

The Resonance Method: Evaluation of Self-Sustainment Voltage

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High-voltage laboratory

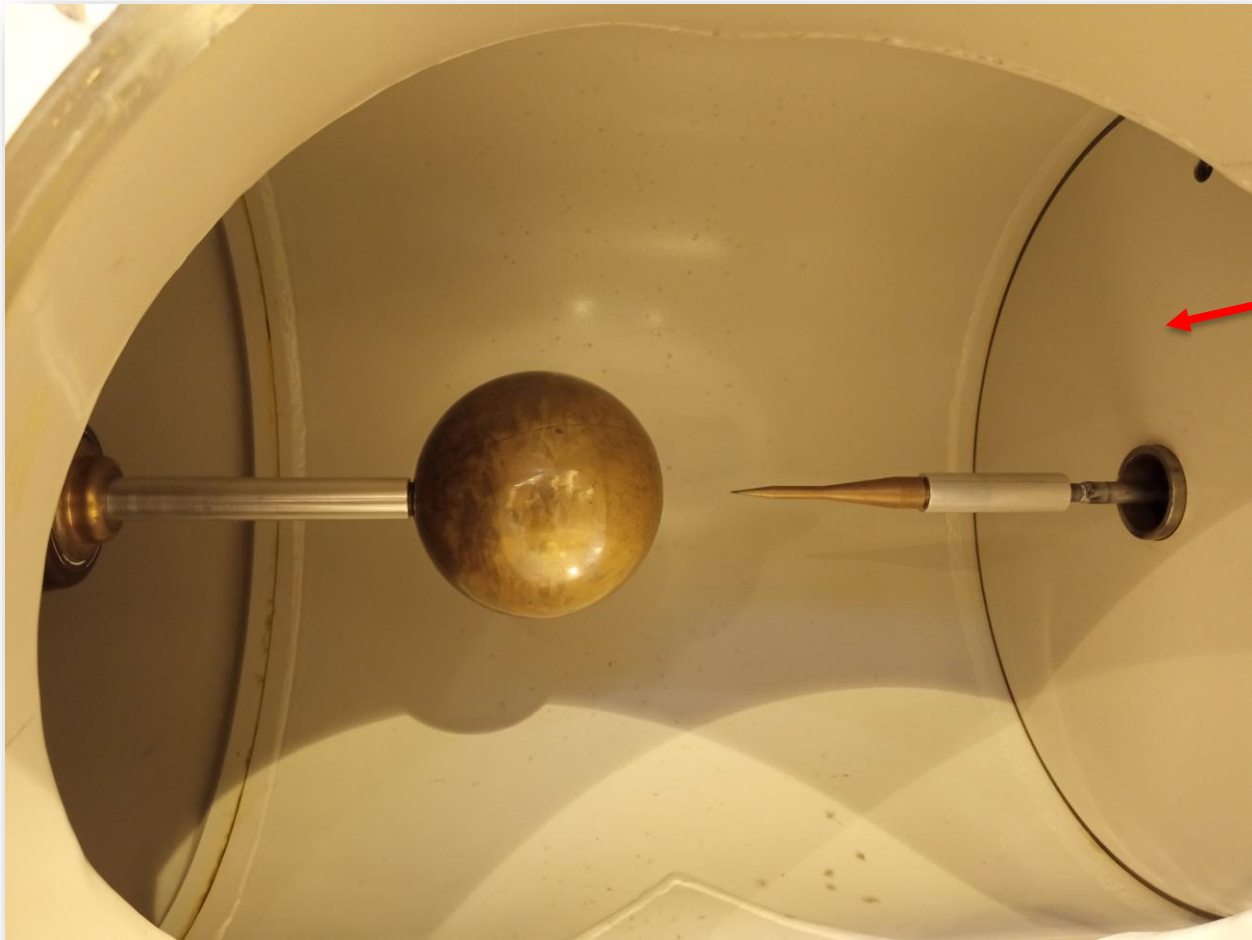


Fig. 1: Pressure vessel interior and electrodes.

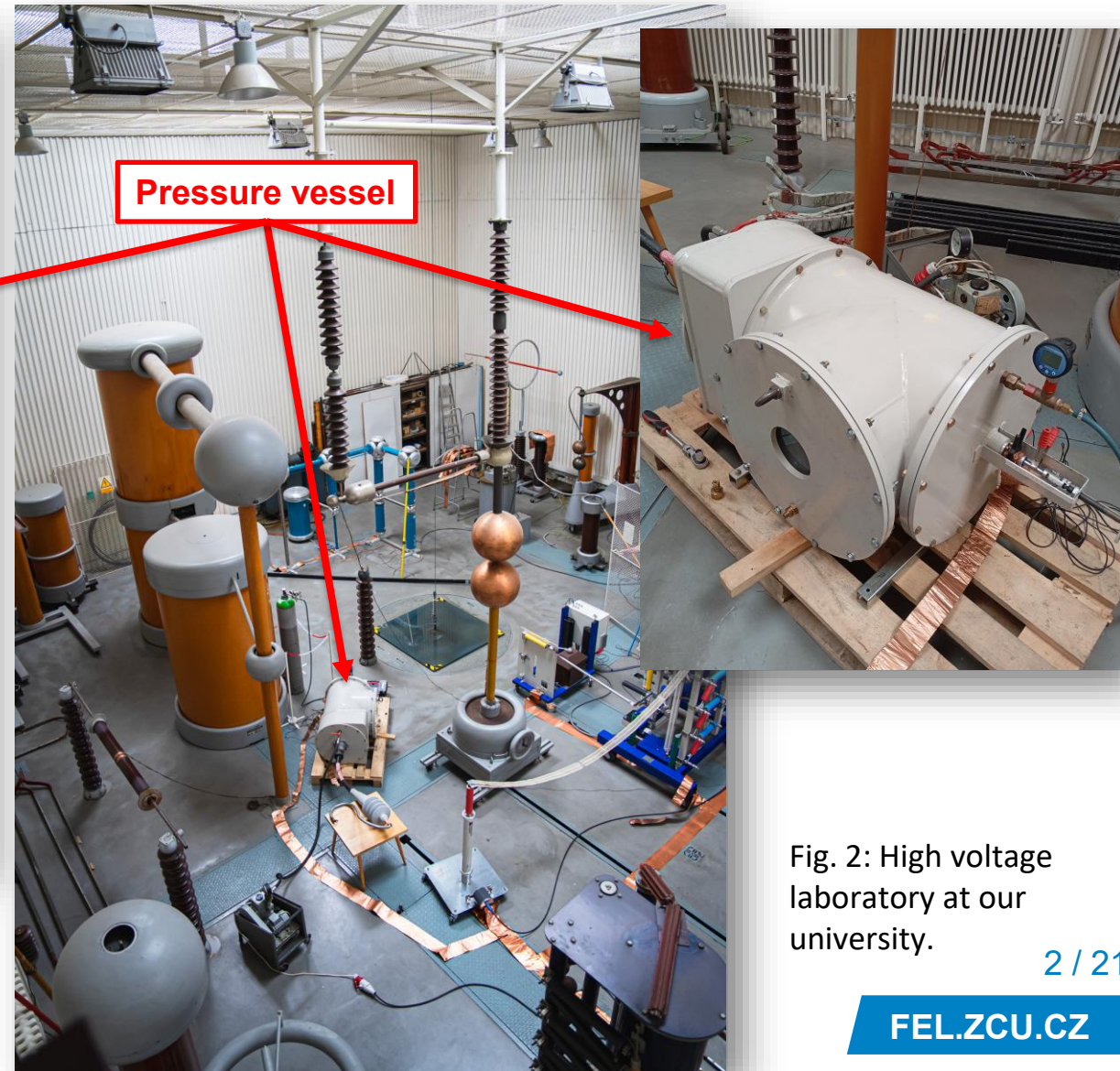


Fig. 2: High voltage laboratory at our university.

High-voltage laboratory

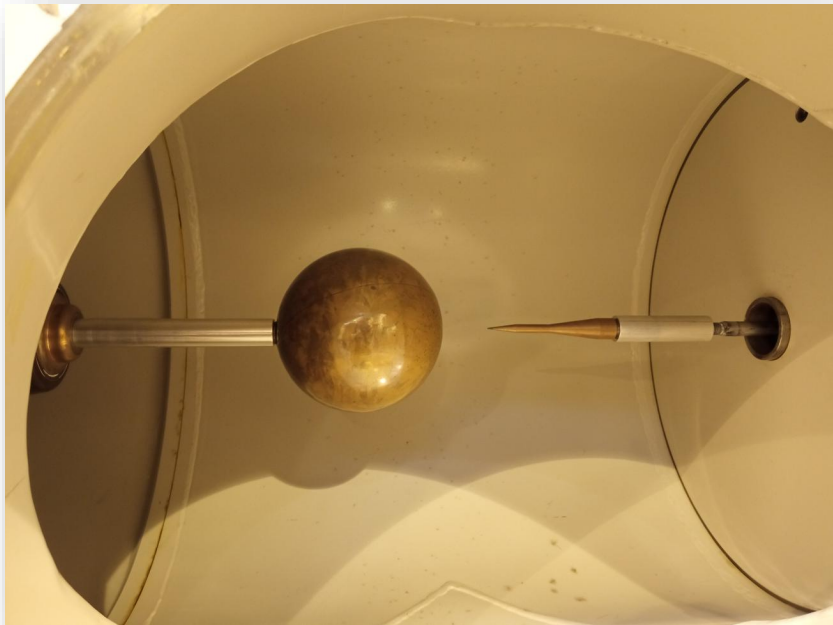


Fig. 1: Pressure vessel interior and electrodes.

