

# „Investigation of a Piezoelectric Plasma Generator as a Charging Source for Aerosols”

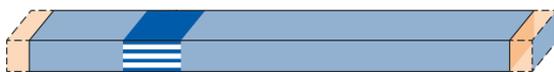
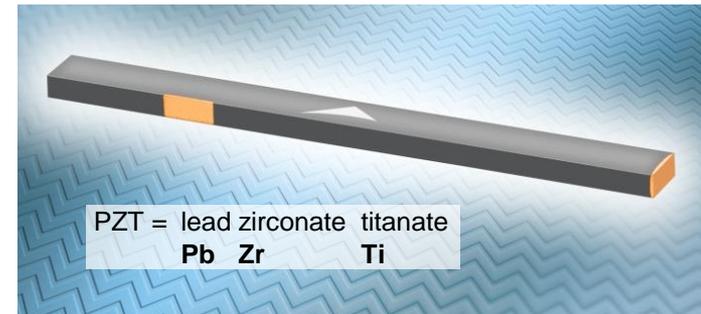
Mario Schrieffl<sup>1</sup>, Alexander Bergmann<sup>1</sup>, Alexander Melischnig<sup>2</sup>,  
Markus Puff<sup>2</sup>

<sup>1</sup> Institute of Electronic Sensor Systems, Graz University of Technology

<sup>2</sup> **EPCOS OHG**, A TDK Group Company

# The CeraPlas™

- TDK developed low-cost piezoelectric cold plasma generator “CeraPlas”
- Properties of the CeraPlas
  - cold plasma generation with ion densities up to  $2E13$  ions/m<sup>3</sup> for each polarity
  - no special plasma generating electrode required
  - no high-voltage plugs or cables needed
  - low plasma temperature (< 50 °C)
  - dimensions [mm]: 45 x 4 x 2.8 or 70 x 6 x 2.8
- Applications
  - surface activation
  - ozone generation
  - sterilization of medical equipment
  - wound disinfection

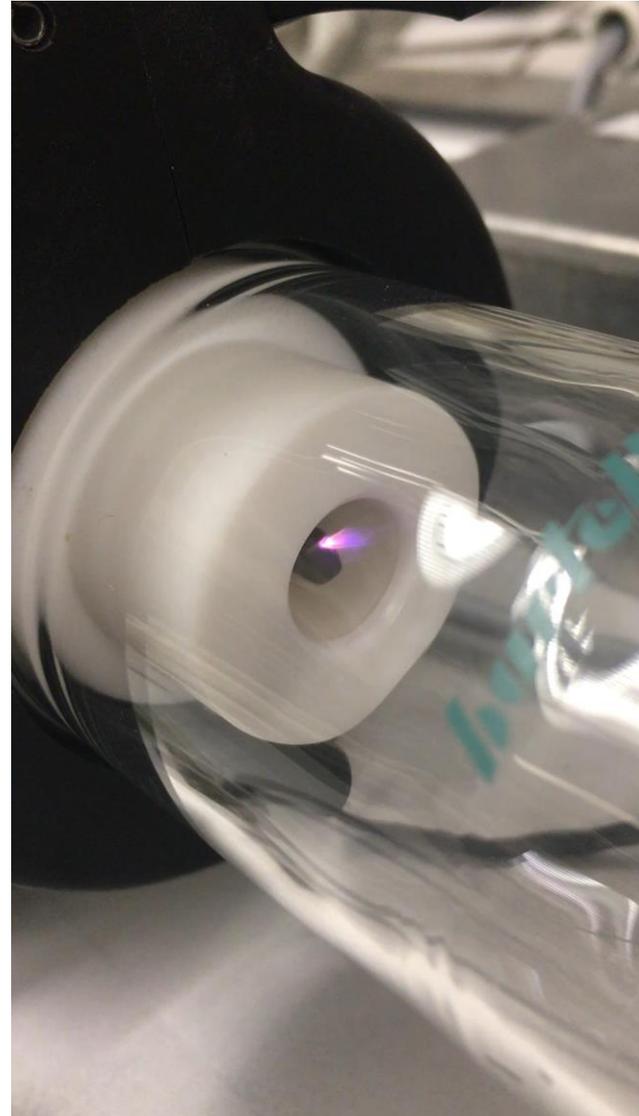


Low input voltage (12V)

High output voltage  
(10 – 15 kV)



# The CeraPlas™



# Can the CeraPlas™ act as ion source for bipolar aerosol charging?

Why is this a relevant question?

