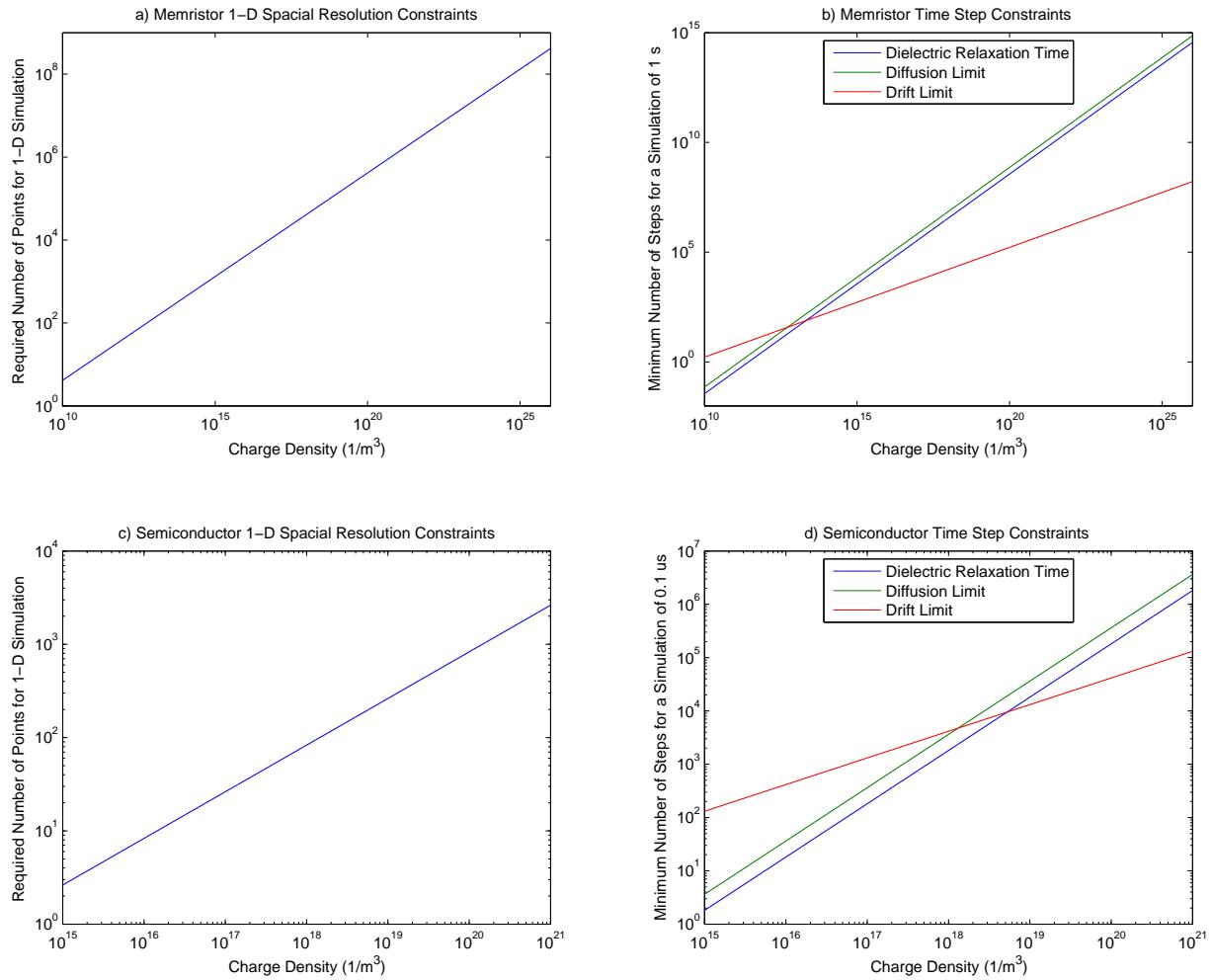


FIGURE 5.2: Spatial and temporal requirements for simulation

### 5.3 1-D Memristor Approximation

The horizontal cross section from figure 5.1 has the most crucial elements of the memristor but its simulation in 1-D is not straight forward. This cross section through the PEDOT does not include the vertical movement of lithium. Without this effect, PEDOT is just a regular conductor with a uniform current density. In order to overcome this problem a generation/recombination term for lithium ions, calculated at every time step, was added to capture the vertical movement in addition to regular drift diffusion equations which represents the horizontal movement. This generation/recombination term can be symbolized as a current source with a resistor connected to all the nodes (figure 5.3). Perchlorate ions were not included in the simulation since they do not move into the PEDOT.

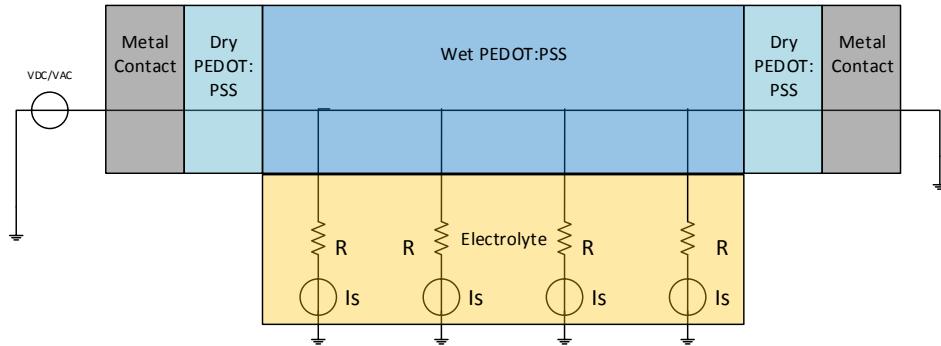


FIGURE 5.3: 1.5-D Memristor Structure

The lithium source has two different terms, one for drift and another one for diffusion. It was assumed that the concentration of lithium was always constant in the electrolyte. This way the vertical diffusion current density can be calculated using the difference between the lithium density in PEDOT and electrolyte. For the diffusion term an electric field had to be estimated between the PEDOT and the electrolyte. First the potential of the electrolyte was assumed to be half of the net applied potential. Then an electric field was calculated using the electrolyte and the instantaneous potential of the PEDOT at different positions.

Since the main characteristic of memristor is the change in resistivity over time it is important to develop a standard approach for measuring it. For the simulations in this chapter and the following one, first all the simulations were run until steady state without the movement of lithium/perchlorate ions. Lithium and perchlorate ions start to move after the steady state has been reached. The movement of ions create another transient which includes changes in resistivity. The current density (at steady state) obtained

from the initial simulation was used as a normalizing factor in order to determine the changes in resistivity after the ion movement has started.

## 5.4 1-D Memristor Simulations

(add intro)

### 5.4.1 1-D Memristor Simulation Using a Pulse Train

For the following simulation a potential pulse train, slow enough to let the memristor reach steady state, was applied on the left contact. Following plot 5.4 shows the resistivity measured using both contacts separately. As expected the resistance of the device more than doubled as the lithium ions move in. Additionally, it can be seen from the graph 5.4 the resistivity measurements from left and right contacts are not always the same over the duration of the simulation. This is normal since this is due to PEDOT layer losing holes on one side and gaining holes on the other which produces a difference in measured resistivity between contacts.

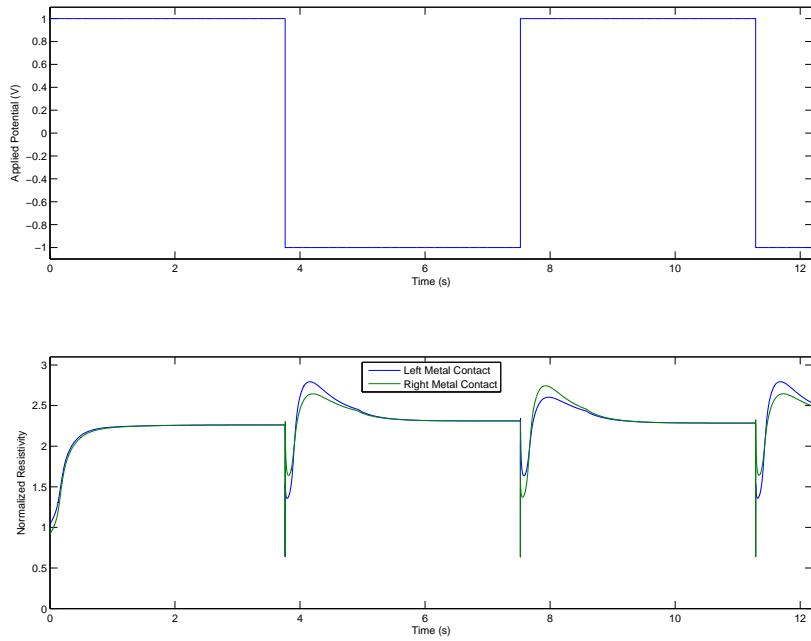


FIGURE 5.4: Change in resistivity over time due to applied potential

The resistivity in figure 5.4 shows a sudden drop when the potential is switched from 1 to 0 and vice versa. This sudden drop occurs because of the accumulation of lithium ions and holes near the negative contact which opposes the electric field generated due to the applied potential (see figure 5.5). When the potential changes suddenly, previously opposing electric field now helps the movement of holes and lithium towards the other

end of the device. This additional electric field momentarily reduces the resistivity of the device.

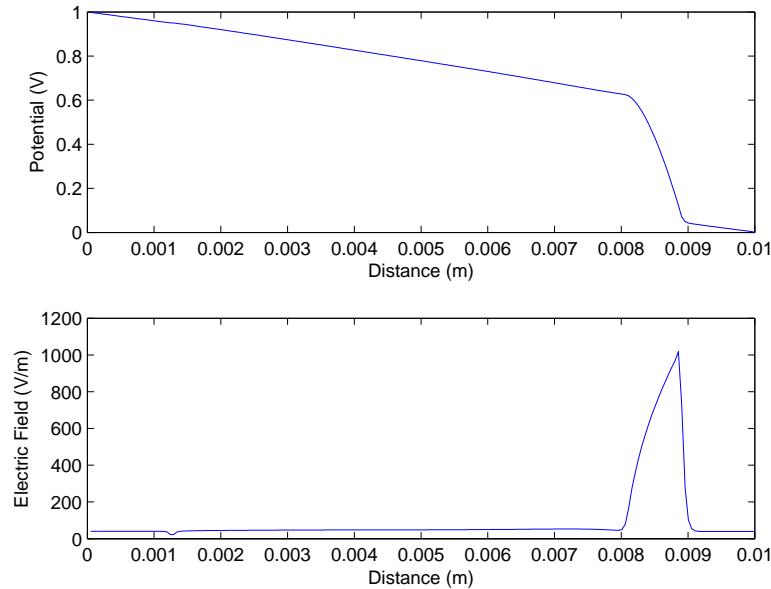


FIGURE 5.5: Potential and electric field at steady state

Figure 5.6 demonstrates the replacement of holes by lithium ions over time which directly effects resistivity seen in figure 5.7. As lithium ions get pulled in from the electrolyte toward the contact they accumulate inside PEDOT and push holes out via coulomb forces. Decreased hole concentration in the PEDOT increases the resistivity of the material. This change in resistivity over time is illustrated in figure 5.4.

