

Abstract

We characterized size and chemical composition of ions generated by a corona-needle charger of a Neutral cluster and Air Ion Spectrometer (NAIS) by using a high resolution differential mobility analyzer and a time-of-flight mass spectrometer. Our study is crucial to verify the role of corona-generated ions in the particle size spectra measured with the NAIS, in which a corona charger is used to charge aerosol particles down to the size range overlapping with the size of generated ions. The size and concentration of ions produced by the corona discharging process depend both on corona voltage and on properties and composition of carrier gas. Negative ions were <1.6 nm ($0.8 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ in mobility) in all tested gas mixtures (nitrogen, air with variable mixing ratios of water vapour), whereas positive ions were <1.7 nm ($0.7 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$). Electrical filtering of the corona-generated ions and not removing all charged particles plays an important role in determining the lowest detection limit. Based on our experiments, the lowest detection limit for the NAIS in the particle mode is between 2 and 3 nm.

1 Introduction

In aerosol size measurements of sub-5 nm particles based on electrical mobility analysis, the most crucial step is the particle charging process. Electrostatic charging by corona dischargers is commonly used in electrical mobility classification. Different corona charger designs have been developed and tested in various conditions to optimise particle charging efficiency and to minimise losses of ions and particles inside the chargers (Hernandez-Sierra et al., 2003; Unger et al., 2004; Biskos et al., 2005; Qi et al., 2007; Qi et al., 2008; review by Intra and Tippayawong, 2009). The charging efficiency, average charge of particles and penetration of particles through the charger depend on corona voltage, particle diameter and residence time (e.g. Tamm et al., 1992; Hernandez-Sierra, 2003; Biskos et al., 2005; Intra and Tippayawong, 2009, 2010). The size and number concentration of small ions formed inside the charger have

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been characterised for some corona chargers (Tamm et al., 1992; Nagato and Owaga, 1998; Hernandez-Sierra et al., 2003; Alonso et al., 2009). The chemical composition of these ions depends on trace gas compounds and residence time (e.g. Skalny et al., 2004; Nagato et al., 2006; Sekimoto and Takayama, 2007). Experiments with corona dischargers have focused on particles in the diameter range from 4 nm to $\sim 1 \mu\text{m}$ (e.g. Pui et al., 1988; Unger et al. 2004; Tan and Wexler, 2007).

Our focus is to study particle charging down to nanometer sizes with a Neutral cluster and Air Ion Spectrometer (NAIS, Kulmala et al., 2007). Controlled charging of the aerosol sample with a corona-needle charger, together with electrical filtering of the corona-generated ions, enables the detection of both neutral and charged aerosol particles. Although the NAIS is a recently developed instrument, it has been already widely used in field studies of atmospheric particle formation (Manninen et al., 2010; Hirsikko et al., 2011 and reference therein). The NAIS has been calibrated and found to be in good agreement with reference instruments (Asmi et al., 2009). It was concluded that NAIS can be reliably used as the first instrument to measure ions and particles near sizes ($\sim 1.5\text{--}2 \text{ nm}$) where atmospheric particle formation takes place (e.g. Kulmala et al., 2007). The NAIS measures the distribution of both charged and neutral particles in the mobility range $3.16\text{--}0.0013 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$, which corresponds to a mobility diameter range of $0.8\text{--}42 \text{ nm}$ (Mäkelä et al., 1996; Ku and de la Mora, 2009). However, according to Asmi et al. (2009) the ions produced by corona discharge have a mobility range of $1.3\text{--}1.6 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$, which results in a lower detection limit of approximately 2 nm for the NAIS when neutral particles are charged for electrical mobility classification. Particles below mobility diameter $\sim 2 \text{ nm}$ cannot be reliably distinguished from corona-generated ions.

This paper presents a verification of the performance of the corona-needle charger used in the NAIS. Efficient filtering of corona-generated ions but not the charged particles at different operating conditions is essential for electrical detection of charged particles down to as small sizes as possible. The lower detection limit of the NAIS is determined by the charging probability, particle concentration and ion mobilities. To estimate

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how much the corona-generated ions hinder the NAIS particle measurements at the size range overlapping with these ions, we experimentally characterized (1) corona-generated ion number size distribution for different carrier gas mixtures (e.g. filtered compressed air, nitrogen, humidified air, and air mixed with test particles), (2) chemical composition of the ions, and (3) charging of sub-4 nm neutral particles while post-filtering the ions away electrically.