- Established neutralizers for aerosol size distribution measurements (radioactive, x-ray, Corona) suffer from some drawbacks
  - ineffective for highly charged or small particles and high particle concentrations
  - permission needed for laboratory usage → expensive
  - high voltage generation (Corona)
- CeraPlas benefits
  - high ion concentrations at low power
  - safe operation
  - low-cost
  - small form factor (comparable to lambda sensor)

# Can the CeraPlas™ act as ion source for aerosol charging?

## How can this question be answered?

- determination of charging parameters based on ion density measurements
- measurement of charge distributions
- tuning of charge distributions to be comparable to reference (Wiedensohler approximation<sup>[1]</sup>)
- Scanning Mobility Particle Sizer (SMPS) spectra: DBD-charger vs. <sup>85</sup>Kr charger

**Ozone-free post-DBD aerosol bipolar diffusion charger: Evaluation as neutralizer for SMPS size distribution measurements**

Rémi Mathon, Nicolas Jidenko, and Jean-Pascal Borra

Laboratoire de Physique des Gaz et des Plasmas, CNRS, Univ Paris Sud, CentraleSupélec, Université Paris-Saclay, Gif-sur-Yvette, France

## Our strategy

- Quick-and-dirty SMPS measurements
  - replace neutralizer by CeraPlas
  - can aerosol particles principally be charged by the CeraPlas?
- Charge distribution measurements
  - charge distribution comparison to reference (<sup>85</sup>Kr, Wiedensohler)?
- Determination of charging parameters
  - measurement of ion densities
  - $N_i t$  product → estimation of steady-state charge distribution
- Tuning of charge distribution
  - tuning of operating parameters to become comparable to reference

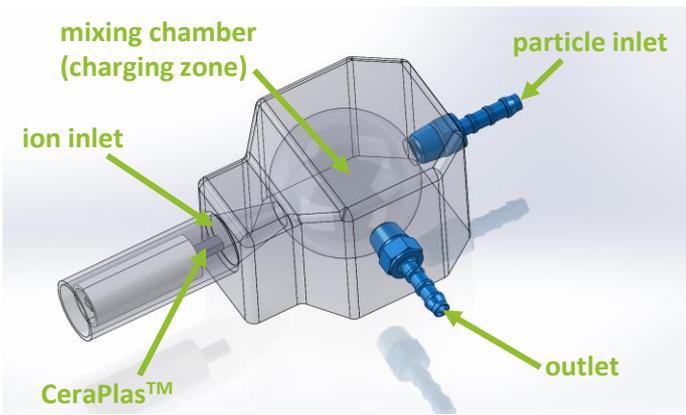
[1] Wiedensohler A., "An approximation of the bipolar charge distribution for particles in the submicron size range," *J. Aerosol Sci.*, vol. 19, no. 3, pp. 387–389, 1988