It is important to note that lithium ions are free to move in x and y direction. In figure 5.6 it can be seen that after the potential is switched as the lithium density moves from side to the other. Most of the lithium movement happens through the exchange of ions between PEDOT and the electrolyte since the distance between them is far less than the length of the PEDOT. So lithium ions, traveling from the positive to the negative contact, are pulled into the electrolyte before they reach the other side. Near the negative contact lithium ions are quickly pulled into the PEDOT and accumulate at the wet/dry interface. Figure 5.7 shows the changes in resistivity throughout PEDOT due to the accumulation of lithium. The resistivity is increased in regions where there is high lithium accumulation.

By examining the resistivity plot of PEDOT it is possible to conclude that it is composed of 3 distinct regions. 2 dry regions, where there is no contact with the electrolyte, have constant uniform resistance. Between these two resistances there is a variable non uniformly distributed resistance controlled by hole/lithium concentration and hole mobility.

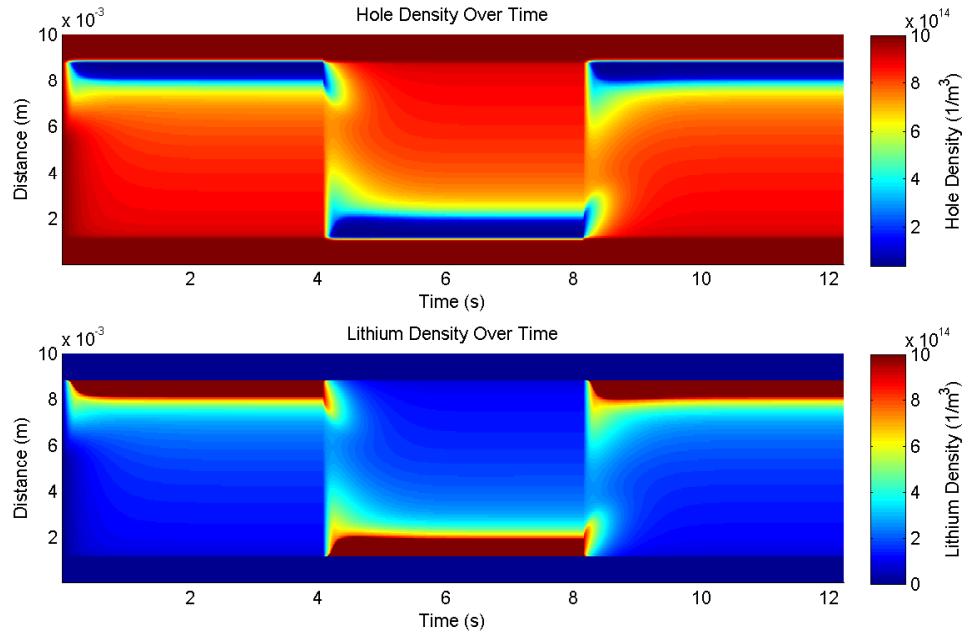


FIGURE 5.6: Lithium and hole density distribution over time

So this model captures the main characteristic of the memristor which is a variable resistance where the resistance at any time depends on the past of the device. Following equation gives the total resistance/memristance for the memristor model developed for this thesis:

$$M(q(t))_{tot} = 2R_{dry} + R(Li, p, \mu_{hole}) \quad (5.1)$$

The minimum resistance of this device is just the total resistance of the PEDOT without the lithium ions. The maximum resistance depends on different factors such as applied potential and the distribution and concentration of lithium ions inside PEDOT.

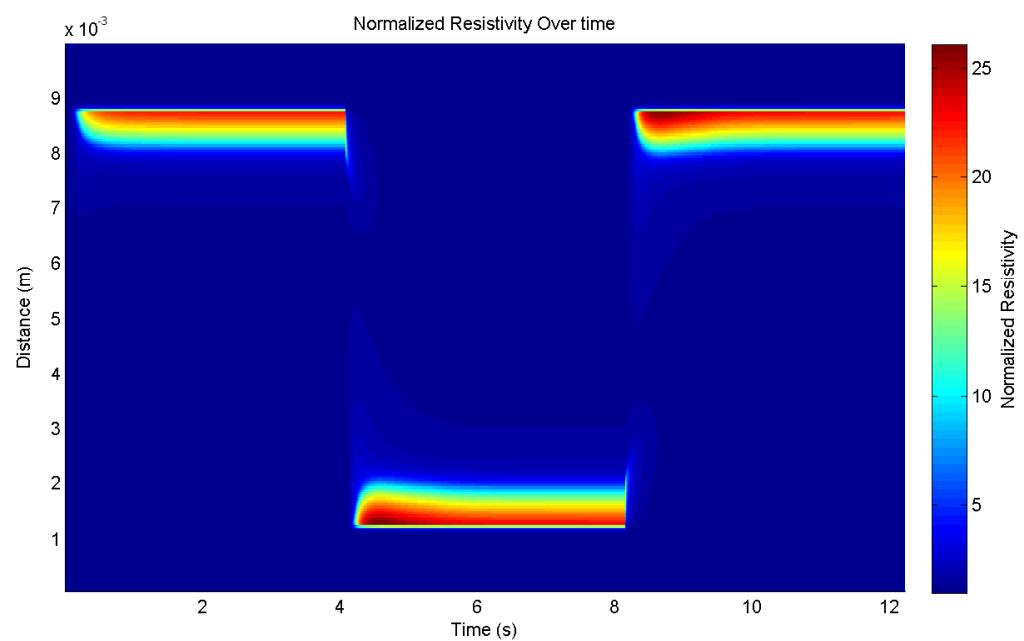


FIGURE 5.7: Normalized resistivity over time

### 5.4.2 1-D Memristor Simulation Using a Sinusoid

Memristor with a pulse train simulation showed the change in resistivity over time due to an applied potential but it does not demonstrate any memory effects. The memory effect of the memristor can be clearly demonstrated in an I-V curve using sinusoidal potential. Following four graphs (figure 5.8) were created using an AC potential with different frequencies at the contacts. All the plots show that current can have more than one value for the same potential at different times. This means that simulated memristor's past states affects its present output, therefore the device has memory. There is a pinch in the current for negative applied potential because it is measured from one contact which sees a higher resistance for a period of time during transient.

The sudden changes in current around 0 V can be attributed to the density limiting mechanism for lithium. Instead of a slowdown in the movement of lithium ions near maximum concentration, the current flow into that region is completely blocked. Even though this makes the lithium density more responsive to changes, sudden movement block combined with large time steps can result in fast changes which can be directly seen as abrupt changes in the hole/lithium current density during transient simulations.

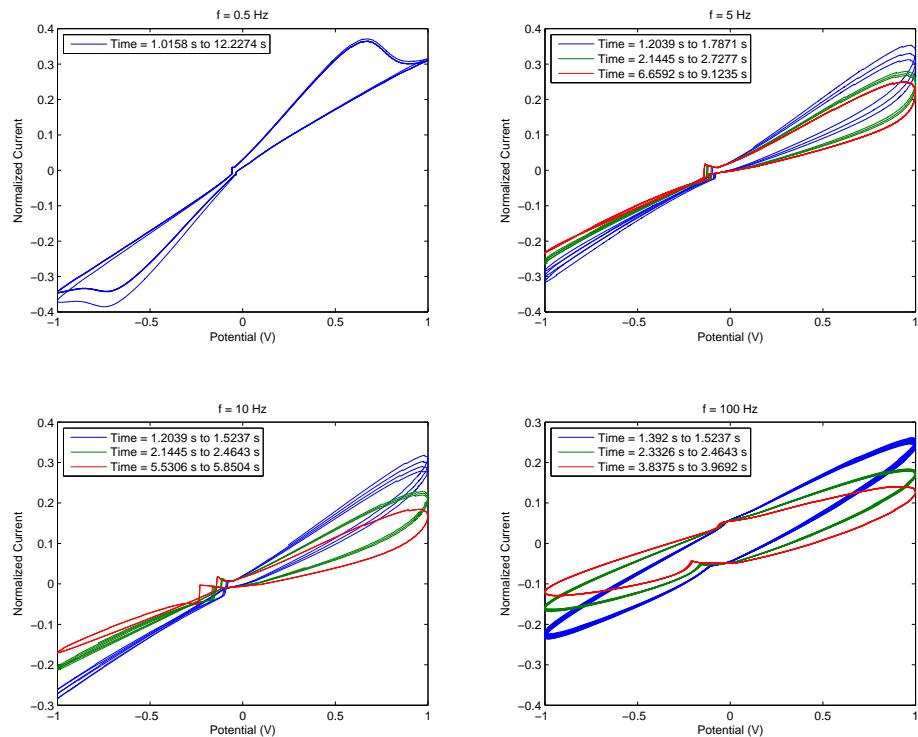


FIGURE 5.8: Normalized Current vs. applied potential at different frequencies

As AC potential pushes lithium ions from one side to the other, an equilibrium is reached over time. The changes in resistivity starts to repeat itself and after a while it becomes periodic. Figure 5.9 shows the changes in memristor current over time for the right metal contact. At first, as lithium moves into the PEDOT there is a drop in current on average. Once enough lithium settles into the PEDOT the increase in resistiviy simply shifts from left to right but it stays fairly constant on average. The final figure (5.10) show the changes in resistivity of PEDOT over time. The maximum value of the resistivity increases over time until it reaches a steady value on either side.

