Vacuum deposition chamber

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Outline



2 The model

3 Mathematical model and simulations

4 Modifications of the problem

5 Conclusions

Overview of the problem

Problem statement

- In industry vacuum deposition is used for the generation of coating.
- Example: reflecting part of projectors for cars
- The deposition occurs in a chamber with two electrodes creating a gradient of potential.
- This modifies the repartition of the ions and electrons present in the chamber.
- The ions act as catalyser for the polymere reaction involved in the coating.

Main objective: Obtain in the chamber a large region with a high concentration of ions.



Our work : model and simulations

- Develop and study a model describing the process in the chamber
- 1D model (cartesian and cylindric)
- 2D model
- Influence of parameters (gas pressure, temperature, electrical voltage)

The simulations are all made with the software COMSOL.



Outline

Description of the problem

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The model: Hypothesis

- Electrostatic field
- Atmosphere of plasma (polymere) might contain all kind of positive and negative particules. We assume that the only positive particules are ions and the only negative particules are electrons.
- The chamber is cylindrical.
- The cathode is the small cylinder in the center of the chamber, which is used as the anode.



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Equations

- 3 variables : potential V, density of ions n_i and of electrons n_e
- Poisson equation: -△V = ^e/_ε(n_i n_e), where △ is the Laplacian, e is the elementary charge and ε is the dielectric constant of the atmosphere in the chamber.
- Continuity equations of the form: ^{∂ρ}/_{∂t} + divJ = source(x, t). In our stationary case:
 - $\nabla J_i = \mu_i n_e S(V)$
 - $\nabla J_e = \mu_e n_e S(V)$, where J_i and J_e represent the current densities of ions and electrons. μ_i and μ_e are the mobility of the ions and electrons. S(V) is the frequency of ionisation given by Townsend formula.

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• The current densities are described by a model called "drift-diffusion" : $J_i = -\mu_i n_i \nabla V - D_i \nabla n_i$ and $J_e = -\mu_e n_e \nabla V - D_e \nabla n_e$

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Dimensionless equations

We denote :

- $V = V_0$
- $n_i = n_0 n'_i$
- $n_e = n_0 n'_e$

and get the dimensionless equations:

•
$$-\triangle u = n_i - n_e$$
,
• $\nabla (-\pi \nabla u = \nabla \pi) - \mu_e$

•
$$\nabla (-n_i \nabla u - \nabla n_i) = \frac{\mu_e}{\mu_i} n_e S(u)$$

•
$$\nabla .(n_e \nabla u - \nabla n_e) = n_e S(u)$$

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Boundary conditions

For the electrical potential, electron and ion densities:

- At the anode:
 - The potential is $u = u_0$
 - Electrons are attracted and effectively absorbed, $J_e.\underline{n} = -n_e \nabla V.\underline{n}$
 - No emission of ions, $J_i \cdot \underline{n} = 0$
- At the cathode:
 - The potential is $u = -u_0$
 - Ions are attracted and effectively absorbed, $J_i \cdot \underline{n} = -n_i \nabla V \cdot \underline{n}$
 - No emission of electrons, $J_e.\underline{n} = 0$

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Model test

In the cartesian coordinates the equations become:

•
$$-u_{xx} = n_i - n_e$$

•
$$(-n_iu_x-n_{ix})_x=\alpha n_eS(u),$$

•
$$(n_e u_x - n_{ex})_x$$
,

with the boundary conditions

at the anode (x=1)

•
$$u = V_0 > 0$$

•
$$n_i u_x + n_{ix} = 0$$

• at the cathode (x=0)

•
$$u = -V_0$$

•
$$n_{ix} = 0$$

•
$$n_e u_x - n_{ex} = 0$$
,

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Results in 1-D

 $V_0 = 2500 \text{ V}$



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1D case

We consider a chamber with L >> R.

Then we have the cylindrical symmetry, i.e. 1D case.

• Gradient:
$$\nabla = \frac{\partial}{\partial r}$$

• The divergence:
$$\nabla = \frac{1}{r} \frac{\partial}{\partial r} (r \frac{\partial u}{\partial r})$$

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Results

Similar to the 1D cartesian case but much more sensitive to perturbations and need a higher external voltage to get to the same voltage profile.



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2D model : Boundary domain and symmetries

The boundary shape is more complex than in the 1-D case.



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Results in 2D



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Results in 2D







Potential

Density of ions

Density of electrons

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Modifications of the problem

We varied temperature and pressure: for an increase of 100°C we observed only slight changes.



For a small increase of P the density of ions increase considerably but stay localized in the same region.

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Conclusions

- The external voltage influences the distribution of ions
- With a potential large enough we obtain a localized distribution of ions
- A change in pressure changes the concentration and spread of ions
- Temperature doesn't have a notable influence

Next step: model with two electrodes.