## Strategy

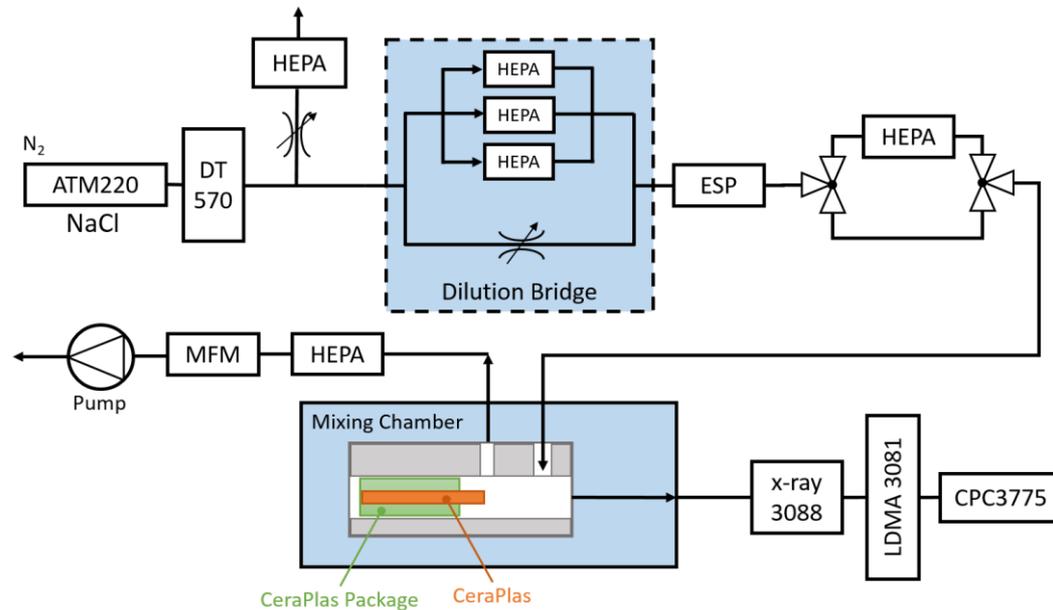
- **Quick-and-dirty SMPS measurements**
- Charge distribution measurements
- Determination of charging parameters
- Tuning of charge distribution

# Quick-and-dirty SMPS Measurements

- CeraPlas was mounted inside a pipe
- A charging chamber was designed for mixing of ions and particles to avoid particle contamination of the device
- With an electrostatic precipitator the charging effect of the CeraPlas was compared to a TSI 3088 x-ray source



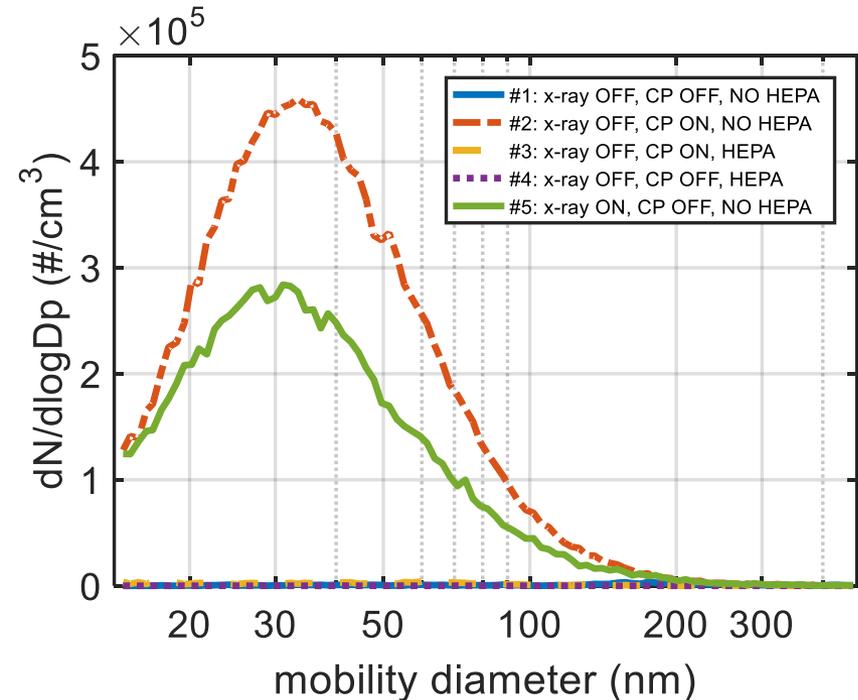
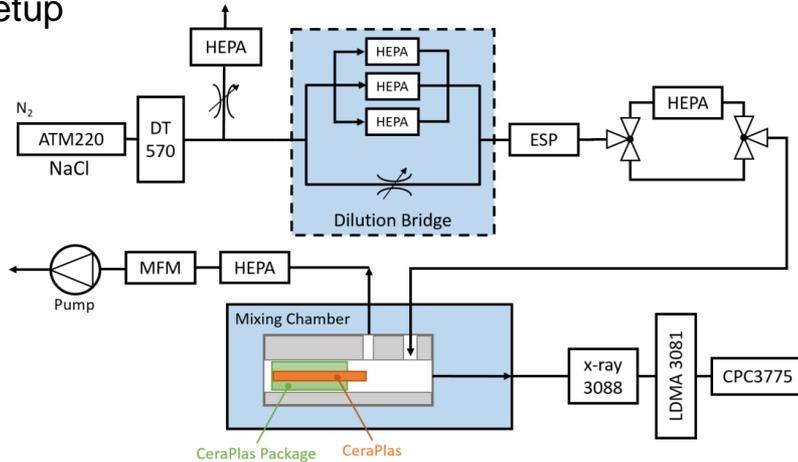
Experimental Setup