2 Methods and instrumentation

The following paragraphs give an overview of the experimental set-up and instruments used to characterize the performance of the unipolar corona charger used in the NAIS.

2.1 Diffusion corona-needle charger of NAIS

Although the ion spectrometers, such as an Air Ion Spectrometer (AIS, Mirme et al., 2007), detect freshly formed charged particles in the size range well below 1.5–2 nm, the majority of these freshly formed particles are uncharged (Kulmala et al., 2007; Manninen et al., 2010; Lehtipalo et al., 2010). Therefore, measurements of neutral particles below 3 nm are necessary. The NAIS is a modified version of the AIS instrument and includes several improvements: controlled charging of the aerosol sample and the filtering of the corona-generated ions. The NAIS consists of two independent spectrometer columns, one of each polarity, where the ions are classified by a cylindrical Differential Mobility Analyzer (DMA). The NAIS may be divided into two units: aerosol processing and aerosol detection unit.

The processing unit consists of two corona charger and electrostatic filter sections, where the aerosol sample is treated according to the operation mode prior to the mobility analyzers. The aerosol processing unit includes two sections: (1) the discharger + pre-filter, and (2) the main charger + filter section (Fig. 1). The NAIS aerosol processing unit has three operation modes: (1) an offset (zero) mode, when only the discharger

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is used to charge the aerosol sample in the opposite polarity of the analyzer and the pre-filter is used to remove all the charged particles, (2) a particle mode, when only the main charger is used to charge the aerosol sample to the analyzer polarity and the main filter is used to remove ions generated by the charger, and (3) an ion mode, when both sections are switched off.

In the detection unit the charged sample is classified and collected in the 21 electrometer rings in the outer cylinder according to their electrical mobility, and the ion currents are measured with the electrometers. The channels up to 12 are located quite densely on the mobility scale to provide higher mobility resolution at the cluster ion range (negative cluster ions $>1.7 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$, positive $>1.3 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$). The sample and sheath flows of the analyzers are 30 and 60 l min^{-1} . A detailed operation principle and a schematic diagram of the NAIS are presented in Manninen et al. (2009).

The main body of the corona chargers in the NAIS instrument is made of conducting aluminium and Teflon insulator parts. A platinum needle with 0.08 mm diameter and 4 mm length maintained at positive/negative high-voltage is used to produce a positive/negative corona discharge in the carrier gas. The discharge is produced in an electrostatic field established in an air stream between the concentric needle and a grounded cylinder (Fig. 1). The corona discharge causes a very rapid increase in the number of free electrons and ions in the immediate vicinity of the needle. Due to collisions with the free electrons, molecule bound electrons receive sufficient energy to escape from the molecules. The electrons and positive ions move in opposite directions in the electric field, colliding with other molecules and subsequently producing more ions in the corona discharge area. The corona-generated ions and aerosol particles of nanometer sizes collide due to thermal motion and the particles can become charged. This process is called diffusion charging. By monitoring and controlling the ion current arriving at the walls of the cylinder, the ion production can be determined and maintained at a constant level.

2.2 High resolution DMA

The mobility distribution of ions produced in the NAIS corona charger was measured with a high resolution Herrmann Differential Mobility Analyzer (HDMA, Eichler, 1997; de Juan and Fernández de la Mora, 1998; Fernández de la Mora et al., 1998, Herrmann et al., 2000). In the HDMA the aerosol flow rate was approximately 15 l min^{-1} , with a sheath flow of approximately 2000 l min^{-1} . For these operating conditions, the HDMA has an improved mobility resolution with a full width at half maximum (FWHM) of its transfer function of 2% at 1.5 nm (Asmi et al., 2009). The mobility range of the HDMA is $7.5\text{--}0.1 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$, which corresponds to a mobility diameter range of 0.5–4.5 nm. Corona generated ions and charged test particles were classified with the HDMA and detected with an aerosol electrometer (TSI 3068B).