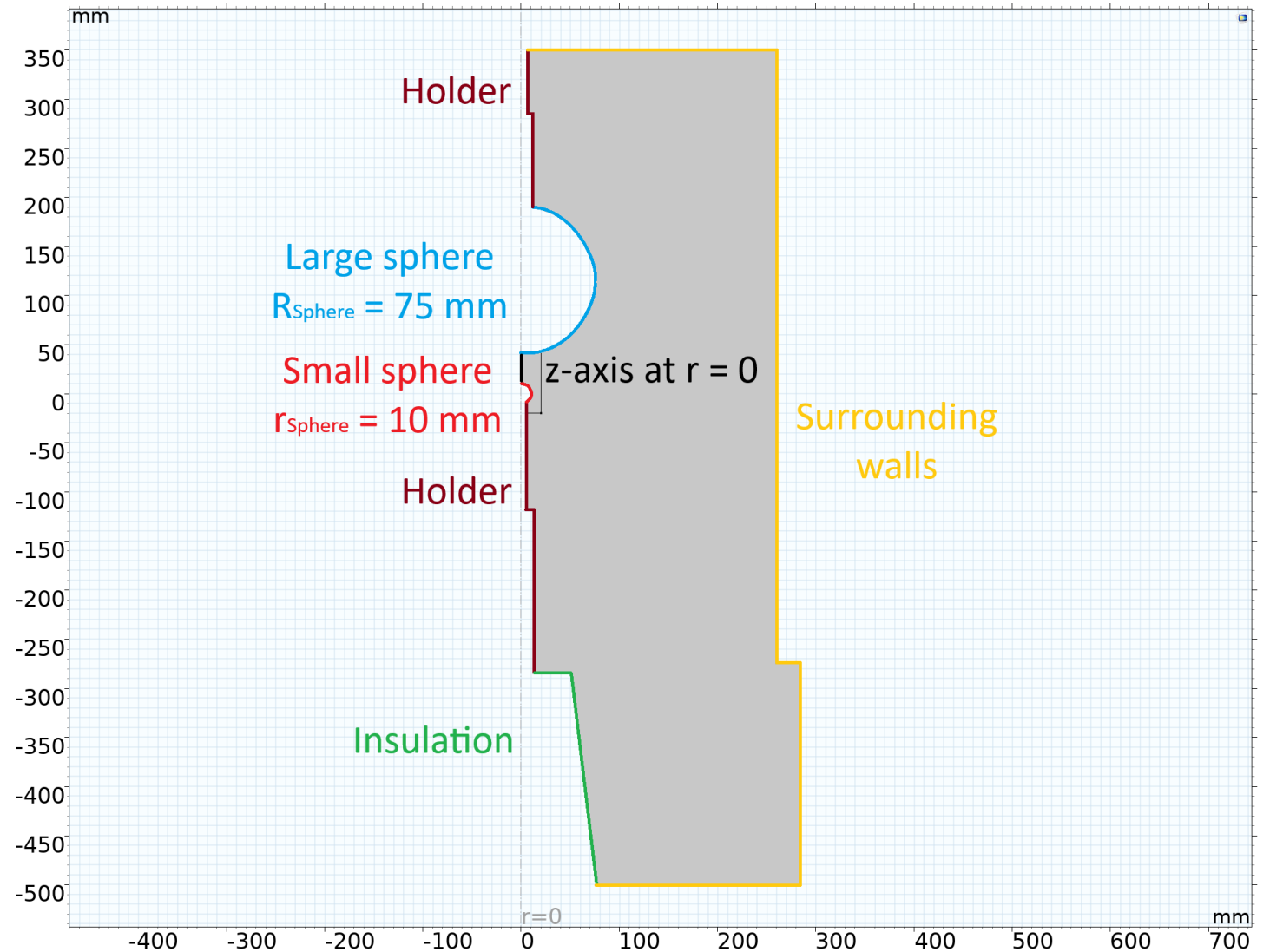


Fig. 3: Axisymmetric geometry in COMSOL Multiphysics, created using [1].

Mathematical model – Charged particle transport

Drift-diffusion-reaction equations (*Transport of Diluted Species*):

$$\frac{\partial n_c}{\partial t} + \nabla \cdot \left[-D_c \left(\frac{E}{N} \right) \nabla n_c \mp \mu_c \left(\frac{E}{N} \right) n_c \nabla \varphi \right] = S_c$$

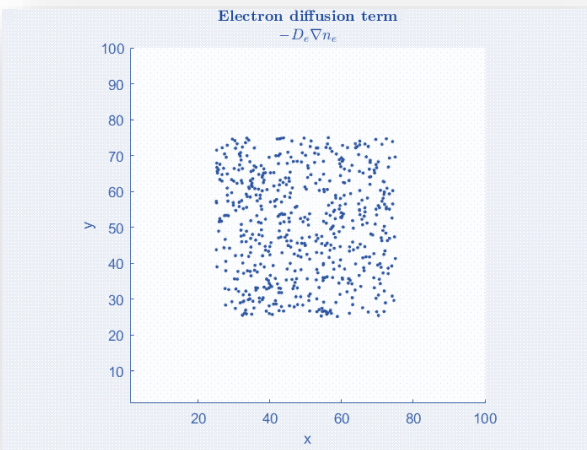


Fig. 4: Particle diffusion, created using [2].

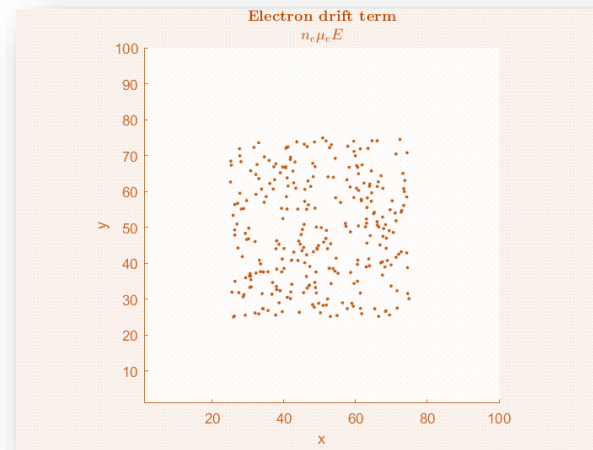


Fig. 5: Particle drift in electric field, created using [2].

Tab. 1: Variable definitions used in the drift-diffusion-reaction-equations.

n_c	Concentrations of particle species c (m^{-3}) – electrons and positive ions
D_c	Diffusivities of particle species c ($\text{m}^2 \text{s}^{-1}$)
μ_c	Mobilities of particle species c – drift ($\text{m}^2 \text{V}^{-1} \text{s}^{-1}$)
N	Neutral gas particle concentration (m^{-3})
E	Electric field intensity (Vm^{-1}) – E/N is reduced el. field (Td)
S_c	Source term of particle species c – reactions ($\text{m}^{-3} \text{s}^{-1}$)

Mathematical model – Charged particle transport

Drift-diffusion-reaction equation for electrons:

$$\nabla \cdot [-D_e(E/N)\nabla n_e + \mu_e(E/N)n_e\nabla\varphi] = S_e$$

Boundary conditions are set as follows:

- Flux** with convection term for upper larger sphere ($R_{\text{Sphere}} = 75 \text{ mm}$), holder and surrounding walls

$$J_{0,n_e} = \gamma\mu_e U(es.Er \cdot nr + es.Ez \cdot nz)n_p - \frac{n_e}{2} \sqrt{\frac{8k_B T_e}{\pi m_e}}$$

See slide 8

- Outflow** for lower smaller sphere ($r_{\text{Sphere}} = 10 \text{ mm}$) and holder
- Axial symmetry** at $r = 0 \text{ mm}$
- No flux** on the insulation

Tab. 2: Variable definitions used for boundary conditions.

γ	Secondary electron emission coefficient (0.0001)
T_e	Electron temperature (K)
m_e	Electron mass (kg)
k_B	Boltzmann constant (JK^{-1})
nr, nz	Normal components
$es.Er(z)$	Electric field in $r(z)$ direction (Vm^{-1})

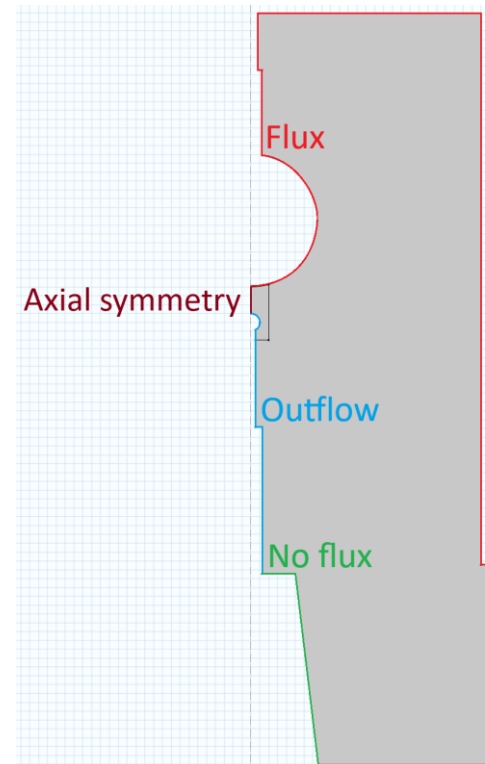


Fig. 6: Domain and boundary conditions, created using [1].

Mathematical model – Charged particle transport

■ Drift-diffusion-reaction equation for positive ions:

$$\nabla \cdot [-D_p \nabla n_p - \mu_p n_p \nabla \varphi] = S_p$$

■ Boundary conditions are set as follows:

- **Flux** with convection term for lower smaller sphere ($r_{\text{Sphere}} = 10 \text{ mm}$) and holder

$$J_{0,n_p} = -\frac{n_p}{2} \sqrt{\frac{8k_B T_p}{\pi m_p}}$$

- **Outflow** for upper larger sphere ($R_{\text{Sphere}} = 75 \text{ mm}$), holder and surrounding walls
- **Axial symmetry** at $r = 0 \text{ mm}$
- **No flux** on the insulation

Tab. 3: Variable definitions used for boundary conditions.

T_p	Positive ion temperature (K) – set equal to 300 K
m_p	Positive ion mass (kg)

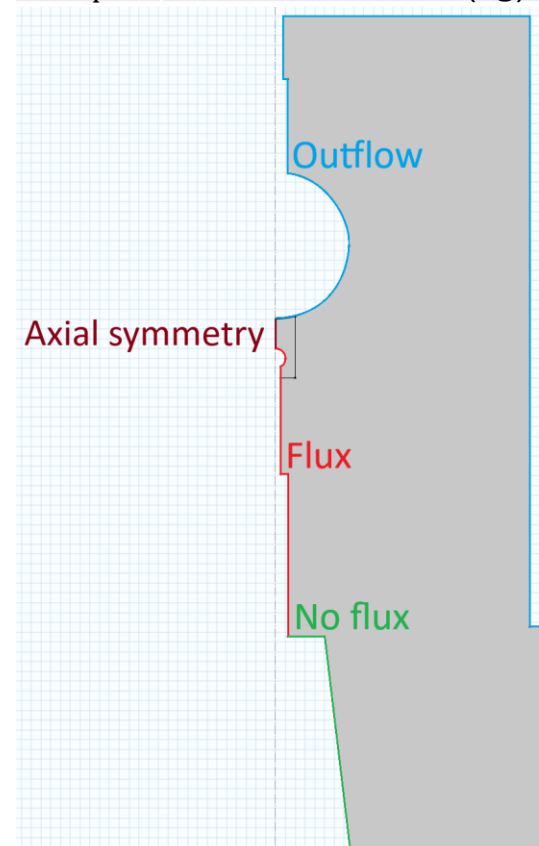


Fig. 7: Domain and boundary conditions, created using [1].