# First Results

- No particles “visible” without charging source (#1, #4)
- Charging of neutral particles with CeraPlas (#2)
- Charging of neutral particles with TSI 3088 x-ray neutralizer (#5)
- Particle emissions from device (tested with HEPA filter): no particle contamination from CeraPlas (#3)

## Experimental Setup

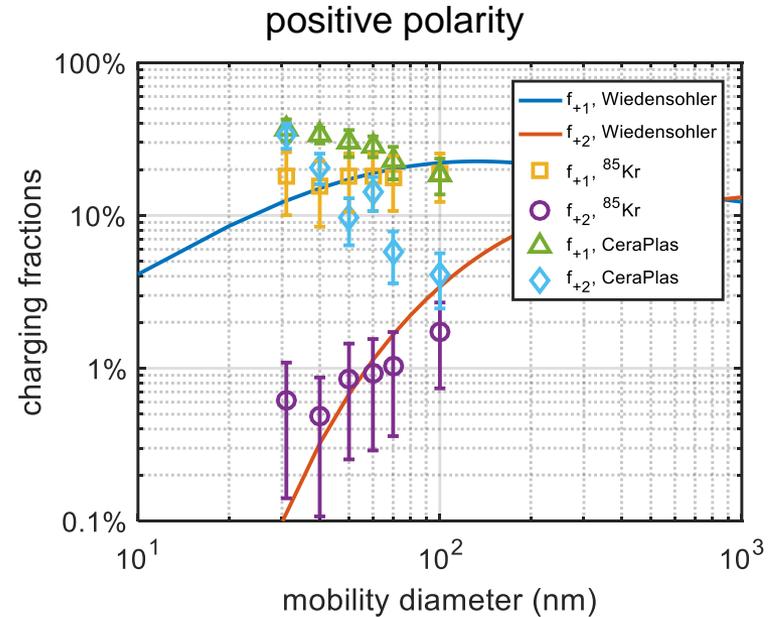
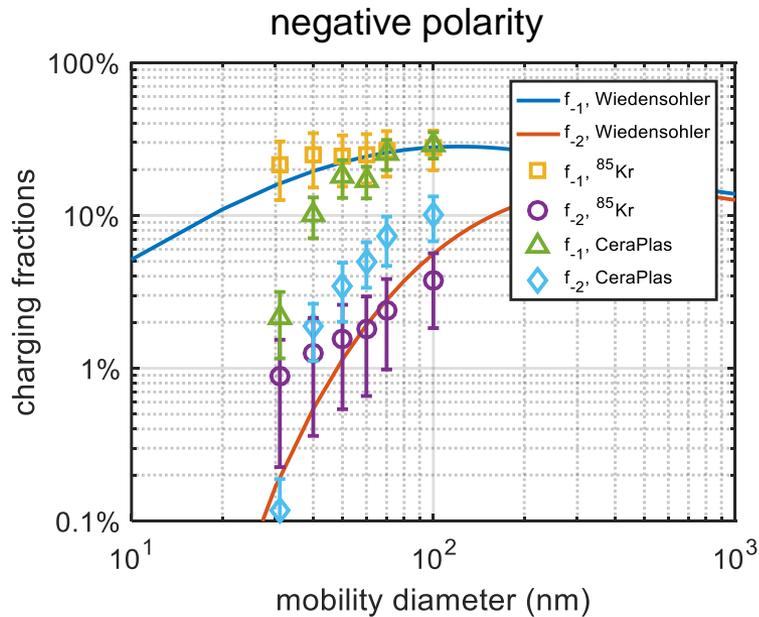