2.3 APi-TOF mass spectrometer

An atmospheric pressure interface time-of-flight mass spectrometer (APi-TOF, Ehn et al., 2010; Junninen et al., 2010) was used to measure the composition of corona-generated ions in both negative and positive polarity. The APi-TOF measures mass spectra of small ions below a few nanometers with minimal perturbation of the sample. The operating mass range of the instrument was roughly 8–2500 Th (corresponding to 0.2–2.25 nm, Ehn et al., 2011). The TOF is the same as used in the Aerodyne high-resolution aerosol mass spectrometer (DeCarlo et al., 2006) and Ionicon PTR-TOF (Jordan et al., 2009). The APi-TOF was connected in parallel with the electrometer, both sampling from the HDMA outflow (Fig. 2).

2.4 Experimental set-up

To examine the performance of the NAIS main charger, we measured simultaneously the size distribution and composition of the ions produced by corona discharge. Figure 2 shows a schematic diagram of the experimental set-up for characterizing the

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ions. The aerosol processing unit (Fig. 1) was disconnected from the NAIS and moved to the experimental set-up. The flow rate through the processing unit was 30 l min^{-1} , corresponding to the sample flow rate of the NAIS. For the sample air mixture we used (1) filtered compressed air, (2) humidified filtered compressed air, and (3) instrument grade (5.0) nitrogen. For preliminary characterisation of ions, the set-up illustrated on the right side of the dashed line in Fig. 2 was used. To test the generation of positive and negative ions in the main charging unit, the DC voltage (corona voltage) was applied to the corona-needle in the range from -5 to $+5$ kV. The efficiency of the electrical filters was tested by varying the voltage applied to the filter electrode (filter voltage) in the range of 0 – 150 V.

For evaluation of the corona charger with test particles, particles in the size range of 1.5 – 5 nm were generated with a WO_x generator (Tungsten Particle Generator, GRIMM 7.860), shown on the left side of the dash-line in Fig. 2. Since the particles exiting the generator are electrically charged, an ^{241}Am (360 MBq) neutralizer together with the electrical pre-filter of the aerosol processing unit were used to obtain neutral test particles. The test aerosols were diluted with filtered makeup air to provide 30 l min^{-1} flow through the aerosol processing unit.

The humidifier used these experiments consists of a 6 mm Gore-Tex tube inside a bigger steel tube with a layer of water between the tubes. A heating cable is wrapped around the outer steel tube. By heating the water between the tubes, more water vapor diffuse to the sample flow inside the Gore-Tex tubing, thus increasing the relative humidity (RH) of the aerosol sample. The RH of the sample flow is measured after the humidifier with an integrated humidity and temperature transmitter (Vaisala Humitter 50U, $\pm 5\%$ RH). A PID controller was used for an active feedback to adjust the heating cable current to set and keep the desired RH.

Finally, we investigated charging of particles in the main corona charger and the filtering efficiency of the ions generated in the corona charger. In the particle measurements with the NAIS, the main filter section removes corona-generated ions which overlap with nanometer sized particles. In other words, the aim is to remove generated

ions but not charged particles. However, the filter removes all charged particles smaller than a certain size (i.e. charged particles with large enough mobility) from the sample air. Therefore, correct adjustment of the main filter is essential for particle detection in the mobility analyzer following the aerosol processing unit.

3 Results and discussion

3.1 Size distribution of corona-generated ions

The size distribution of negative corona-generated ions (Fig. 3) spreads out over a large diameter range having 5 maxima between mobility diameters of 0.98 nm and 1.3 nm ($2.3 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ and $1.2 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$), whereas the distribution of positive ions had two maxima at 1.1 nm and 1.2 nm ($1.7 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ and $1.4 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$). The mean mobility diameter was 1.1 nm ($1.6 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$) for positive ions and 1.0 nm ($1.8 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$) for negative ions. The mean residence time of corona-generated ions was approximately 100 ms between the corona needle tip and the inlet of the DMA (Fig. 2). It should be noted that ion distributions change rapidly on a μs time scale after being produced, and it is difficult to duplicate exact instrument times in the experimental set-up. However, we believe that residence times here are long enough that changes are no longer rapid; thus, our results are applicable to the system we are attempting to characterize. The relatively small ion concentrations measured with our set-up reflect the high flow rate of the aerosol, electrostatic losses in the charging region, diffusion losses in the sample lines, short life time of the ions and the HDMA transmission efficiency. The difference in concentration between polarities is due to higher loss rate of more mobile negative ions. Typically, the number concentration of charger ions varies from 10^4 to 10^7 ions cm^{-3} depending on the applied conditions in each particular case (Hernandez-Sierra, 2003; Intra and Tippayawong, 2010).