Mathematical model – Charged particle transport

Source term for electrons:

$$S_e = W + S_{\text{coll(eff)}} + S_{\text{ph}}$$

$$S_{\text{coll(eff)}} = \alpha_{\text{eff}}(E/N) \cdot N \cdot \mu_e(E/N) \cdot E \cdot n_e$$

Source term for positive ions:

$$S_p = W + S_{\text{coll}} + S_{\text{ph}}$$

$$S_{\text{coll}} = \alpha(E/N) \cdot N \cdot \mu_e(E/N) \cdot E \cdot n_e$$

W is an independent source term set equal to $100 \text{ m}^{-3} \text{ s}^{-1}$

S_{ph} is photoionization rate, see slide 9

α and α_{eff} are ionization coefficients defined according to [3]

All transport coefficients for electrons and electron temperature are defined according to [4]

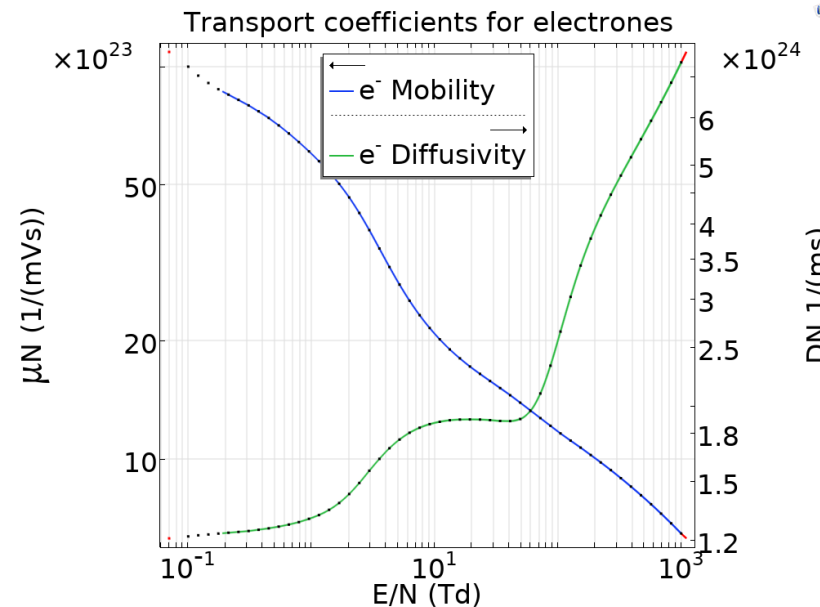


Fig. 8: Transport (electron e mobility μ_e and diffusivity D_e) coefficients as a functions of reduced electric field E/N (Td) [4].

Tab. 4: Transport coefficients for positive ions p [5].

μ_p	$2.0\text{e-}4 \text{ [m}^2\text{/V/s]}$	Positive ion p mobility
D_p	$5.05\text{e-}6 \text{ [m}^2\text{/s]}$	Positive ion p diffusivity

Mathematical model – Electric field

- **Laplace equation for electrostatic potential**

(*Electrostatics*): $\Delta\varphi = 0$

- **Electrostatic field intensity:** $E = -\nabla\varphi$

- **Space charge** does not disturb external electric field
– **linear problem**

- **External electric field** is calculated only once and then scaled by voltage U

- **Boundary conditions** are set as follows:

- **1 V** for lower smaller sphere ($r_{\text{Sphere}} = 10 \text{ mm}$) and holder
- **Ground** for upper larger sphere ($R_{\text{Sphere}} = 75 \text{ mm}$), holder and surrounding walls
- **Axial symmetry** at $r = 0 \text{ mm}$
- **Zero charge** on the insulation

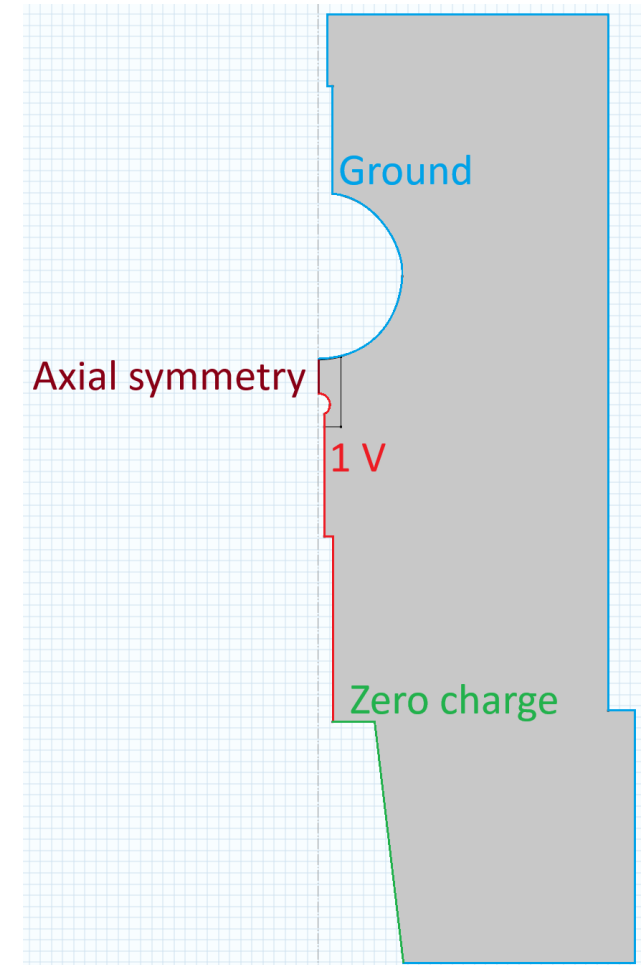


Fig. 9: Domain and boundary conditions, created using [1].

Mathematical model – Photoionization

- **Three-exponential Helmholtz** model for photoionization rate $S_{ph,i}$ proposed by A. Bourdon, et al. in [6] (*Coefficient Form PDE*):

$$\Delta S_{ph,i} - (\lambda_i \cdot p_{O_2})^2 \cdot S_{ph,i} = -A_i \cdot p_{O_2}^2 \cdot \chi_{ext}(E/N) \cdot f_q \cdot S_{coll}$$

$$S_{ph} = \sum_{i=1}^3 S_{ph,i}$$

- Quenching factor f_q , partial pressure of molecular oxygen p_{O_2} and parameters λ_i and A_i (where $i = 1, 2, 3$) are summarised in the Table 5 and collisional ionization S_{coll} is defined as follows:

$$S_{coll} = \alpha(E/N) \cdot N \cdot \mu_e(E/N) \cdot E \cdot n_e$$

$$E/N = \left(1e21 \cdot U \cdot \frac{es.normE}{N} \right) \text{ and } E = U \cdot es.normE$$

- Boundary conditions are set according to Fig. 10

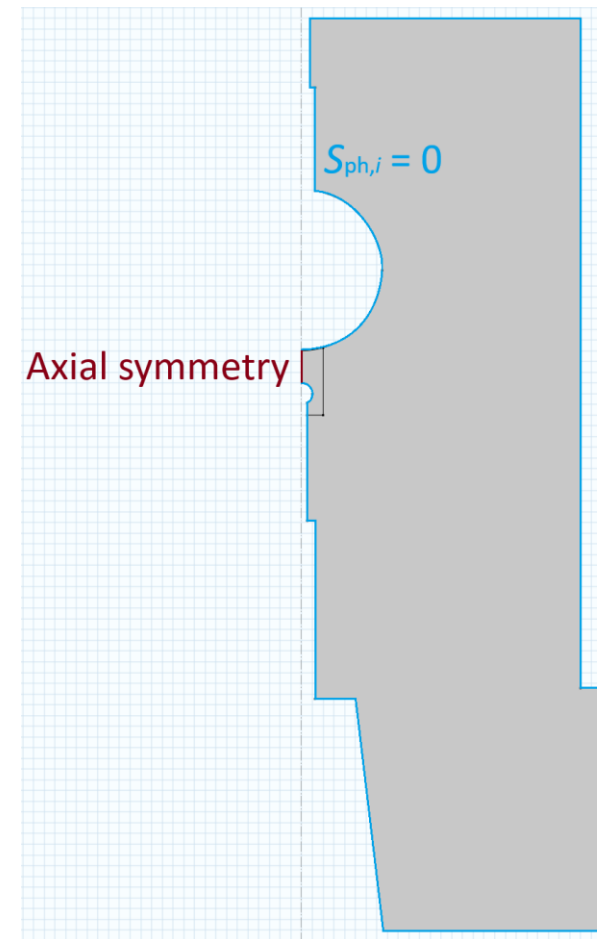


Fig. 10: Domain and boundary conditions, created using [1].

Tab. 5: Photoionization model parameters [6].

A_1	1.986e-4 [cm ⁻² *Torr ⁻²]
A_2	0.0051 [cm ⁻² *Torr ⁻²]
A_3	0.4886 [cm ⁻² *Torr ⁻²]
λ_1	0.0553 [cm ⁻¹ *Torr ⁻¹]
λ_2	0.1460 [cm ⁻¹ *Torr ⁻¹]
λ_3	0.89 [cm ⁻¹ *Torr ⁻¹]
p_{O_2}	150 [Torr]
f_q	0.037975

Mathematical model – Photoionization

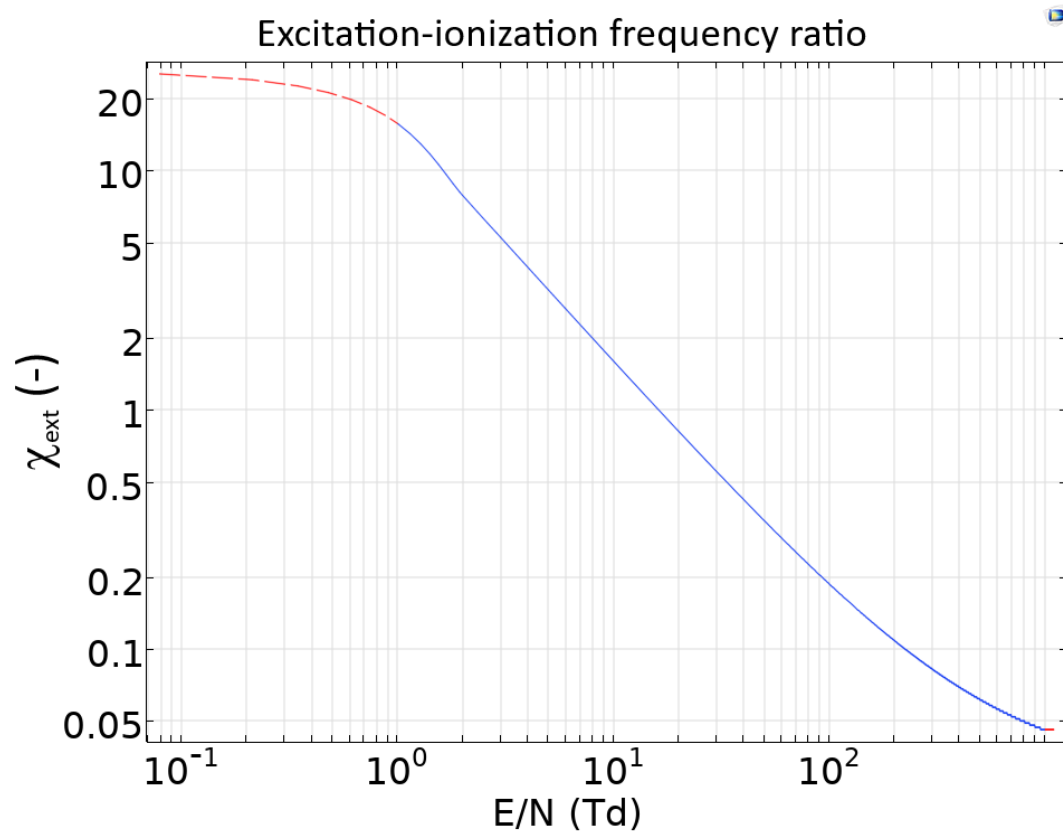


Fig. 11: Excitation-ionization frequency ratio χ_{ext} from slide 9 as a function of reduced electric field E/N (Td) for the photoionization model [6].