→ Basically it is possible to charge particles using the CeraPlas

## Strategy

- Quick-and-dirty SMPS measurements
- **Charge distribution measurements**
- Determination of charging parameters
- Tuning of charge distribution

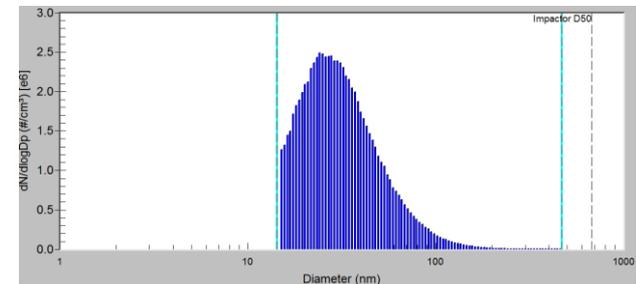
# Results: Charge Distributions CeraPlas vs. Reference



- negative charging fractions show correct trend
- big deviations for positive charging fractions
- for bigger particle sizes results fits better to reference  
→ steady state conditions??

→ investigation of charging parameters!!

Size distribution of test aerosol



## Strategy

- Quick-and-dirty SMPS measurements
- Charge distribution measurements
- **Determination of charging parameters**
- Tuning of charge distribution

# Determination of Charging Parameters

- Ion densities measurements
  - important to characterize CeraPlas
  - input parameter for calculation of  $N_{it}$  product
  - determination of  $N_i/N_p$  which should be  $> 10$  [3]
- $N_{it}$  product
  - product of ion density times the residence time of particles in that ion atmosphere
  - a value of  $6E12 \text{ sm}^{-3}$  necessary to achieve steady state charge distribution which is independent from aerosol concentration and initial charge [2]
  - high  $N_{it}$  product important for reliability and reproducibility of SMPS measurements
- Methods
  - measurement of **inlet ion concentrations** with Ionometer IM806V2
    - $N_i^+ = (3.48E6 \pm 0.05E6) \text{ cm}^{-3}$
    - $N_i^- = (3.15E6 \pm 0.08E6) \text{ cm}^{-3}$
  - calculation of  $N_{it}$  product with multiphysical simulation

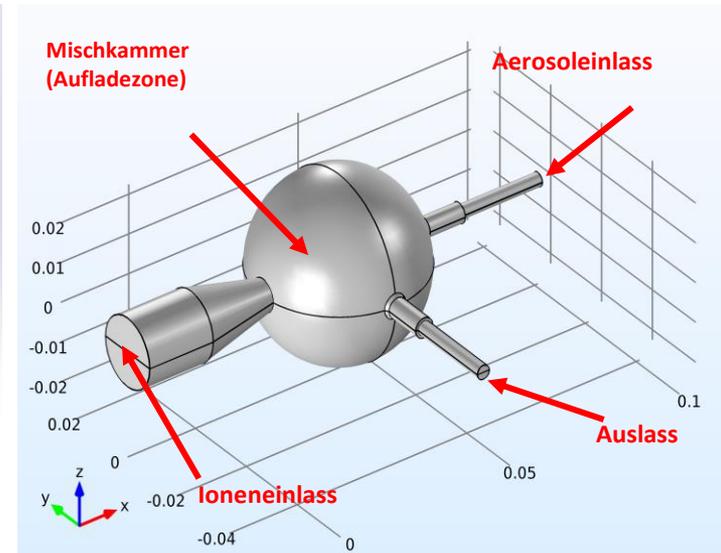
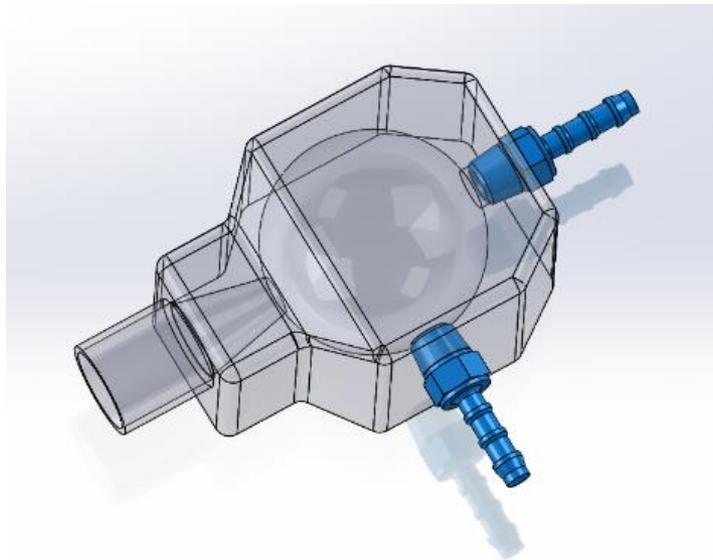