We studied the size spectra of ions generated by the corona discharge as a function of applied voltage and polarity. The mobility diameter distributions of negative ions are

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shown in Fig. 4a as a function of corona voltage. All spectra were measured with a fixed flow rate of filtered compressed air through the charger. The negative ion mobility diameter spectra seem to change significantly with the corona voltage: the peak diameter value shifted from 1.2 to approximately 1.0 nm (1.4 and $2.1 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$, respectively).

5 Four peaks in the negative ion distribution were usually distinguished between 1.0 and 1.3 nm (2.1 and $1.2 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$). With low corona voltage the peak diameter is in the middle of this range, and increasing the voltage reveals a tendency of the negative ions to evolve and have only two peaks: at 1.0 nm and 1.3 nm. The positive ion size spectra had three peaks at 1.1, 1.2 and 1.4 nm (1.6 , 1.3 and $1.1 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$), and increasing the corona voltage elevated the total ion concentration but the peak sizes did not change. In general, the negative ions were smaller than 1.6 nm ($0.8 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$) in all conditions when measured with the HDMA, whereas the positive ions were below 1.7 nm ($0.7 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$). These results are in good agreement with the NAIS corona charger experiments by Asmi et al. (2009), and other corona charger experiments (Nagato and Owaga, 1998; Hernandez-Sierra et al., 2003; Alonso et al., 2009).

The efficiency of the electrical main filter as a function of voltage applied to the filter electrode is illustrated in Fig. 4b. The figure suggests that the main filter is able to remove all the small corona-generated ions from the spectra already with a relatively low voltage (± 30 – 40 V). Increasing the filter voltage from 10 V to 20 V decreased the concentration of the ions equally for all the sizes. In normal operating conditions of the NAIS the corona-needle voltage is in the range of 2–3 kV and the filter operates with a voltage of 70–100 V, therefore the ions are removed more efficiently. In these experiments, the pre-filter, which is normally operated during offset measurements, was used to remove all sub-4 nm charged particles with a filter voltage ± 50 V before they reach the main charger region.

3.2 Effect of carrier gas

The presence of water vapour strongly modifies the chemical composition of ions produced by corona discharge (Skalny et al., 2004; Nagato et al., 2006). Therefore, we

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investigated the effect of carrier gas mixture and in particular the role of RH to ion mobility spectra. Also our results indicate that the corona charger ion production rate and the size distribution depend strongly on the particle/gas mixture and on relative humidity (Fig. 5). The negative ions, upper panel in Fig. 5, shifted towards smaller sizes as the RH increased. Concurrently, the positive charger ions, lower panel in Fig. 5, had an opposite behaviour. The positive charger ion size distribution had a peak at 1.1 nm, and as the RH increased, the maximum shifted to 1.25 nm. In addition, the number of peaks both in the positive and negative ions size distribution changes as a function of RH.

The size distributions of positive and negative ions measured in humidified carrier gas (Fig. 5) agree well with the corona-generated ion size distributions observed by e.g. Nagato and Owaga (1998), Alonso et al. (2009), Asmi et al. (2009), and Junninen et al. (2009). Based on their data, positive and negative ions had approximately five and two peaks in the mobility spectra, respectively, when filtered (also dried with silica gel in Alonso et al., 2009) room air was used as the carrier gas. On the contrary, we detected size distributions with two peaks for positive ions and approximately five peaks for negative ions (Fig. 3) when we used dried, filtered compressed (room) air, which had an RH close to zero. However, the location of the peaks seemed to be more or less the same in all studies. Figures 5 and 6 illustrate clearly how the ion production rate, and thus the size distribution, depends on the carrier gas mixture. The blue solid line in Fig. 6 represents the ion size spectra at low relative humidity, whereas the red solid line represents the humidified size spectra. In these two cases the measured size distributions differ significantly.