The Resonance Method

- Method for calculating self-sustainment voltage U_0 – transition between **non-self-sustaining** and **self-sustaining** discharge – proposed by prof. M. S. Benilov, et al. in [7]
- **Linear stationary ($\partial/\partial t \rightarrow 0$) study** for the resonance of current I
 - Parametric sweep for voltage U
 - Once the current I diverges, the resonance is found at U_0

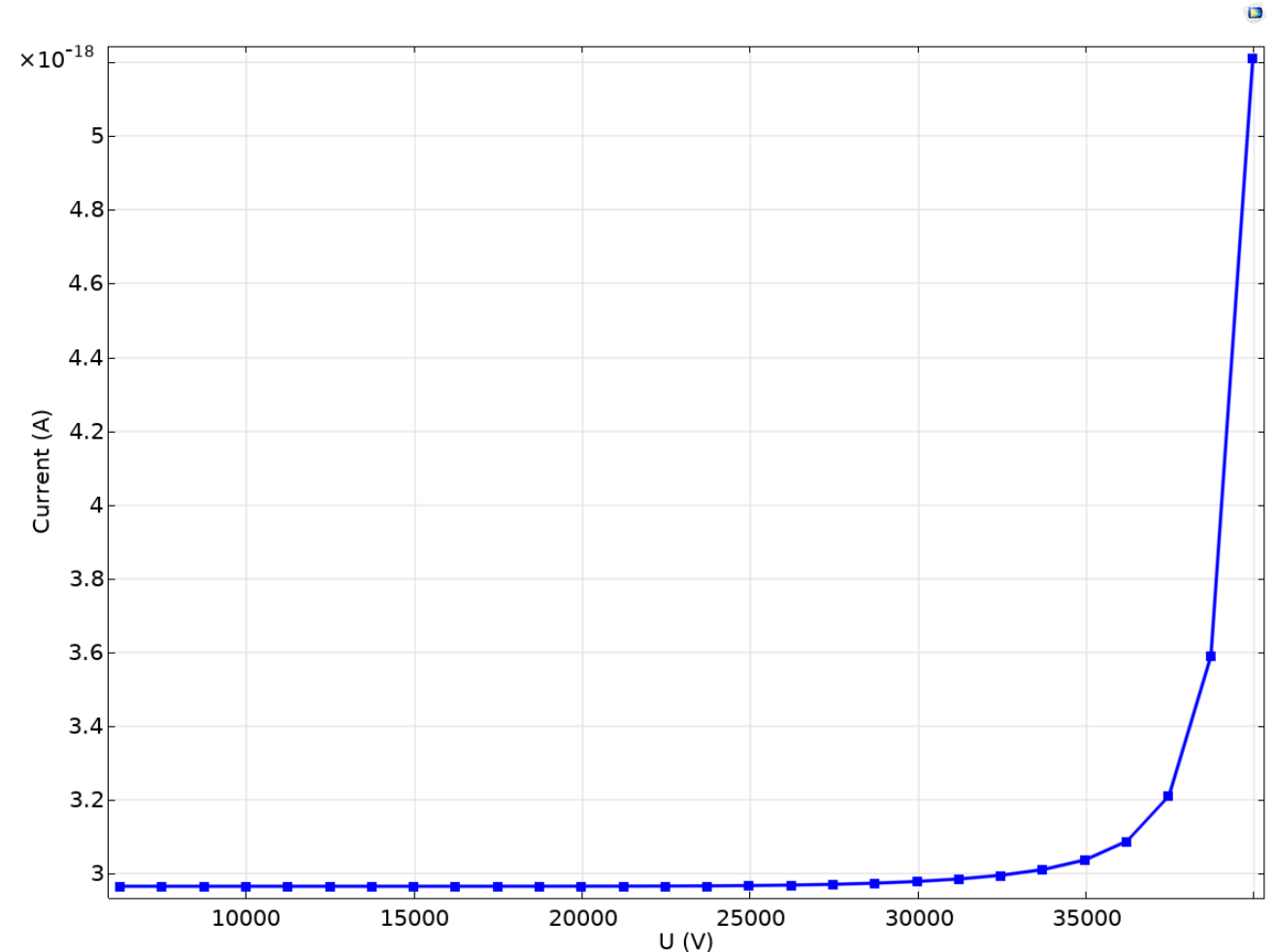


Fig. 12: Current-voltage characteristic of non-self-sustained discharge for air, $d = 30$ mm and $p = 1$ atm, created using [1].

The Resonance Method

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- **Linear stationary ($\partial/\partial t \rightarrow 0$) study** for the resonance of current I
 - Parametric sweep for voltage U
 - Once the current I diverges, the resonance is found at U_0
 - Particle concentrations switches to negative (non-physical) values for $U > U_0$ to ensure the solution continues for any value of U
 - To model discharge behaviour for $U > U_0$ nonlinear coupling must be added

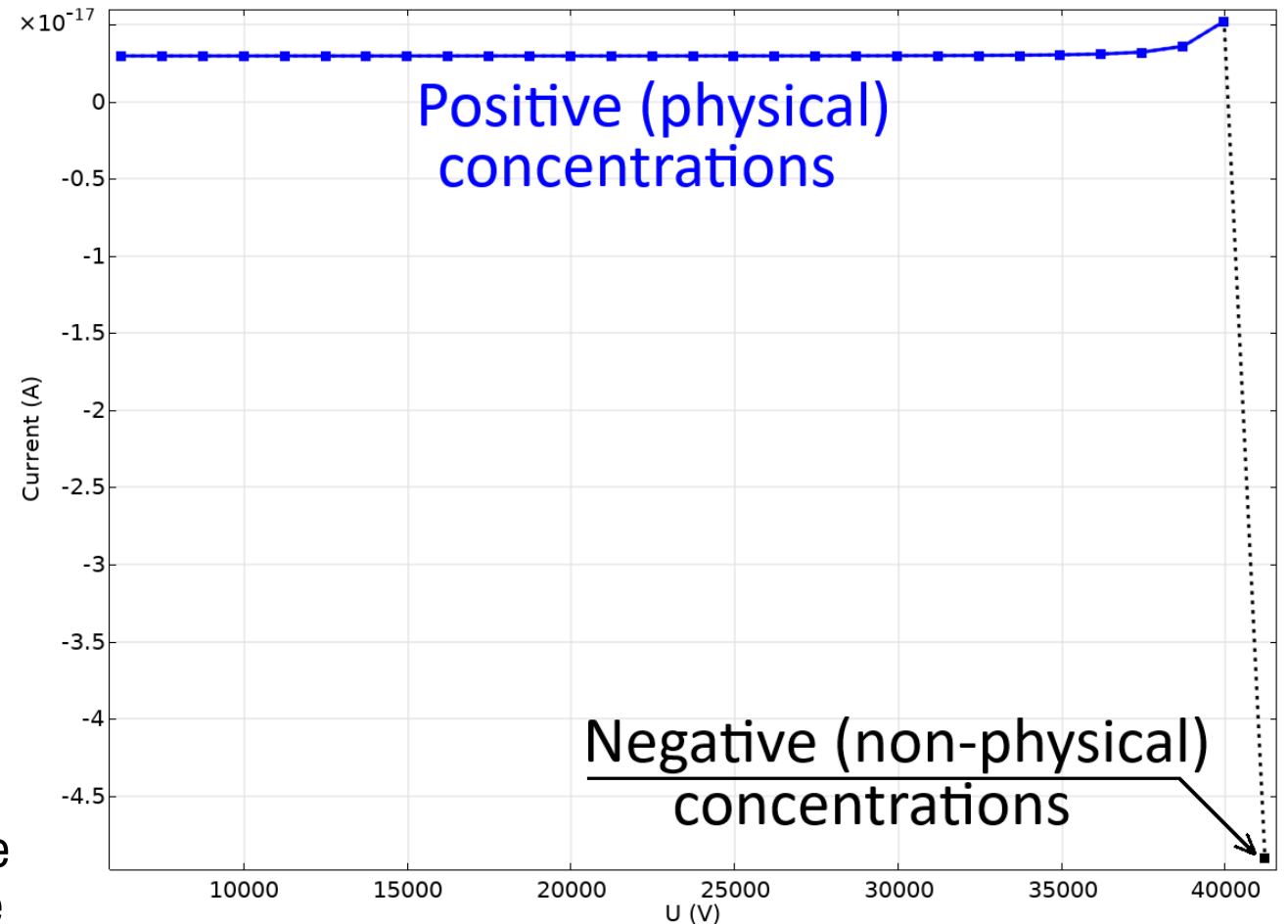


Fig. 13: Current-voltage characteristic of non-self-sustained discharge with resonance point for air, $d = 30$ mm and $p = 1$ atm, created using [1].