[3] M. Adachi, K. Okuyama, Y. Kousaka, H. Kozuru, and D. Y. H. Pui, "Bipolar Diffusion Charging of Aerosol Particles Under High Particle/Ion Concentration Ratios," *Aerosol Sci. Technol.*, vol. 11, no. 2, pp. 144–156, 1989.

[2]... R. Mathon, N. Jidenko, and J.-P. Borra, "Ozone-free post-DBD aerosol bipolar diffusion charger: Evaluation as neutralizer for SMPS size distribution measurements," *Aerosol Sci. Technol.*, vol. 6826, no. January 2017, pp. 1–10, 2016

# $N_i t$ Product

- ...is the product of ion density times the residence time of particles in that ion atmosphere
- ...is an important charging parameter (indicator of steady state charge distribution)
- normally the  $N_i t$  product is determined via
  - measurement of ion density in charging area
  - calculation of residence time from flow rate assuming plug flow
  - not very accurate
- Idea: determination via 3D model → simulation of mixing chamber



## N<sub>it</sub> Product – Modelling of Mixing Chamber

- Determination of the N<sub>it</sub> product
  - turbulent flow profile within mixing chamber
  - ion density profile: transport of positive/negative ion species

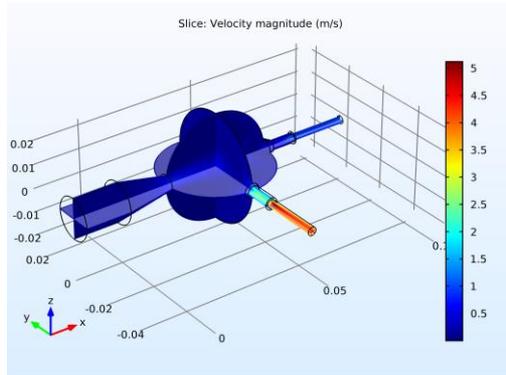
$$\frac{\partial N_i^+}{\partial t} = \nabla(-D_i^+ \nabla N_i^+) + u \cdot \nabla N_i^+ - \alpha N_i^+ N_i^- \quad N_i^\pm \dots \text{pos./ neg. ion density}$$

$$\frac{\partial N_i^-}{\partial t} = \nabla(-D_i^- \nabla N_i^-) + u \cdot \nabla N_i^- - \alpha N_i^+ N_i^- \quad D_i^\pm \dots \text{ion diffusion coefficient (0.357E-5 m}^2\text{/s, [4])}$$

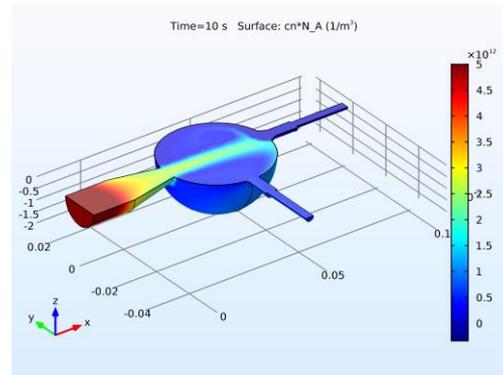
$$\alpha \dots \text{ion recombination coefficient (1.6E-12 m}^3\text{/s [2])}$$