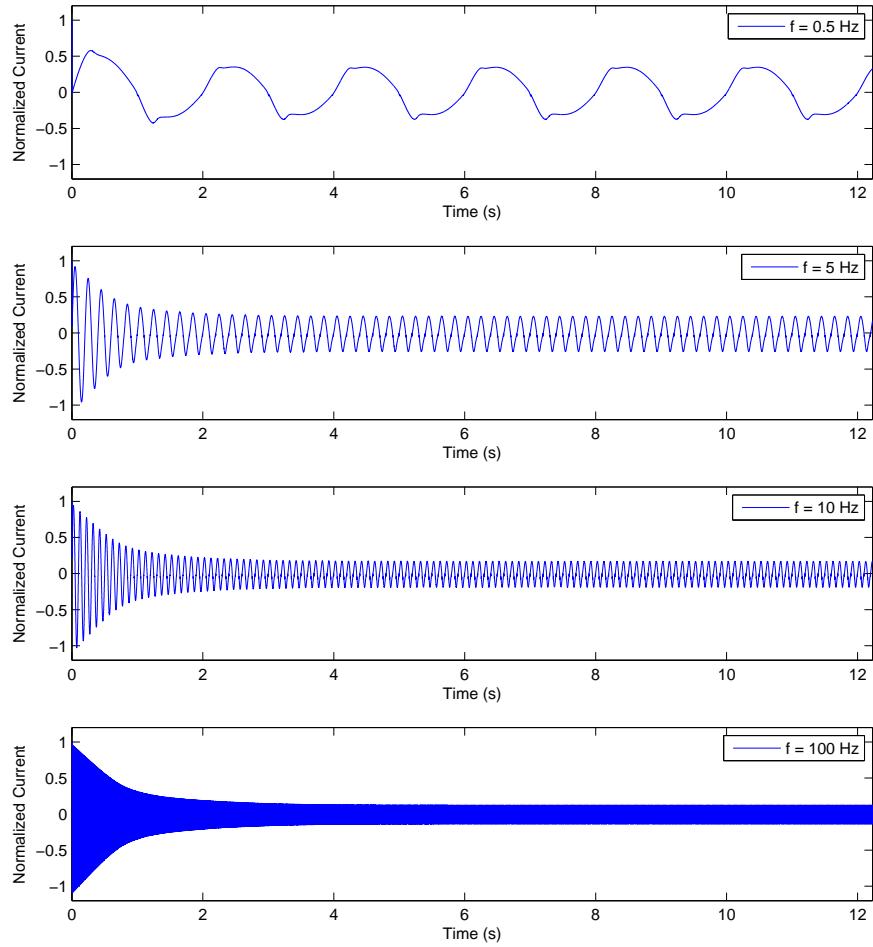


FIGURE 5.9: Normalized current over time

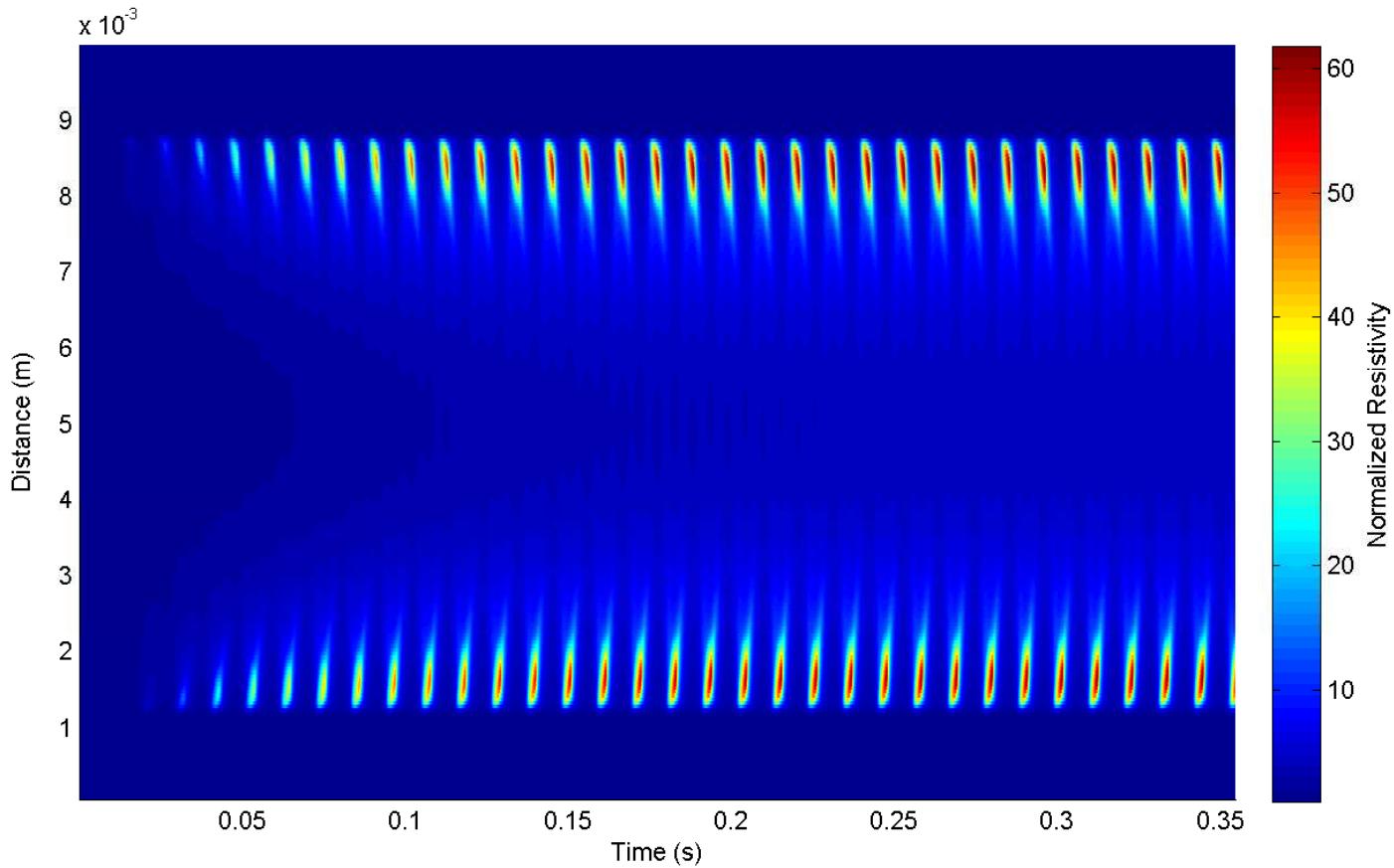


FIGURE 5.10: Normalized resistivity over time ( $f = 5$  Hz)