The Resonance Method

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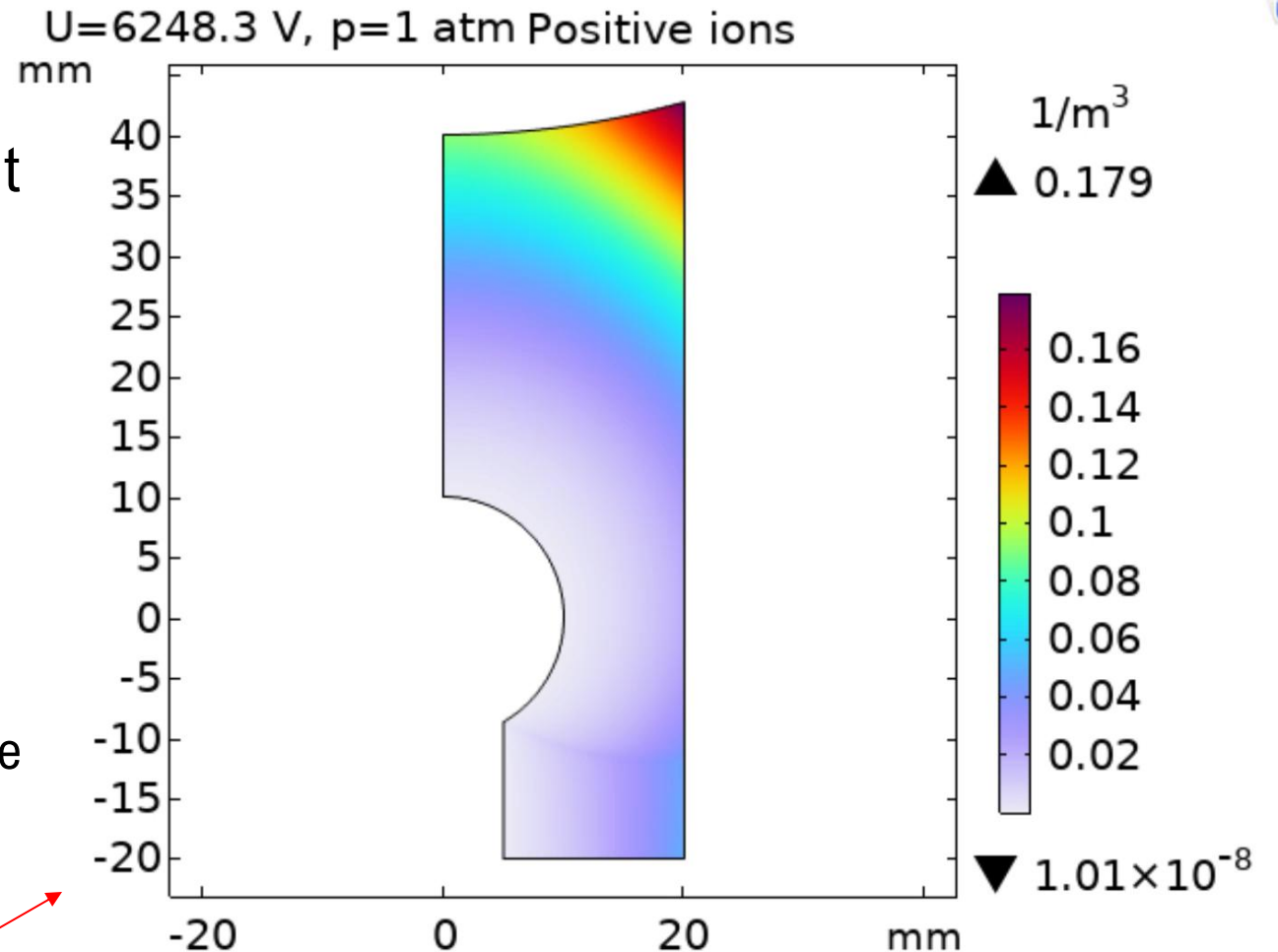
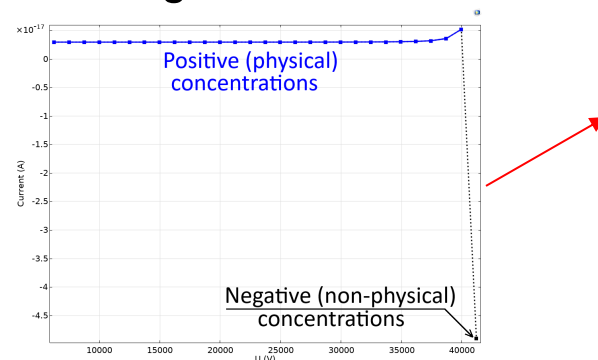


Fig. 14: Concentration distribution of positive ions for voltage values in Fig. 13 for air, $d = 30$ mm and $p = 1$ atm (animation), created using [1].

The Resonance Method

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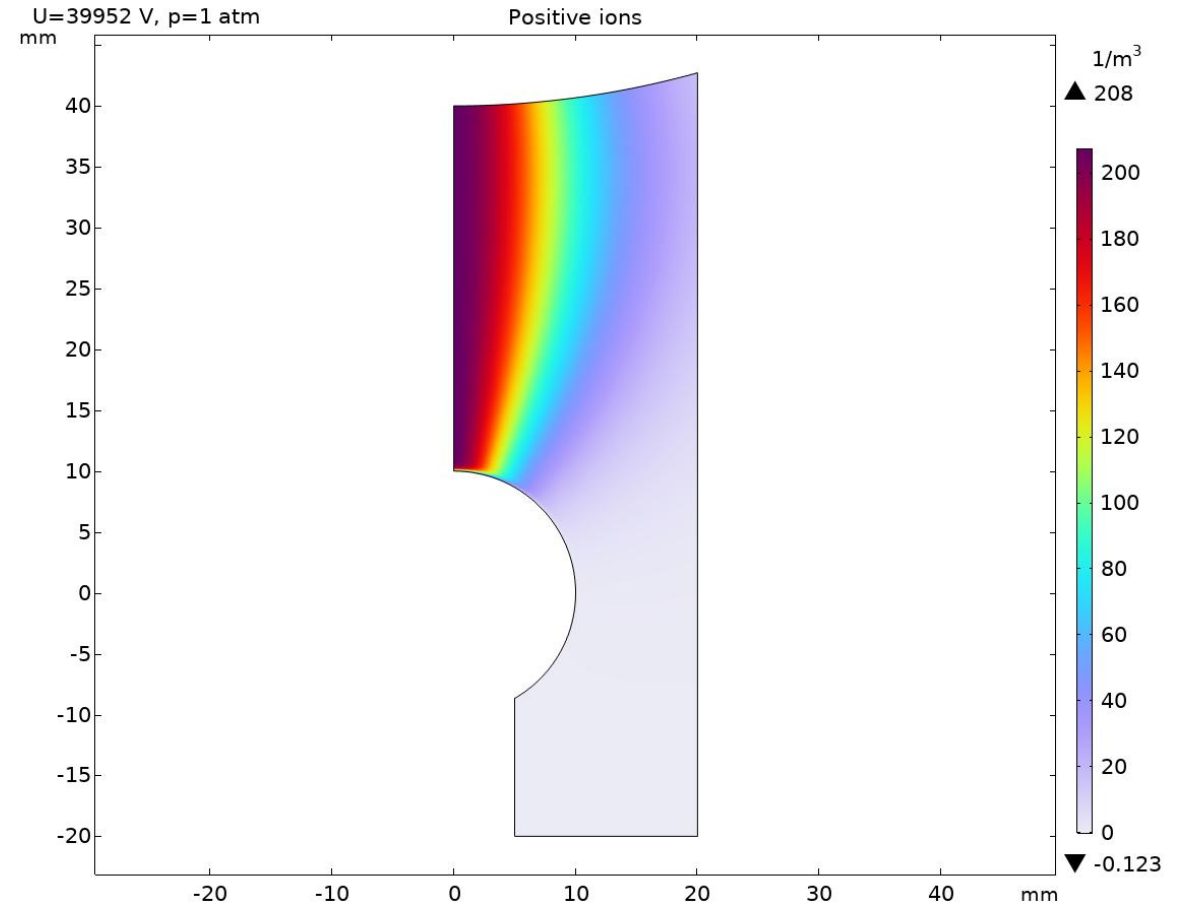


Fig. 15: Concentration distribution of positive ions for voltage just below the resonance point for air, $d = 30$ mm and $p = 1$ atm – positive (physical) values, created using [1].

The Resonance Method

- Method for calculating self-sustainment voltage U_0 – transition between **non-self-sustaining** and **self-sustaining** discharge – proposed by prof. M. S. Benilov, et al. in [7]
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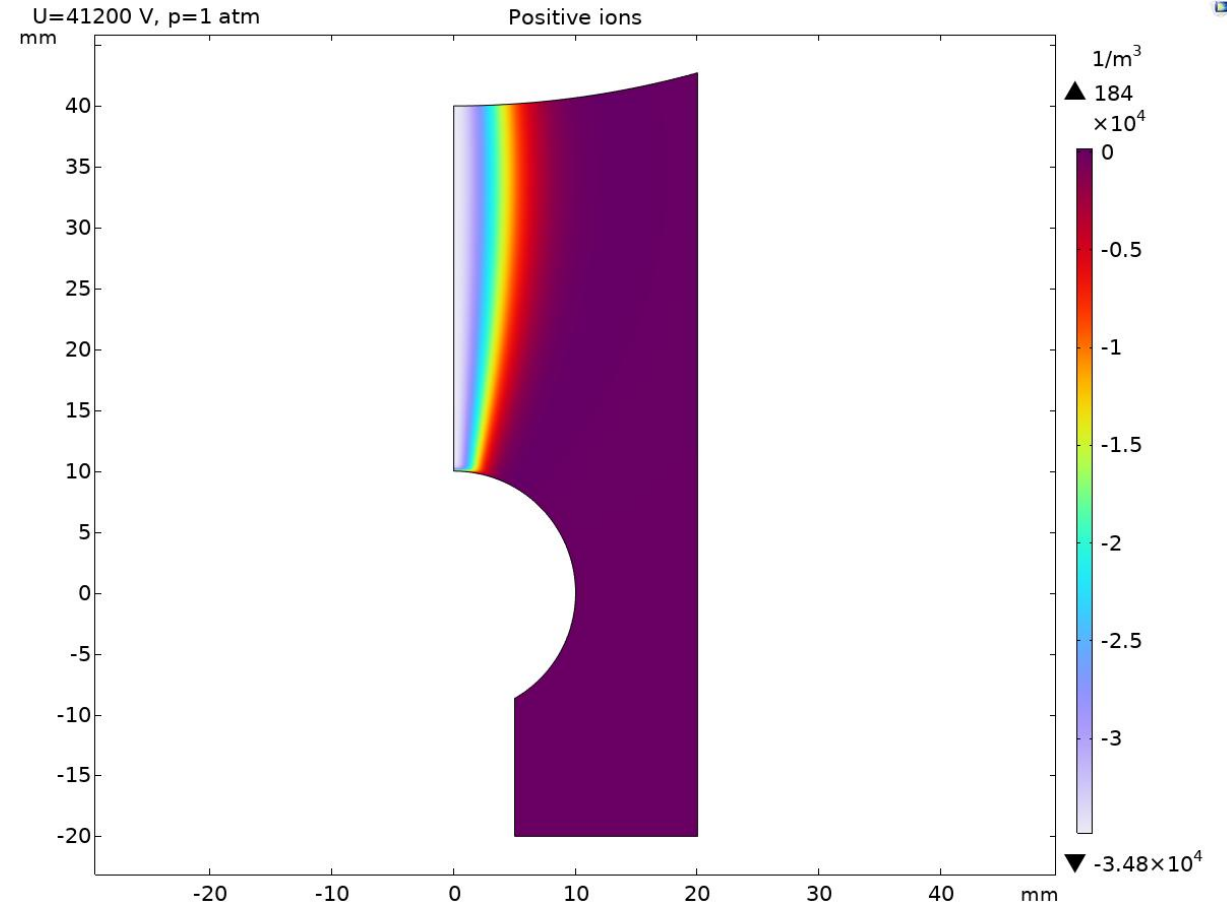


Fig. 16: Concentration distribution of positive ions for voltage at the resonance point for air, $d = 30$ mm and $p = 1$ atm – negative (non-physical) values, created using [1].