- “typical” particle trajectory: drag force acting on aerosol particle
- calculation of N<sub>it</sub> product by integrating the ion density along particle trajectory

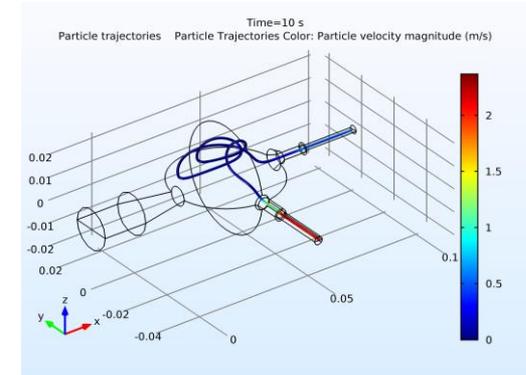
$$N_{it} = \int N_i[\vec{q}_p(t)] dt$$



flow velocity profile (Q<sub>1</sub>=0.3 lpm, Q<sub>2</sub>=1 lpm)



ion density profile (N<sub>i</sub><sup>±</sup>, inlet = 2E13 m<sup>-3</sup>)



particle trajectory, released at midpoint of inlet cross section

[4]... G. Biskos, “Theoretical and experimental investigation of the differential mobility spectrometer,” no. January, 2004

[2]... R. Mathon, N. Jidenko, and J.-P. Borra, “Ozone-free post-DBD aerosol bipolar diffusion charger: Evaluation as neutralizer for SMPS size distribution measurements,” *Aerosol Sci. Technol.*, vol. 6826, no. January 2017, pp. 1–10, 2016

## $N_{i,t}$ Product – Results

- used parameters
  - Flow:  $Q_1=0.3$  lpm,  $Q_2=1$  lpm,
  - ion densities from measurement:  $N_i^+ = 3.48E12$  m<sup>-3</sup>,  $N_i^- = 3.15E12$  m<sup>-3</sup>,
  - particle diameter  $d_p = 100$  nm, number of released particles  $N_p^{rel} = 1000$
- **$N_{i,t}$  Product:**  $N_i^+t = 3.95E+11$  s/m<sup>3</sup>,  $N_i^-t = 3.89E+11$  s/m<sup>3</sup> →  **$N_{i,t} = 4E+11$  s/m<sup>3</sup>**
- Shortcomings of the model
  - diffusion coefficient of ions not known → uncertainty  
(for model hydrated proton ionic clusters with  $D_i^\pm = 0.357E-5$  m<sup>2</sup>/s were assumed)
  - ion recombination coefficient from literature
  - no consideration of other ion loss mechanisms (ion diffusion, space charge, etc.)
- Benefits of the model
  - flow consideration (usually plug flow is assumed)
  - ion density distribution considered
- Conclusions
  - new method of  $N_{i,t}$  product was introduced
  - $N_{i,t}$  values of about  $4E11$  s/m<sup>3</sup> is critically low
  - $N_{i,t}$  product must be increased!!!

## Strategy

- Quick-and-dirty SMPS measurements
- Charge distribution measurements
- Determination of charging parameters
- **Tuning of charge distribution → t.b.d.**

## Summary and Outlook

- Bipolar aerosol particle charging is principally possible with CeraPlas
- Measured charge distributions show strong deviations from reference results
- Method for calculation of  $N_i t$  product was introduced
- Charging conditions ( $N_i t$  product) need to be enhanced

### Next steps:

- Increase  $N_i t$  product
  - increase residence time  $t \rightarrow$  flow / geometry optimization
  - increase ion density  $N_i \rightarrow$  power / positioning of CeraPlas
- Detailed investigation of charge distribution
  - steady-state conditions: is it possible to be comparable to reference?
  - impact of particle concentration / size

Thank you for your attention!!

Questions?????