In our experiments, the residence time of the corona generated ions in the sample flow (between the tip of the corona needle and the inlet of the DMA) was longer than the time the ions drift inside the NAIS. The ions moved in approximately 50–90 ms assuming a plug flow from the main corona charger to the top of the analyzer where the ion concentration is measured. As mentioned, ion composition changes with time (e.g. Nagato et al., 2006), and thus we studied the changes in generated ion size

distributions using flow rates ranging from 28 to 31 l min⁻¹ through the charger. This 10% variation of the flow rate did not significantly change the size distribution, indicating that no rapid changes occurred on these time scales. A shift to larger sizes with decreasing flow was not observed, which also supports the idea presented by Alonso et al. (2009), suggesting that varying the flow rate of air through the charger is not a proper approach to study the effect of ion aging alone.

3.3 Chemical composition of corona-generated ions

We also measured the chemical composition of the ions generated in the corona discharger. In these experiments the mobility of the corona-generated ions was preselected using the HDMA, and the ions were subsequently directed to an APi-TOF for mass spectral analysis. It should be noted that the APi-TOF has an *m/z* dependent transmission which may slightly distort the distributions, and additionally some fragmentation may occur inside the instrument (Junninen et al., 2010; Ehn et al., 2011). For this application, however, these effects are considered small.

Figure 7 shows two typical ion spectra recorded by the APi-TOF: each for positive and negative corona-generated ions. They were sampled in the experiments using filtered compressed air and standard corona charger voltage (± 3 kV). The corresponding ion mobility size distributions are depicted in Fig. 3. The ion spectra presented in Fig. 7 are time averages over several tens of minutes, taken with the HDMA set to a single voltage. For recording the positive ion mass spectrum (top panel), one HDMA voltage was chosen, corresponding to the single large peak found in the mobility size spectrum at 1.1 nm (Fig. 3). The negative ion mass spectrum (bottom panel) is a composite of three single measurements using the HDMA voltages corresponding to the three main peaks of the corresponding mobility size spectrum at 0.98, 1.06 and 1.33 nm (Fig. 3).

The results show that the positive ion spectrum was dominated by peaks at mass/charge (*m/z*) 100 and *m/z* 199, which are believed to have the elemental formulas C₅H₁₀NO⁺ and C₁₀H₁₉N₂O₂⁺, respectively, with the *m/z* 199 peak being the

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dimer (i.e. C_5H_9NO $C_5H_{10}NO^+$) of the m/z 100 peak. The molecular composition of $C_5H_{10}NO^+$ is not known, but could be either protonated piperidone or protonated methyl-pyrrolidone. Ehn et al. (2010) reported that the positive ion spectra in the ambient air samples were dominated by compounds with structures similar to these, e.g. pyridines and quinolines containing a ring of carbon atoms with one carbon atom substituted by a nitrogen atom causing a very high proton affinity. Some identified compounds in laboratory experiments in this study are listed in Table 1. Note the peaks at high m/z have a spacing of 74 Th, i.e. m/z 610, 684, and 758. These compounds arise from silicone polymers, most likely originating from conductive silicone tubing, a common contaminant in these types of measurements.

The negative ion spectrum was dominated by different acids, ionized by donating a proton. The largest contribution is by the nitrate ion NO_3^- at m/z 62 and dimers thereof at m/z 125 (HNO_3 NO_3^-). NO_3^- is a usual terminal ion, also in ambient air, due to its low proton affinity. It clusters also with carbonic acid (H_2CO_3 NO_3^- , m/z 124), and probably other, heavier acids as well. Other acids identified in the negative ion spectra were e.g. pyruvic acid, lactic acid and benzoic acid. A more detailed list is found in Table 1. Compounds with peaks at m/z larger than ~ 200 Th, remain largely unidentified.