The Resonance Method

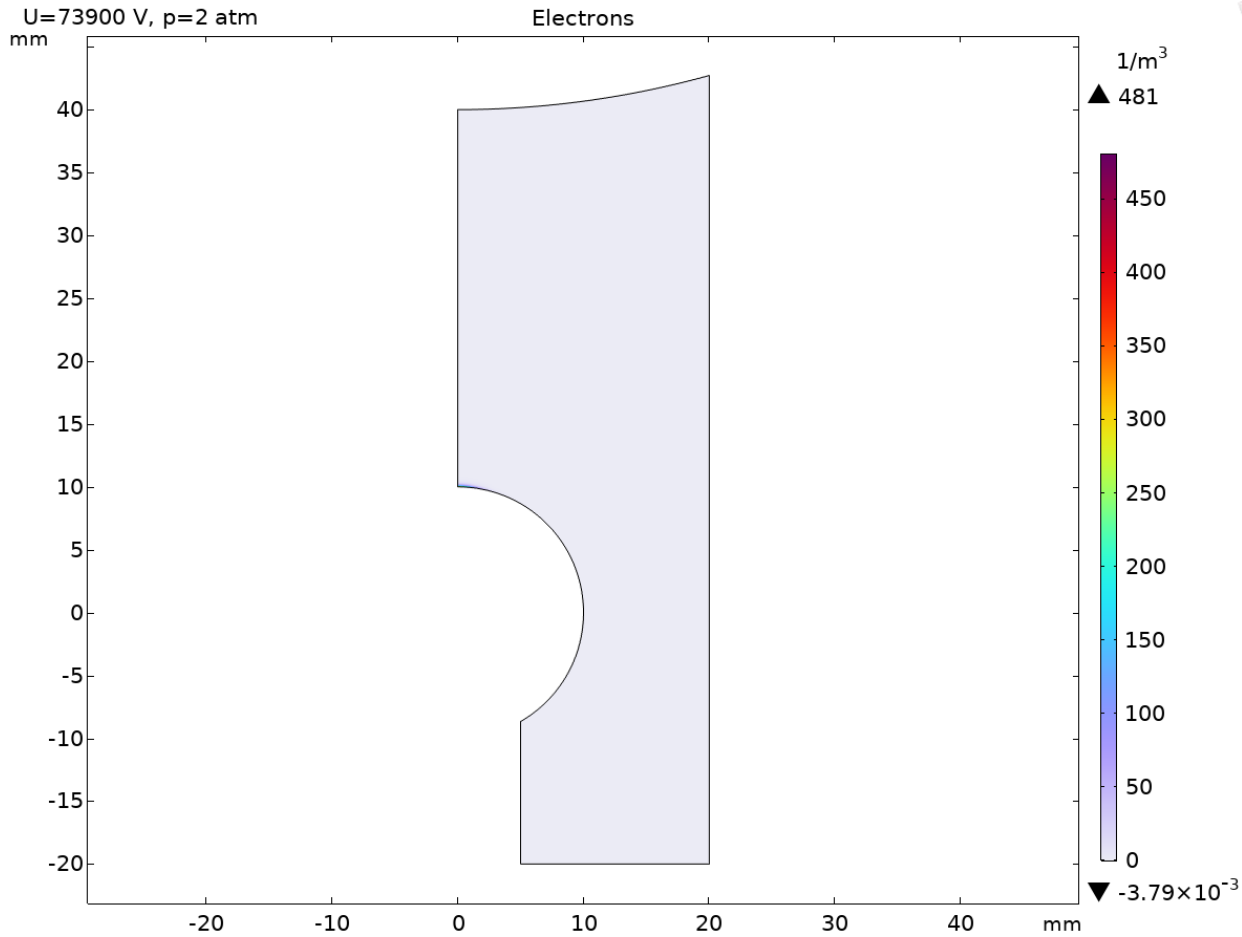


Fig. 17: Concentration distribution of electrons for voltage just below the resonance point for air, $d = 30$ mm – positive (physical) values, created using [1]. (Note that $p = 2$ atm.)

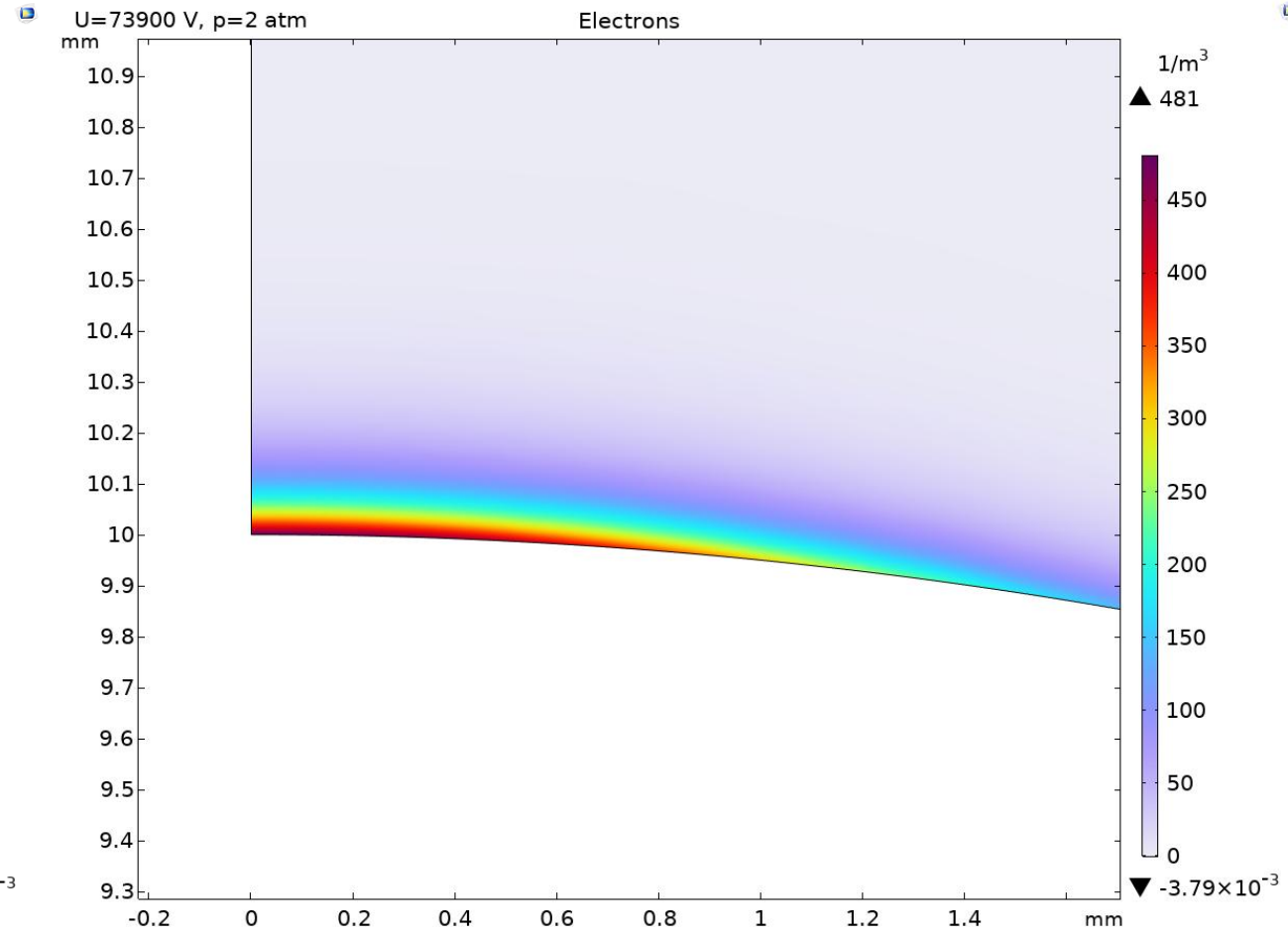


Fig. 18: Zoom of Fig. 17.

The Resonance Method

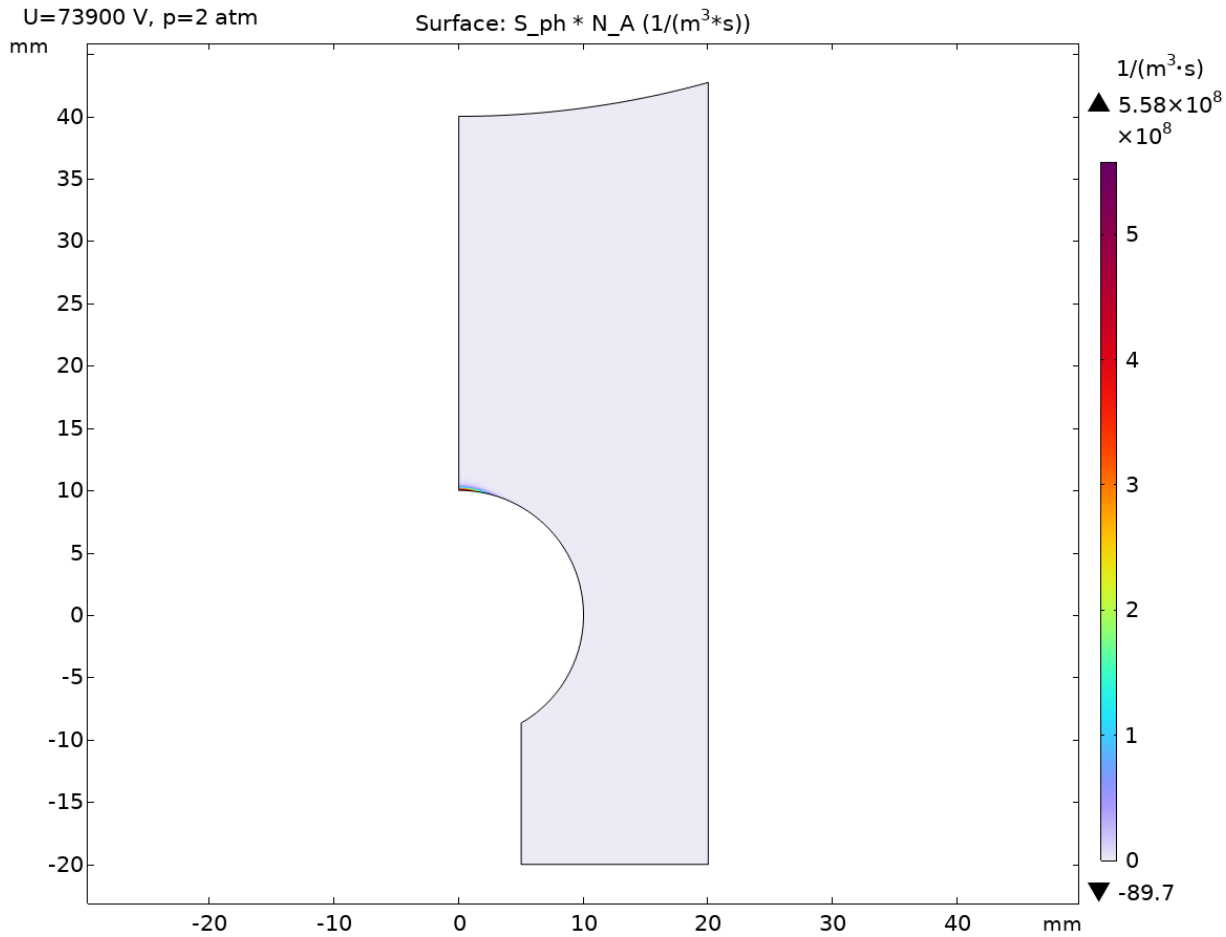


Fig. 19: Distribution of the photoionization rate for voltage just below the resonance point for air, $d = 30$ mm – positive (physical) values, created using [1]. (Note that $p = 2$ atm.)