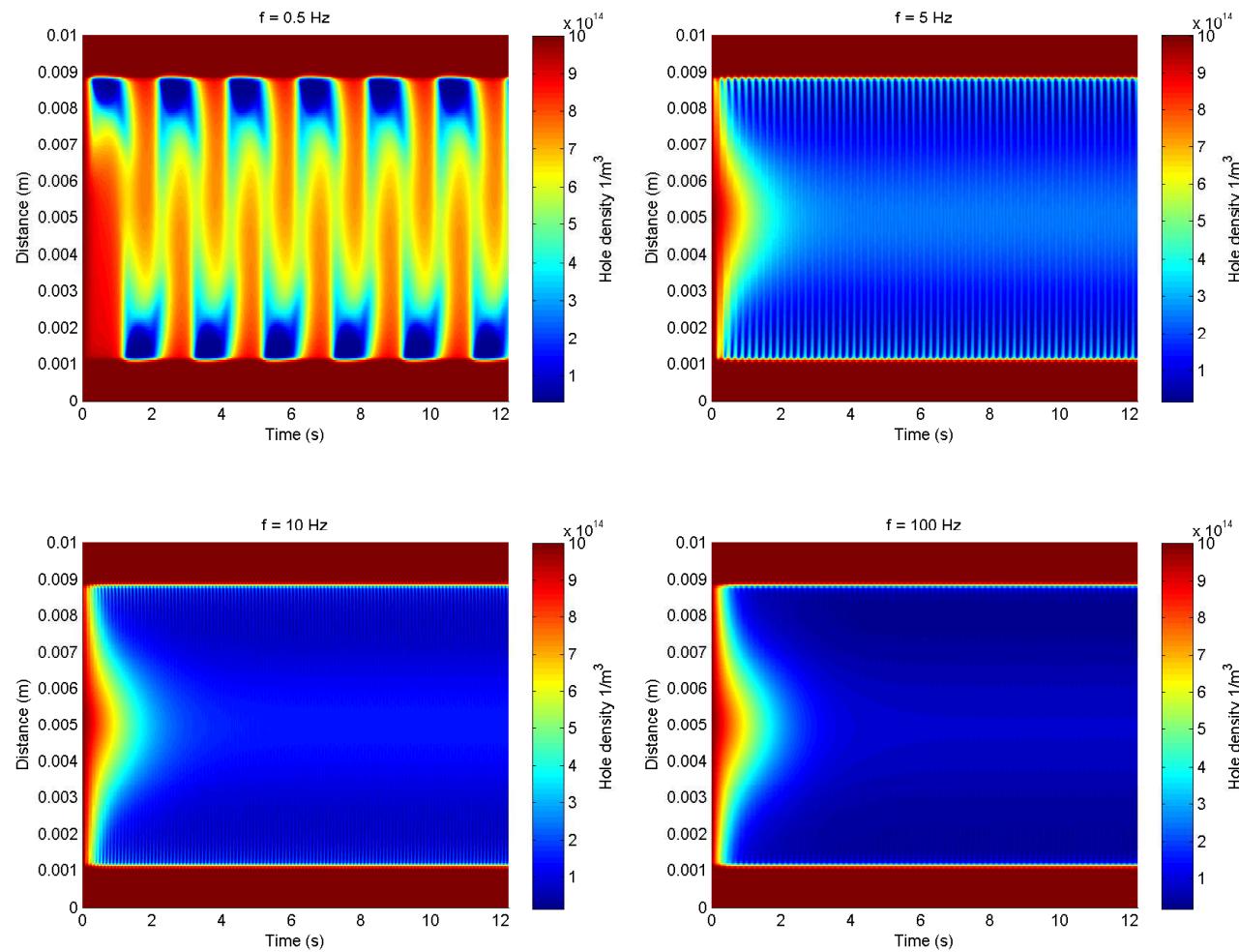


FIGURE 5.11: Lithium density over time at different frequencies

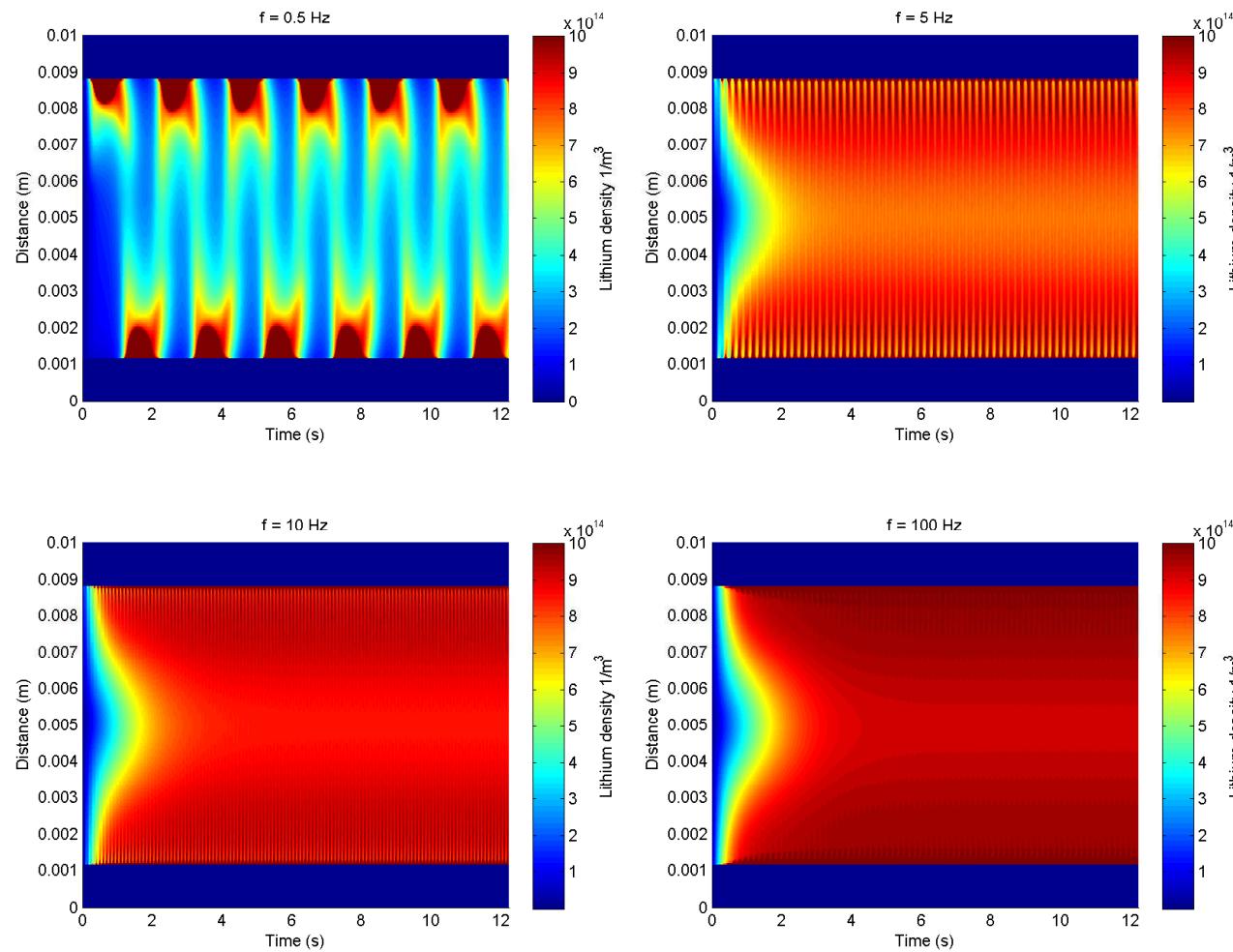


FIGURE 5.12: Hole density over time at different frequencies

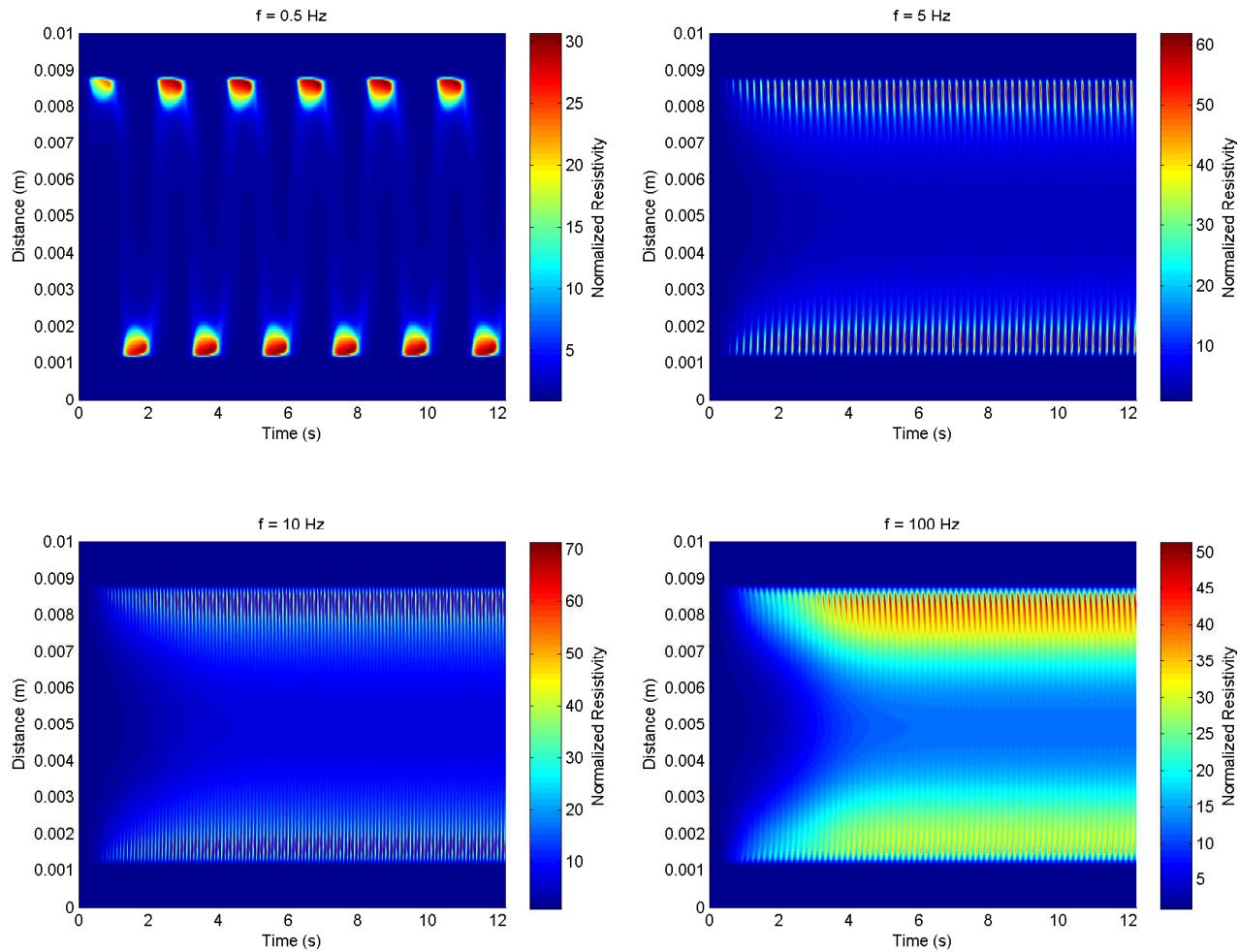


FIGURE 5.13: Normalized Resistivity at different frequencies

### 5.4.3 1-D Memristor Simulation With Increasing Charge Density

After establishing that the simulation behaves appropriately it is important to investigate how charge density effects this model since the size of the device limits the maximum charge density that can be simulated. The actual device has a hole density in the range of  $\approx 10^{22} \text{ m}^{-3}$  but the simulation is restricted to  $\approx 10^{15} \text{ m}^{-3}$ . In theory, if the densities used for simulation does have a limited affect on the behavior of the memristor then simulations in low density can be used as a proxy for the actual device which operates in much higher carrier densities.

Following 3 graphs (5.13, 5.14 and 5.15) show hole and lithium densities at steady state as well as the current density over time at the right metal contact for various charge densities. The simulations were done on high mesh densities to allow stable simulation at high charge concentrations. 20 different simulations were made using densities ranging from  $5 \times 10^{15} \text{ m}^{-3}$  to  $1 \times 10^{17} \text{ m}^{-3}$  but only 5 of them were plotted for illustration purposes. All the values were normalized to the initial plot using respective hole/lithium density ratios. Following equations were used for the normalization procedure.

$$n_{norm} = \frac{n_i}{r} \quad (5.2)$$

$$J_{norm} = \frac{J_i}{r} \quad (5.3)$$

$$r = \frac{n_{initial}}{n_0} \quad (5.4)$$

$n_i$  is the charge carrier density which could be either holes or lithium ions.  $J$  is the current density calculated from the left metal contact.  $r$  is the normalization constant which is calculated using initial carrier density ( $n_{initial}$ ) of a simulation to be normalized and  $n_0$  which is the carrier density of the initial plot ( $5 \times 10^{15} \text{ m}^{-3}$ ). After the normalization of various variables, all the plots were placed in the same graph to visually illustrate how increasing charge density affects the simulation.

Plots for lithium ions (figure 5.14) and holes (figure 5.13) are nearly identical to each other in terms of the plot shapes. They both tend to get sharper around the area where there is a substantial accumulation of lithium. This is not unexpected since the Debye length gets smaller with increasing carrier density concentration. So with increased density it is expected that the density accumulation at the wet/dry PEDOT interface will be more narrow. This effect is not as strong for lithium ions since there is a maximum concentration limit on them. Lithium and hole plots only show the difference carrier

density makes on steady state. The affect of debye length on device operation can be further explored by looking at the current density over time.

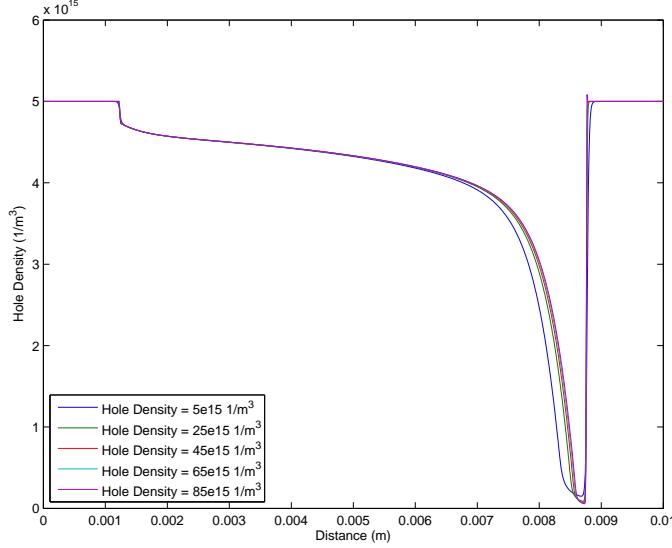


FIGURE 5.14

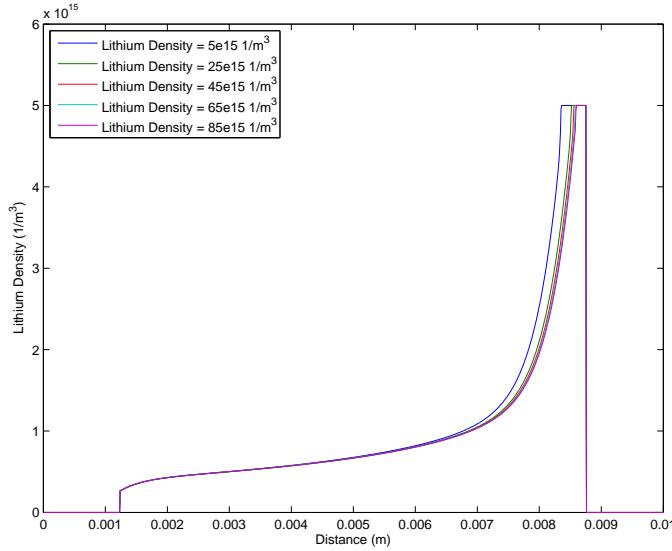


FIGURE 5.15

Like the simulation in section 5.2.2, first few miliseconds of this simulation was run without the movement of lithium ions. At this stage normalized currents are identical to each other which means a simulation with low density can be scaled up to any density without introducing any error. The introduction of lithium ions slowly deviates the current plots from each other over time but the overall behavior of the memristor remains unchanged. The current density decreases over time and reaches a steady state value.