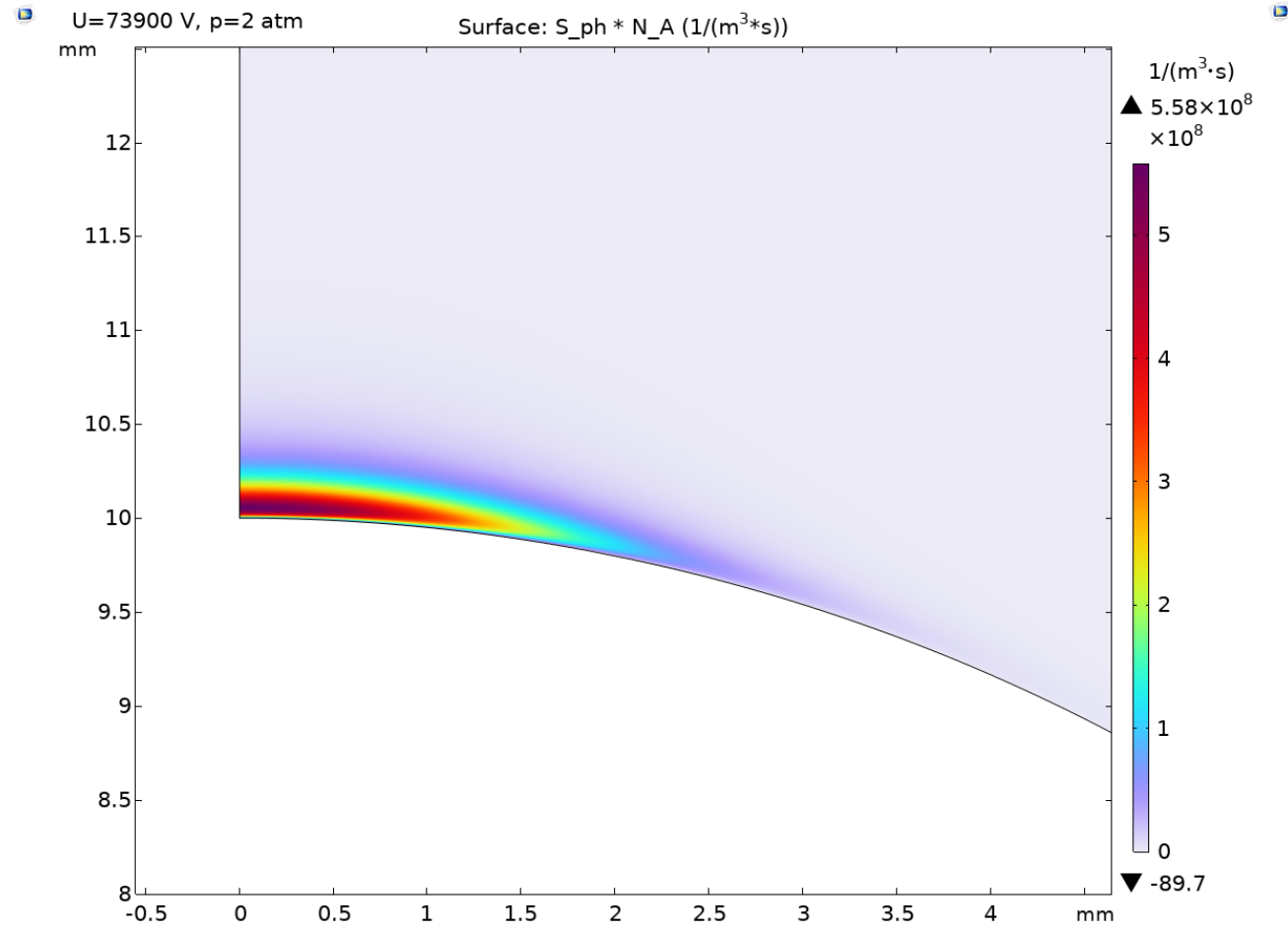


Fig. 20: Zoom of Fig. 19.

The Resonance Method

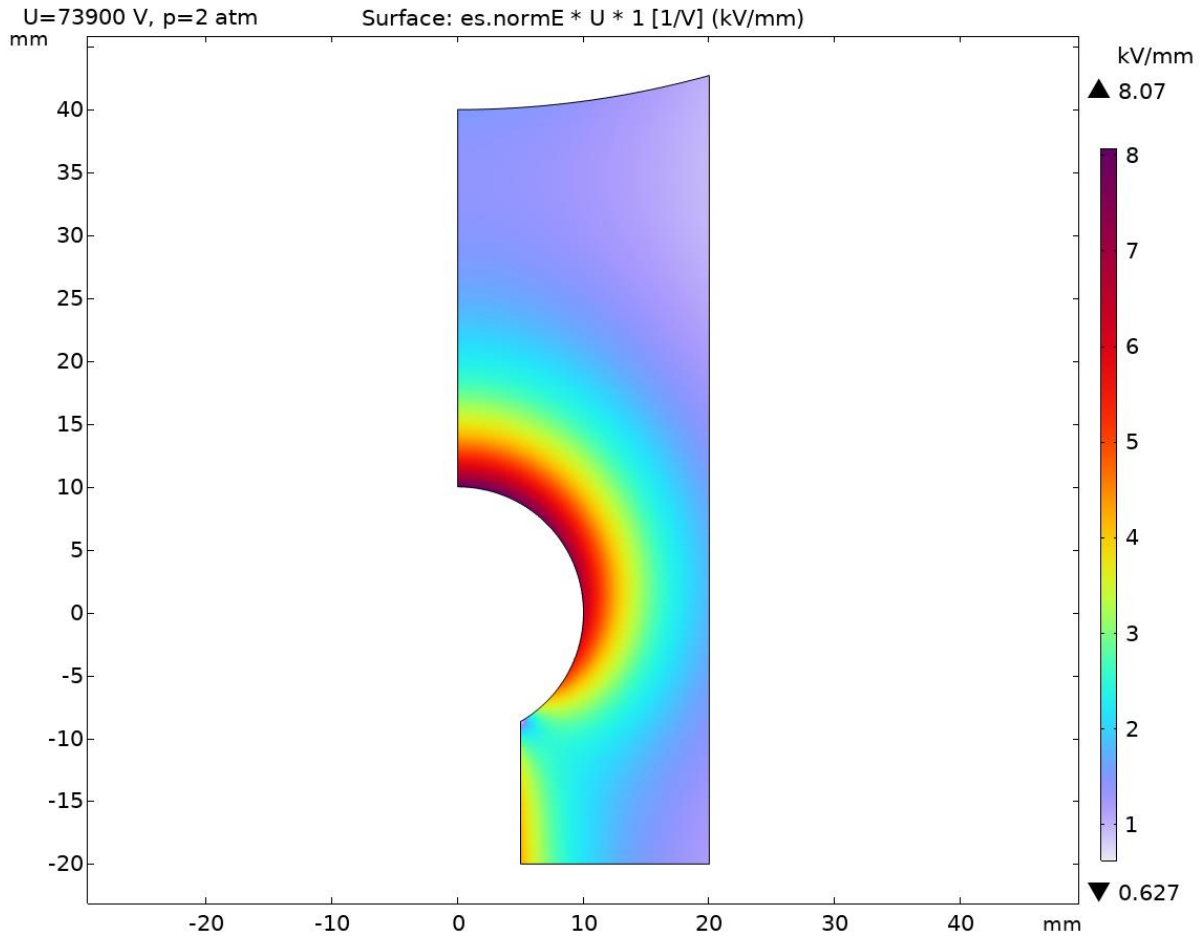


Fig. 21: Electric field intensity distribution for voltage just below the resonance point for air, $d = 30$ mm, created using [1]. (Note that $p = 2$ atm.)

The Resonance Method

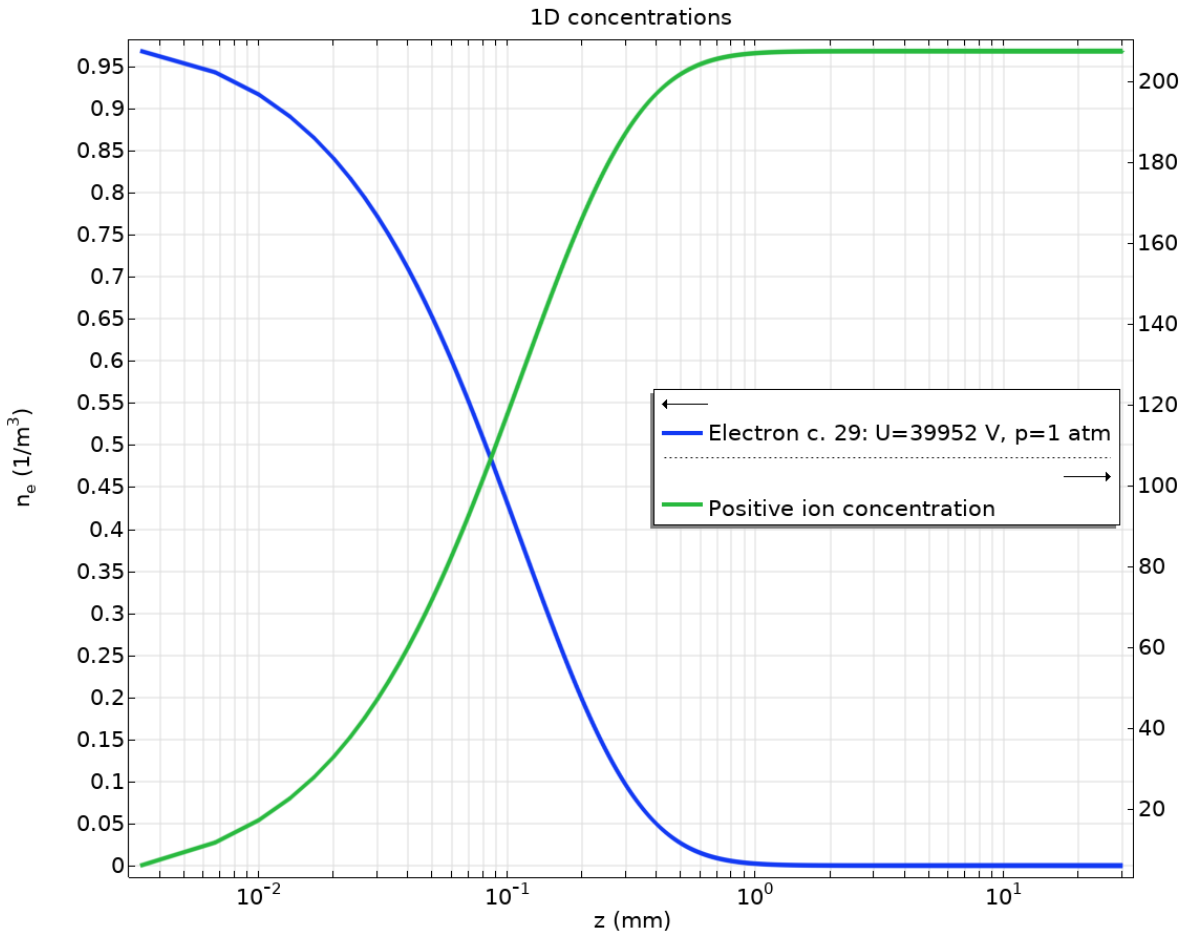


Fig. 22: Concentration distribution of electrons and positive ions along the axis of symmetry for voltage just below the resonance point for air, $d = 30$ mm and $p = 1$ atm – positive (physical) values, created using [1].

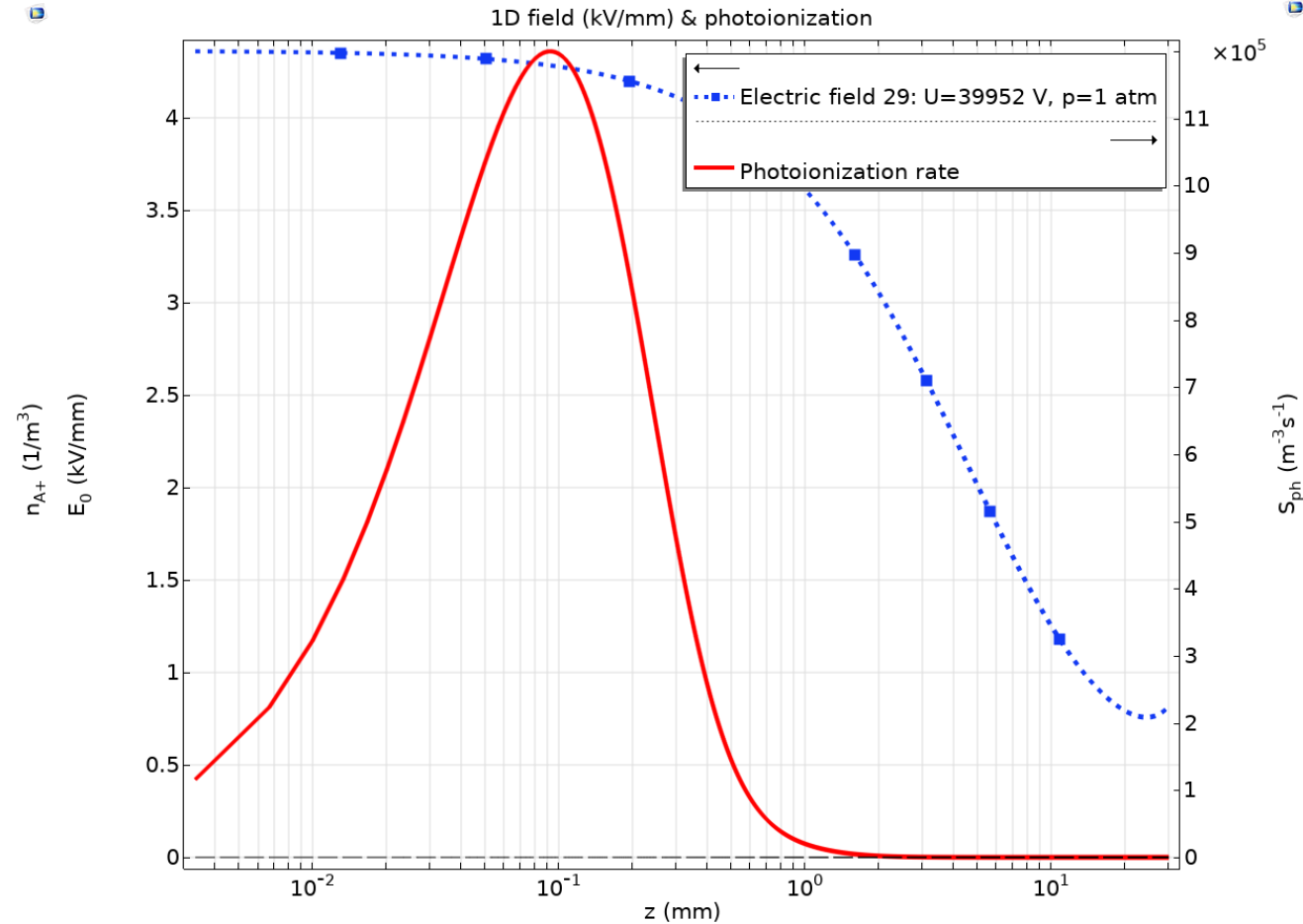


Fig. 23: Distribution of S_{ph} and E along the axis of symmetry for voltage just below the resonance point for air, $d = 30$ mm and $p = 1$ atm, created using [1].

The Resonance Method

- The corona stabilization effects cause a higher breakdown voltage above the calculated U_0
- Corona discharge is localized **self-sustaining** discharge without breakdown
- To calculate breakdown voltage, nonlinear processes must be added:
 - Poisson (with a non-zero right-hand side) and drift-diffusion-reaction equations coupling
 - Nonlinear reactions

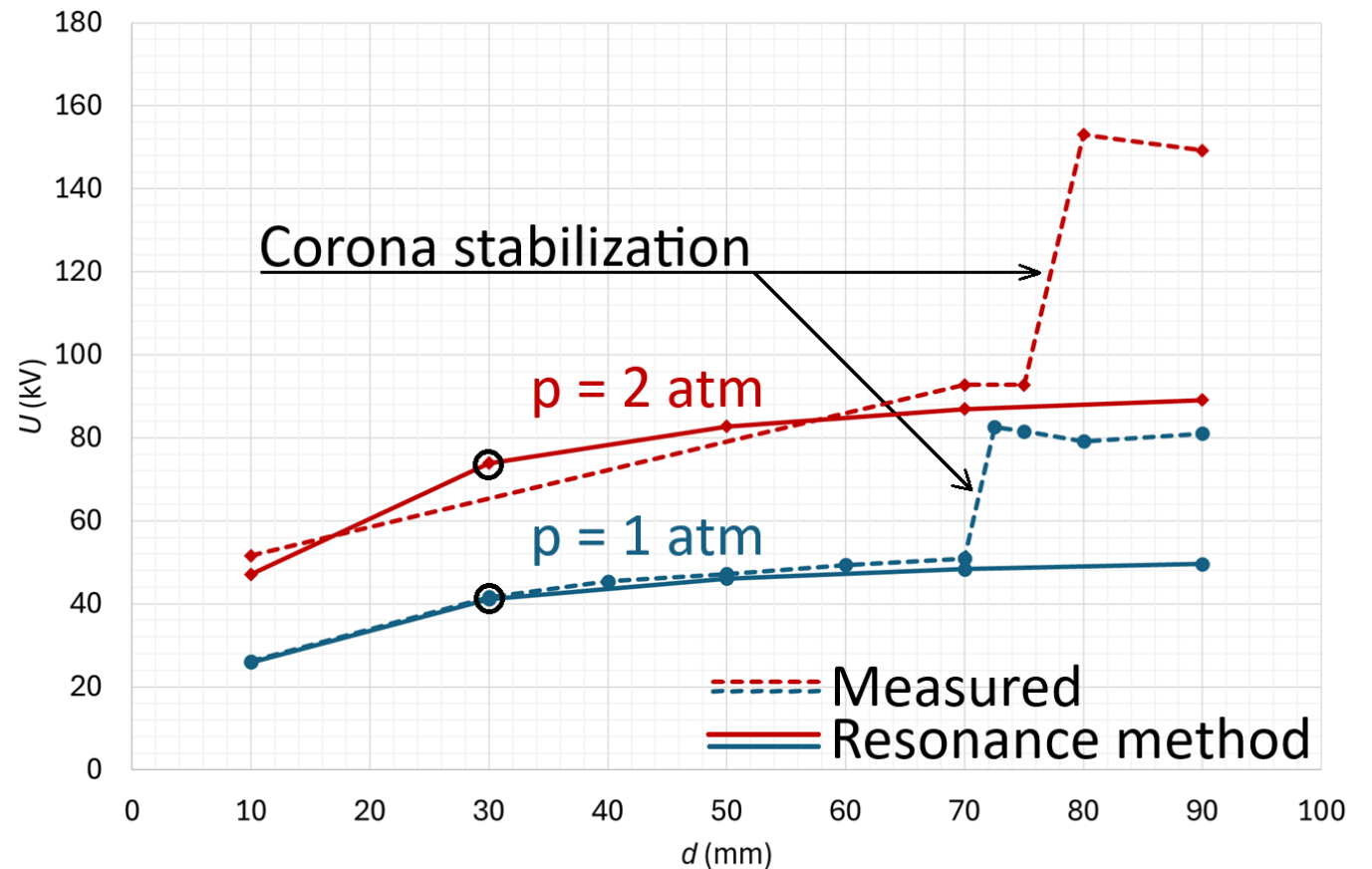


Fig. 24: Calculated self-sustainment voltages U_0 using the resonance method (solid curves) and measured breakdown voltages (dashed curves) as a function of gap distance d for $R_{\text{Sphere}} = 75$ mm and $r_{\text{Sphere}} = 10$ mm and air at $p = 1$ atm (blue) and $p = 2$ atm (red).

Thank you for your attention

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Konference COMSOL Multiphysics 2026

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