After the first plot which has the lowest hole density, consecutive plots get closer and they are almost impossible to distinguish from each other at steady state.

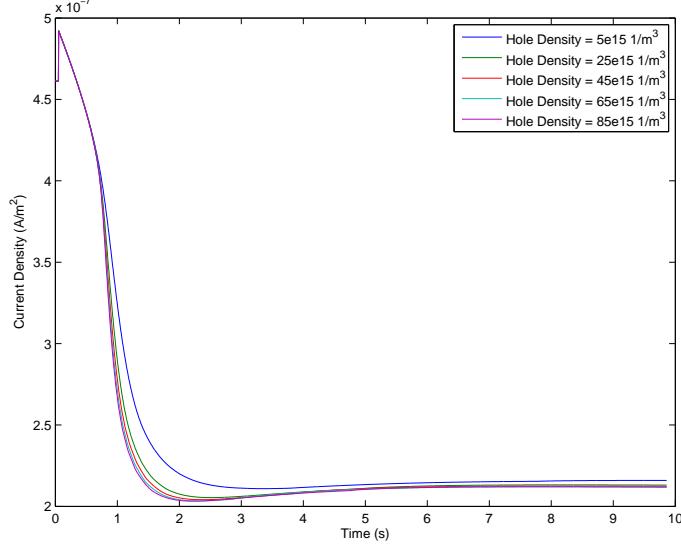


FIGURE 5.16

The impact of increasing charge density can be numerically explored using normalized plots. Following formula was used to calculate a percentage difference between different normalized carrier densities as well as normalized current densities.

$$P_i = \frac{1}{K} \sum_{k=1}^K |100 \frac{n_i(x_k)/r - n_0(x_k)}{n_0(x_k)}| \quad (5.5)$$

$$P_i = \frac{1}{T} \sum_{k=1}^T |100 \frac{J_i(t_k)/r - J_0(t_k)}{J_0(t)}| \quad (5.6)$$

$K$  is for total number of points along x axis and  $T$  is the total number of time steps taken by the simulator.  $x_k$  and  $t_k$  are used to for a point in x axis and a point in time respectively.  $r$  is the normalization constant described in equation 5.4. Both equations start by taking a point in either space (lithium/hole density) or time (current density) for a simulation with a certain carrier density and normalize them. After that point is normalized its difference from the lowest carrier density simulation is calculated as a percentage. Finally all the differences are averaged and used as a measure of the difference between simulations with different charge densities. They can also be used as an error factor for scaling.(Is this paragraph clear enough ?)

Figure 5.16 shows the difference hole/lithium density makes in the simulation. Carrier densities in the dry regions of the PEODT change very little between consecutive simulations since lithium ions do not reach there. For this reason only the densities at the wet PEDOT were used to measure the difference between plots.

In all 3 cases, there is an increase in average difference for increasing carrier density. Hole density shows the most difference between plots. Since holes are free to move inside the PEDOT they directly get effected by the change in debye length due to the increase in hole density. This is not the case for lithium ions because their movement is restricted by the limit on lithium density. In contrast, current density is not as sensitive to density changes as lithium and hole distributions.

An approximate error estimation can be made for current density if a low hole/lithium density is used for simulation and then scaled up to match the actual memristor. Figure 5.16 shows around 3.3 % difference for 2 orders of magnitude increase in hole density. Assuming the error scales linearly after that point a total of 6 orders of magnitude will result in 9.9 % error introduced in order to linearly scale from  $10^{15}$  to  $10^{21}$ . The actual error is likely to be less since the error progresses sub linearly. Following the same procedure 6 orders of magnitude difference produces a 12 % error for lithium distribution and 48 % error for hole distribution. The difference in hole distribution looks very large but the overall shape of the plot is still reserved, most change occurs around the wet/dry PEDOT interface where lithium accumulates. Based on these error margins and previous plots it can be concluded that the behavior of the memristor simulated using low charge density is very similar to the simulation at high carrier density.

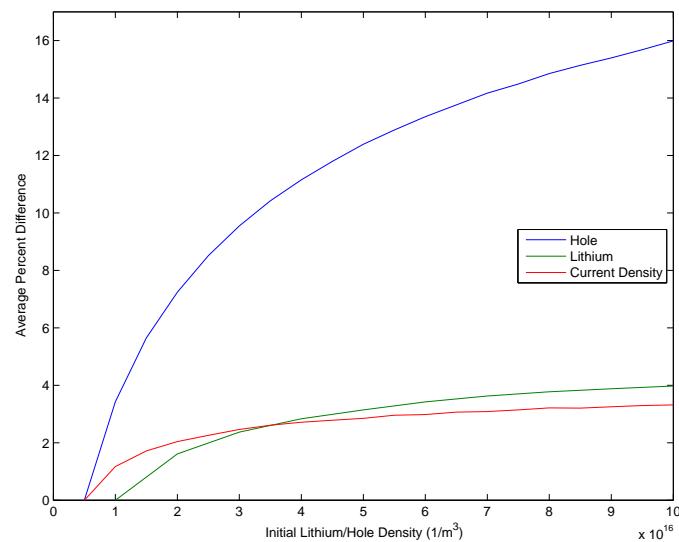


FIGURE 5.17

#### 5.4.4 Experiment vs. Simulation

Hopping distance change due to lithium modifies hole mobility. Can be used to match maximum resistivity.

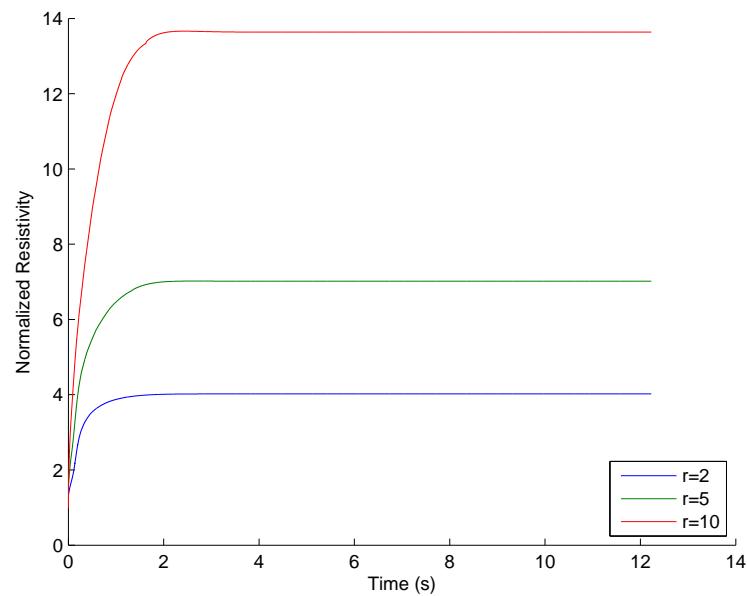


FIGURE 5.18

Lithium mobility can be scaled to match the experimental values

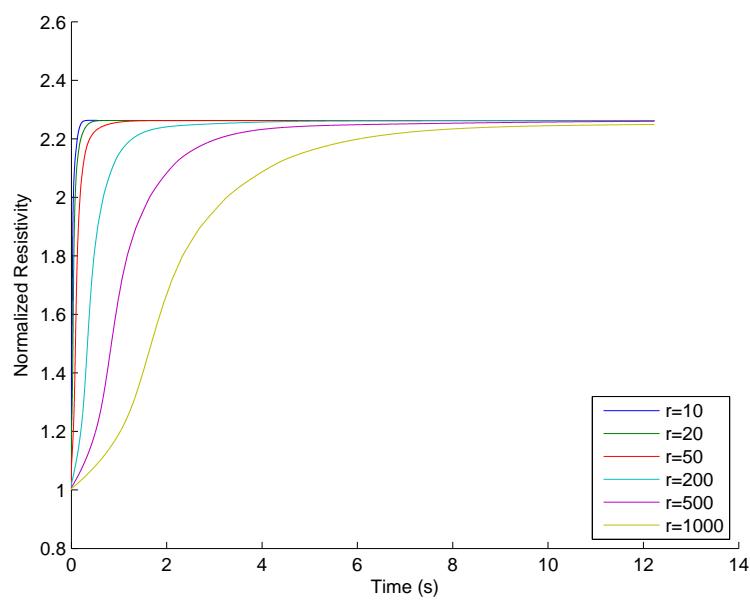


FIGURE 5.19

Resistivity change due to applied potential.(This behaves unexpectedly. I thought the resistivity would go higher with increased applied potential. Probably because of 1 line approximation for the electrolyte, the vertical potential is probably off)

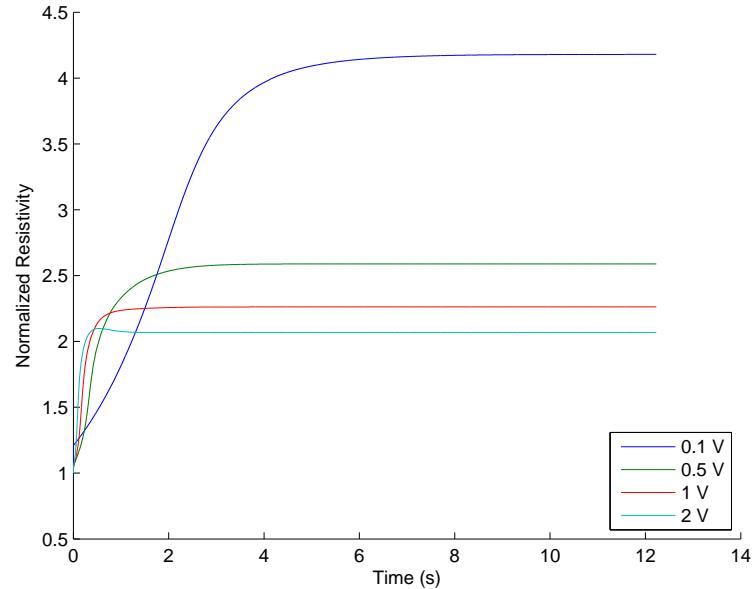


FIGURE 5.20

# Chapter 6

## 2-D Memristor Simulation

### 6.1 Effect of PEDOT:PSS Thickness

### 6.2 2-D Memristor Simulation Using a Pulse Train

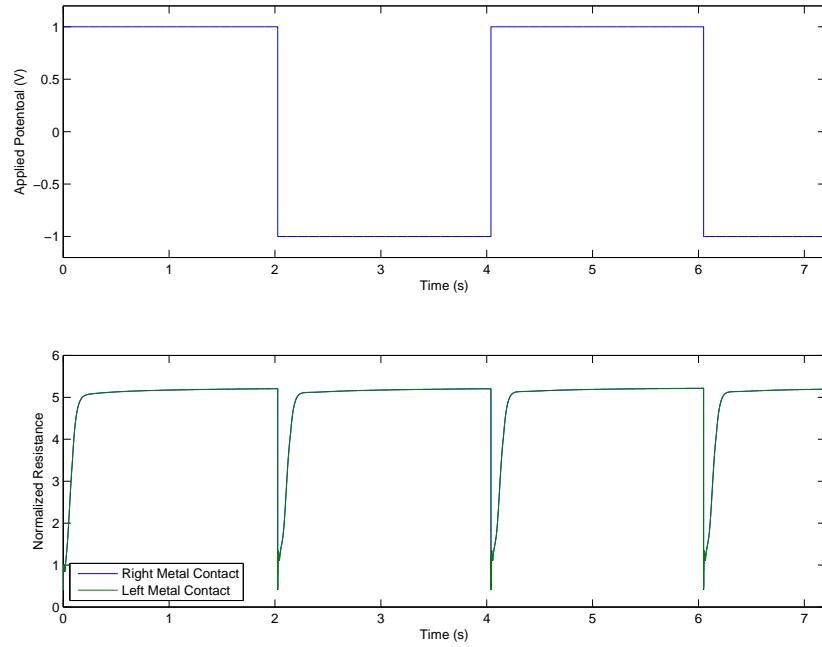


FIGURE 6.1

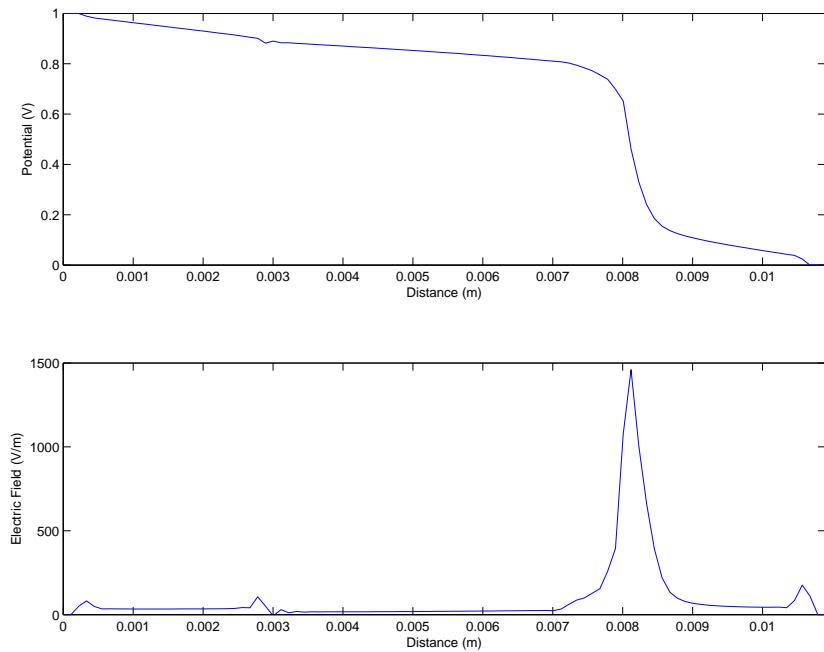


FIGURE 6.2

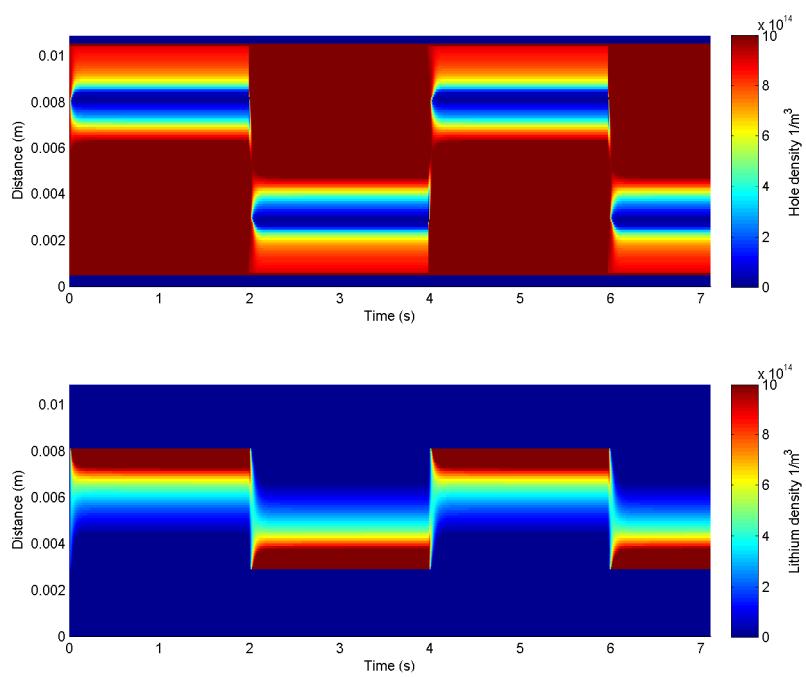


FIGURE 6.3

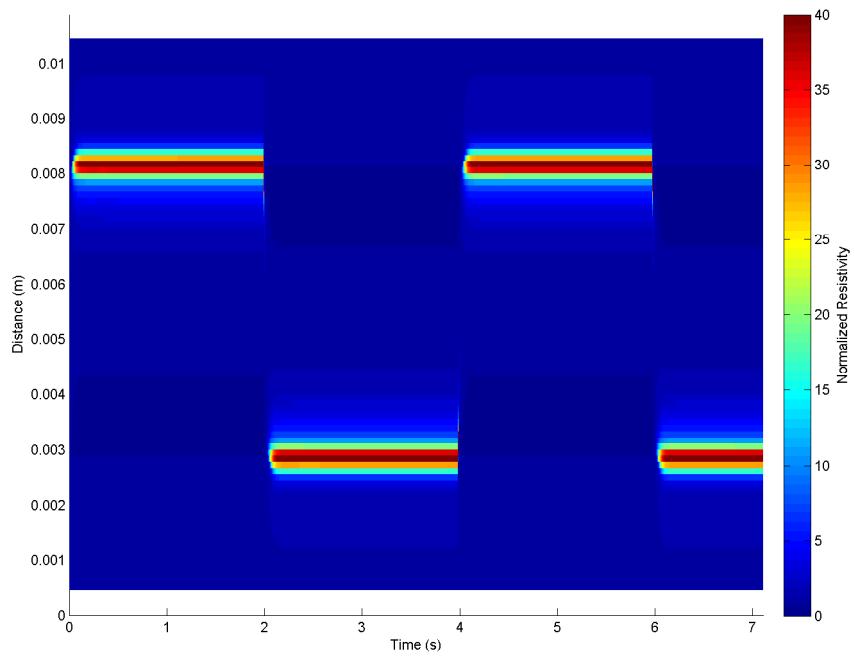


FIGURE 6.4

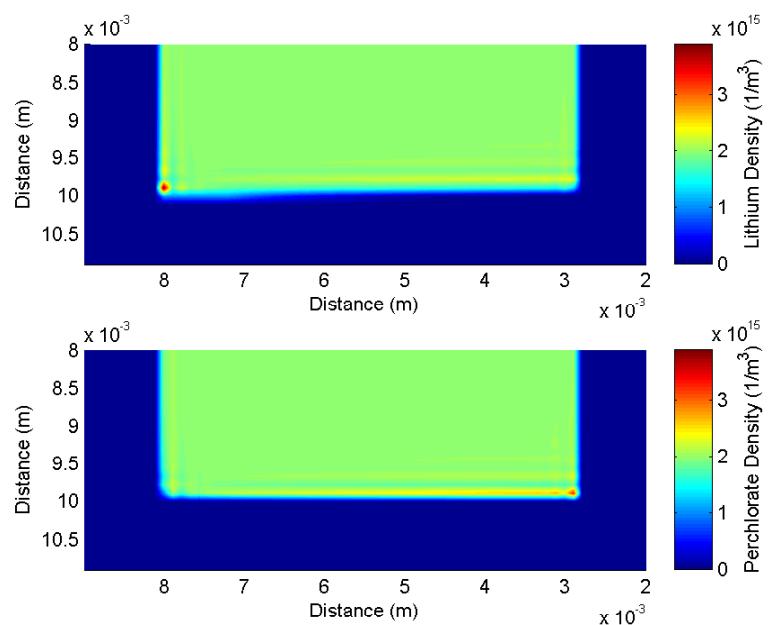


FIGURE 6.5

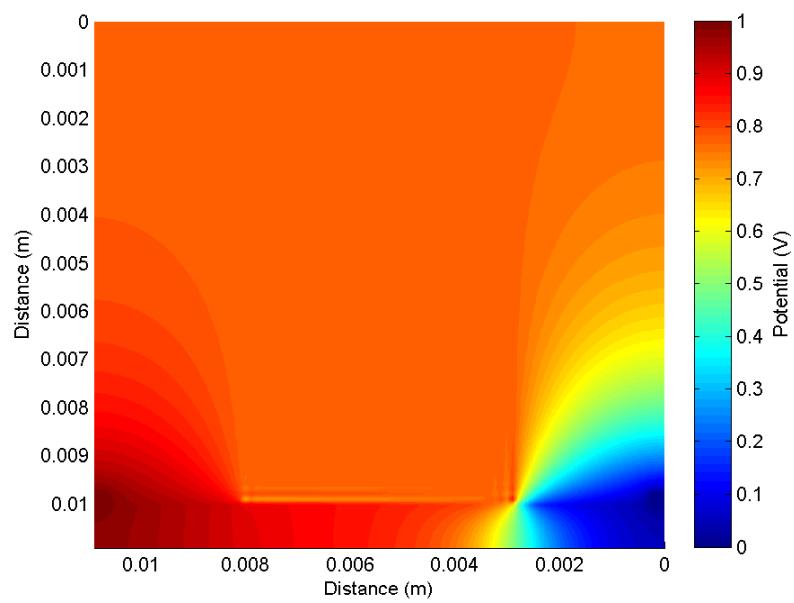


FIGURE 6.6

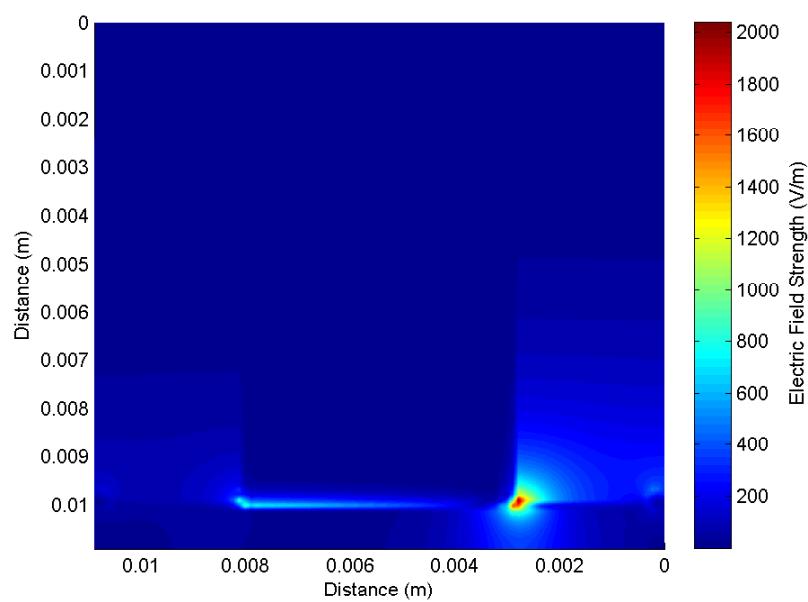


FIGURE 6.7

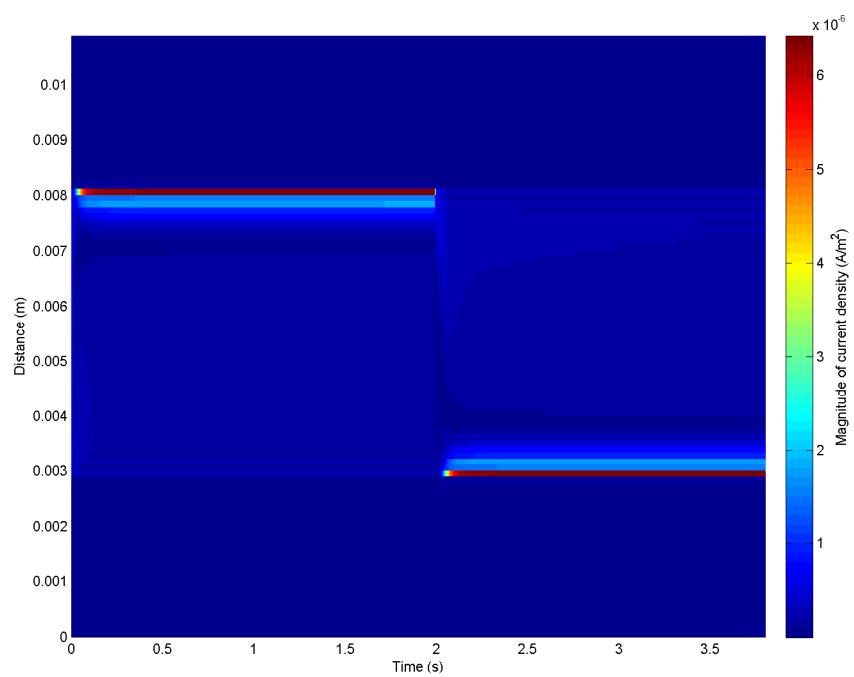


FIGURE 6.8

### 6.3 2-D Memristor Simulation Using a Sinusoid

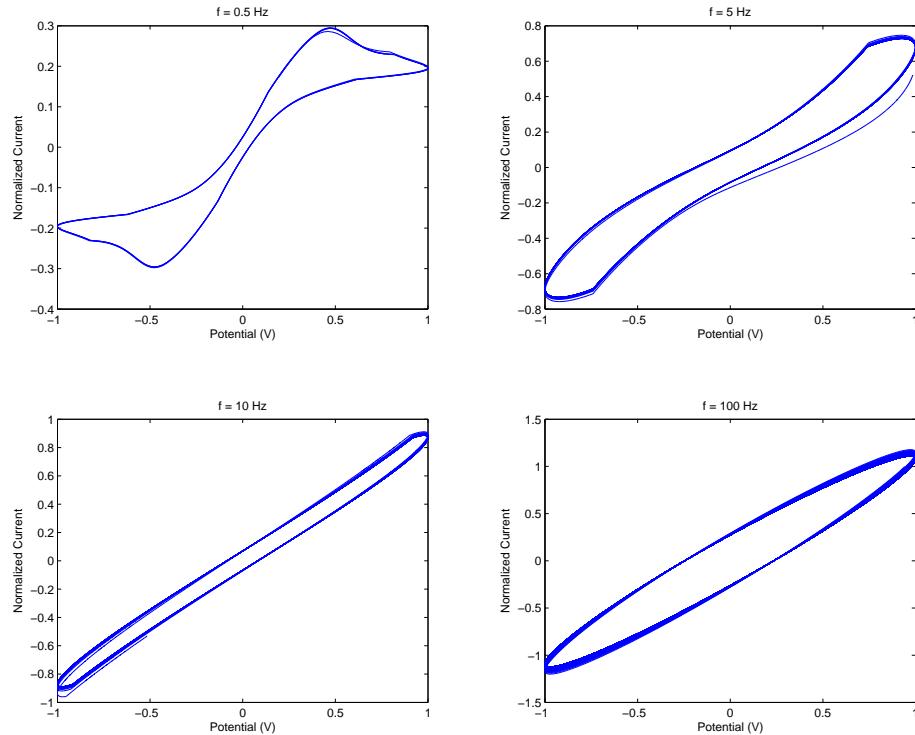


FIGURE 6.9

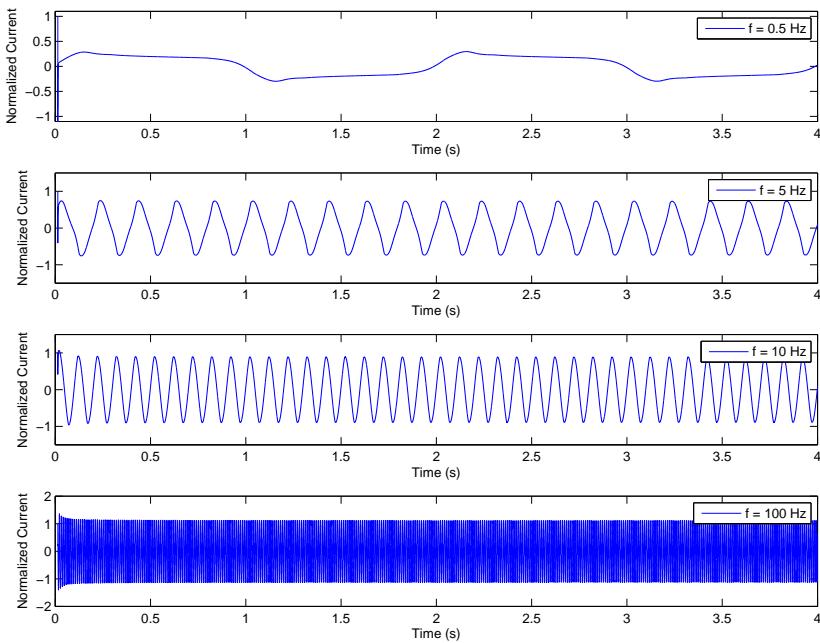


FIGURE 6.10

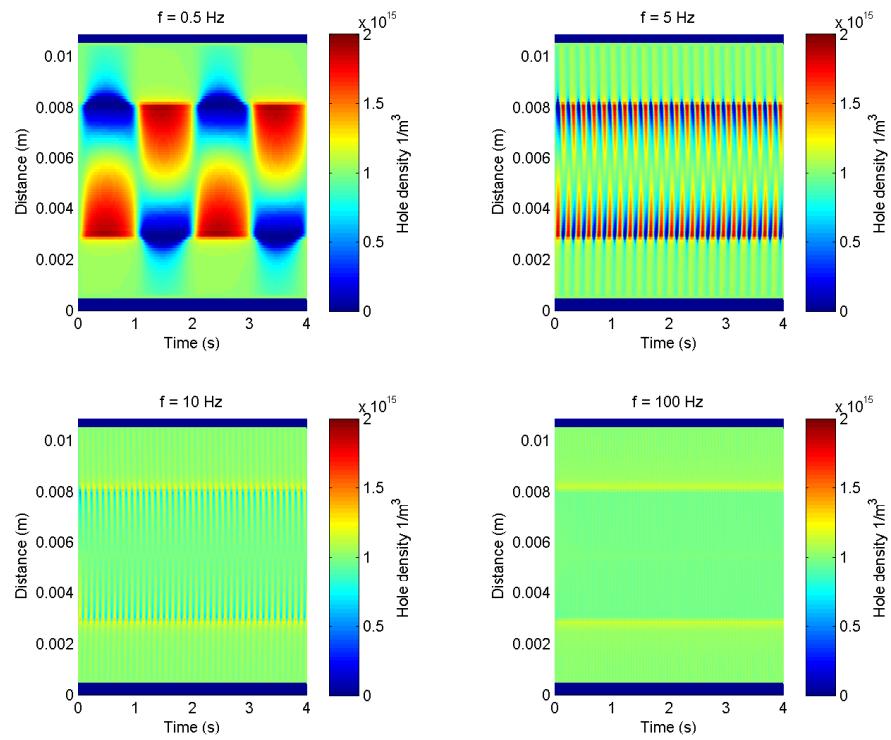


FIGURE 6.11

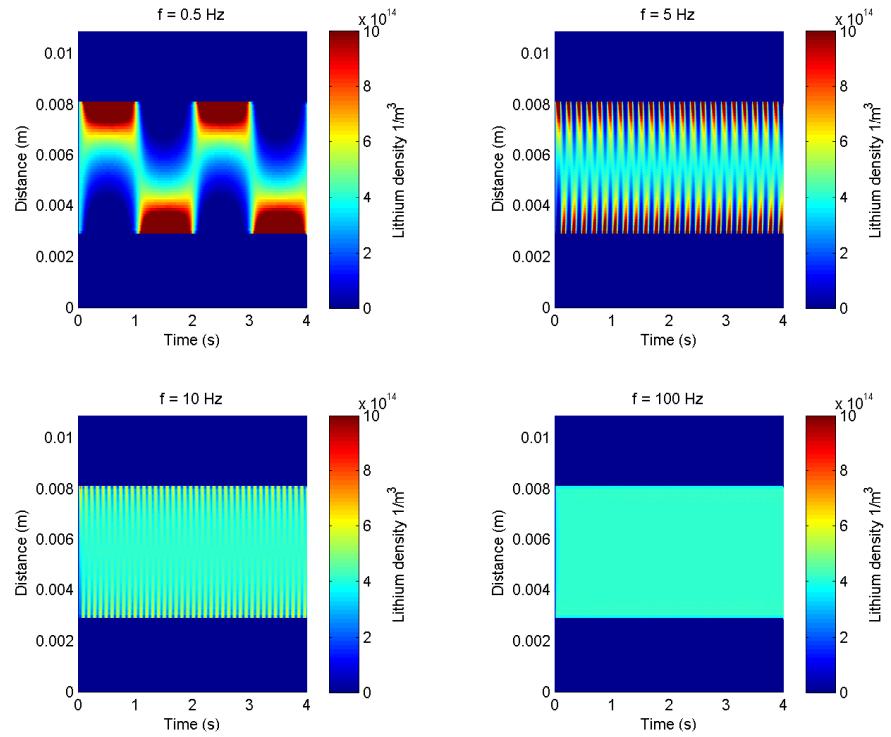


FIGURE 6.12

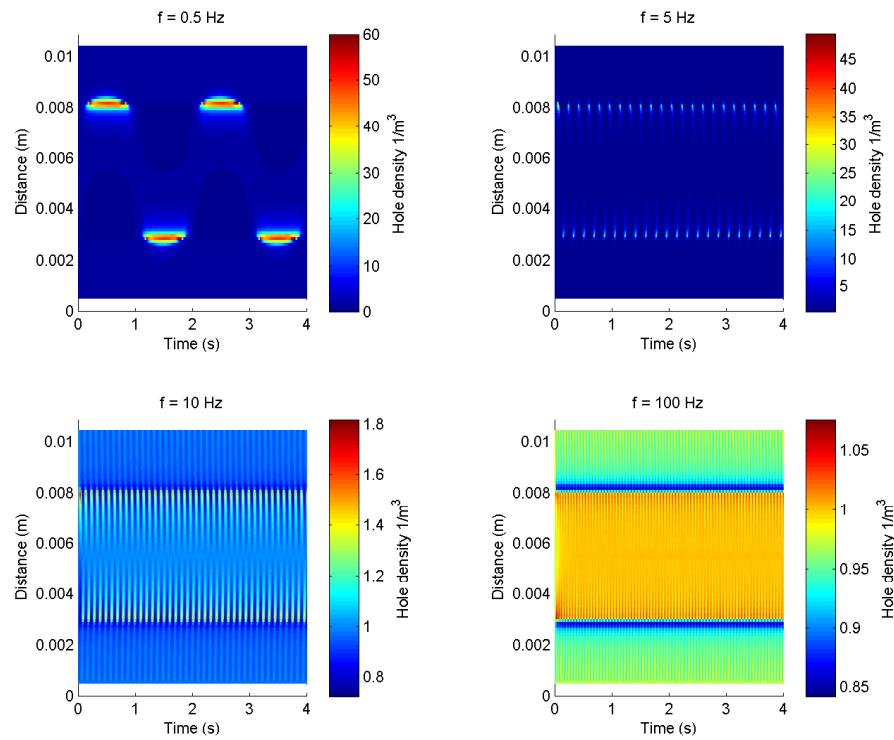


FIGURE 6.13

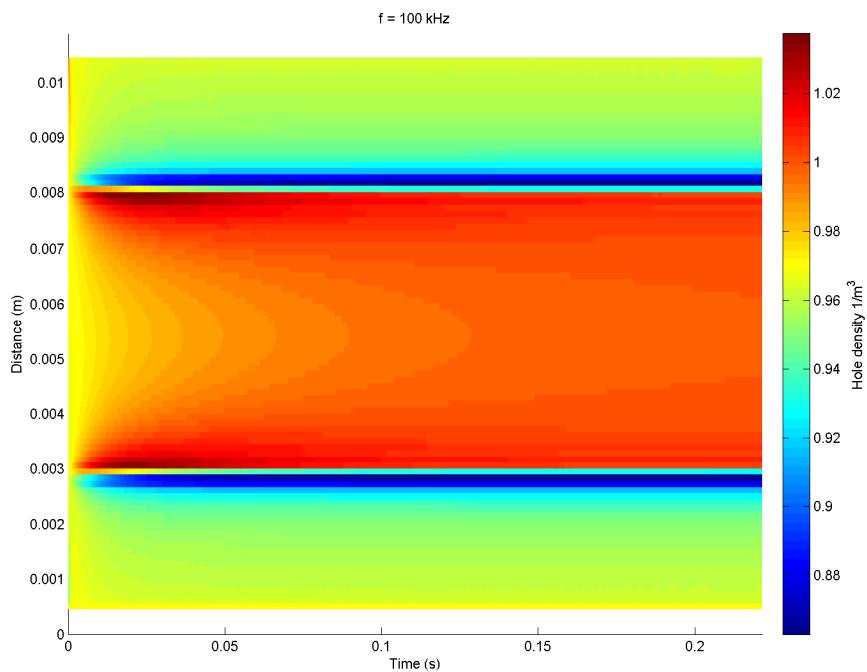


FIGURE 6.14

## 6.4 Experiment vs. Simulation

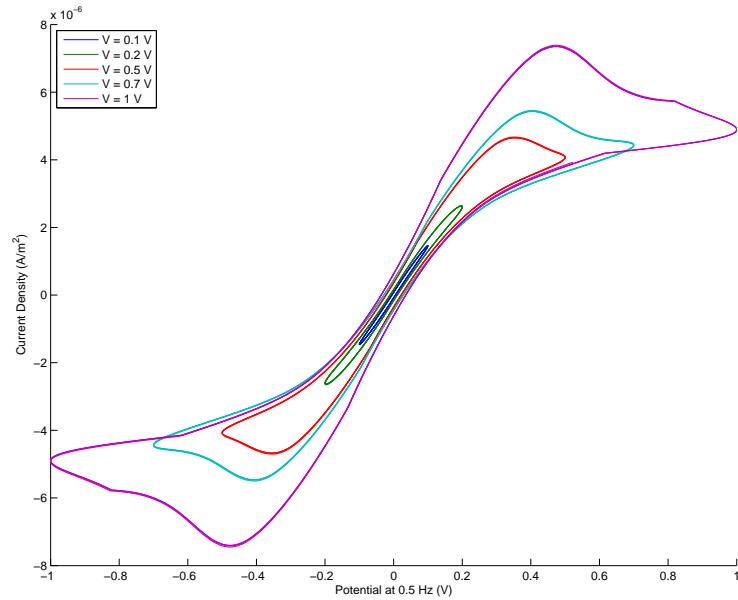


FIGURE 6.15

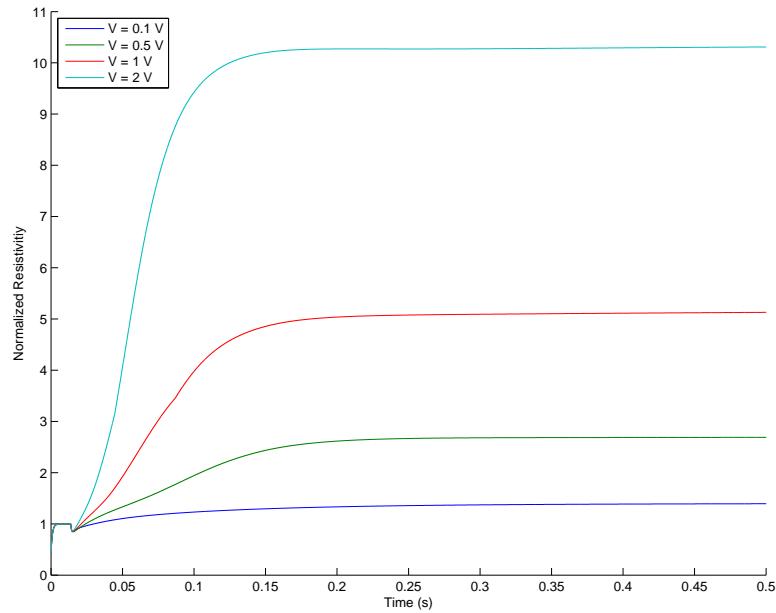


FIGURE 6.16

# Chapter 7

## Conclusion

The purpose of this thesis was to create a memristor model for computer simulation. We were able to model the change in conductivity through the movement of lithium and undoping of PEDOT. The transient simulation results match experimental data visually for both single channel and notched memristor. Although our simulation matched with the experimental data this model requires more work and can be improved through better numerical schemes, physical models and thorough experimentation. There are also few limitations of this model due to its size and physical features.

In terms of numerical methods there are a few down sides of using finite difference. The simulation for this device was done in 2-D instead of 3-D. This did not cause a lot of problems in our case since the structures we used were quite simple and produced reasonable results in 2-D. Unfortunately getting a transient response for devices using finite difference is computationally expensive. Even though we had a 100x100 grid, which is quite coarse, the simulation time was not less than 3 hours on a computer with multiple cores. Addition of another 100 points for a third dimension will make this simulation at least a 100 times longer. This makes 3-D simulations impractically long and very hard to test and optimize.

Another fundamental issue with this simulation arises from debye length. Maximum grid size in finite difference depends on the debye length, which is at least 5 or 6 orders of magnitude smaller than the device size. This means that we need at least  $10^5$  points in each direction in order to simulate this device. At this point the simulation becomes impossibly long so we had to compromise by either reducing the device size or carrier density.

The model we have developed is quite open for improvements on carrier transport models. A constant bulk mobility was used for holes in this model which is not the case in

an actual device. Holes move from site to site via hopping. Addition of lithium into PEDOT not only reduces the number of available holes but also decreases the number of possible sites through holes can move. A variable range hopping mechanism and thermal effects can be added into this model for a more complete simulation. Also PEDOT:PSS is a disordered material and the way it was deposited on a substrate can make a big difference on hole movement. Anisotropic hole mobility can be implemented in order to account for this issue.

Overall we have showed that it is possible to simulate a memristor using a simple finite difference method which can be very useful model in understanding the way ions and holes move in PEDOT. Our results show a very promising start for a vast research and development opportunities on memristor and polymer conductors.

## **Appendix A**

### **Title 1**

## **Appendix B**

### **Title 2**

## **Appendix C**

### **Title 3**

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