The chemical composition of corona-generated ions have been observed to depend on the trace gas composition and time for reactions, as one could expect (e.g. Skalny et al., 2004; Nagato et al., 2006). However, Sekimoto and Takayama (2007) showed that the chemical composition of the ions formed in a corona discharger depends also on the applied voltage.

3.4 Particle charging and filtering of the corona-generated ions

As the ions produced by the corona discharge set the lower detection limit of the NAIS, we investigated filtering efficiency of the ions. First, we generated a polydisperse particle population using a tungsten particle generator, and then the particles were charged in the ^{241}Am neutralizer. Second, the electrical pre-filter (± 50 V) removed all charged

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particles and the main corona section charged the remaining neutral particles. Finally, the main filter of the aerosol processing unit removed small charged particles and the corona-generated ions. The results indicate that the main corona charger used in the NAIS is able to charge small neutral particles in both polarities equally efficiently for electrical classification (Figs. 8 and 9). Figure 8 shows how the size distribution including corona-generated ions and charged WO_x particles shift to larger sizes as the post-filter voltage is increased. Clearly, the figure indicates that the electrical filtering is size dependent.

As shown in Fig. 9, both the ions and the particles charged in the charger were removed by the main filter. The figure shows how positive and negative corona-generated ions of ~ 1.1 and ~ 1.3 nm, respectively, (maximum charger peaks in the size distribution, in Fig. 8) and positively and negatively charged WO_x particles of ~ 3.8 and ~ 3.6 nm, respectively, (maximum WO_x peaks, in Fig. 8) are removed as a function of voltage applied to the filter electrode. The positive and negative corona-generated ions were removed (Fig. 9) when the filter voltage was set to approximately 40–50 V. Although the positive ion concentrations were six times higher at the beginning of the experiment, both negative and positive corona-generated ions were filtered at approximately the same filter voltage. A 50 V a portion of the WO_x particles was also removed. The electrical filtering of the corona-generated ions seemed to be more sensitive to the particle size (mobility) than to the absolute concentration. These results suggest that if we would have generated WO_x particles at 2–3 nm size range the post-filtering with lower voltages would have been more efficient. The dilemma of the post-filter adjusting is how to filter the ions produced by corona discharge without removing the sample particles in the same size range. That voltages used as default values in the NAIS seem to be adequate in filtering out the corona-generated ions and still leaving some of the initially neutral particles at the 2–3 nm size to be detected by the electrometers in the instrument.

4 Conclusions

We present size distribution measurements of both positive and negative ions produced by corona discharge under different conditions. The chemical composition of the ions was measured in parallel with the size measurements. The aim was to characterise both the size and chemical composition of the ions generated by a corona-needle charger from the Neutral cluster and Air Ion Spectrometer (NAIS) with the state-of-art instrumentation. Our results indicate that in the corona discharging process the size and the concentration of ions depend both on the corona voltage and on the physical and chemical properties of the carrier gas mixture. The role of the water vapour is pronounced. The results of this study help in understanding the large scattering of ion mobility values given in the earlier publications (Nagato and Ogawa, 1998; Hernandez-Sierra et al., 2003; Alonso et al., 2009; Asmi et al., 2009).

We characterized the corona-generated ions with a high resolution DMA, which has a narrow transfer function (FWHM of 2% at 1.5 nm), whereas the NAIS transfer functions are quite broad due to low resolution. This has to be considered when determining the lowest theoretical particle detection limit of the NAIS. At normal atmospheric pressure the ratio of sample and sheath flows of the NAIS is 1:2, producing wide transfer functions (Knutson and Whitby, 1975) and spreading the detection of corona-generated ions to several channels (Asmi et al., 2009). Therefore, part of these ions is detected also at bigger sizes by the NAIS. According to our results, the lowest detection limit can be set between 2 to 3 nm, being smaller in the negative and larger in the positive polarity. The limit could be further decreased if the measured small particle concentration is high (when an even larger fraction of the corona-generated ions is used to charge the sample particles).

Further study of the corona discharger and its effect on the NAIS spectra is necessary. Especially, some guidelines for adjusting the electrical filter to remove the corona-generated ions should be given. The results presented in this paper can function as preliminary guidelines until more detailed measurements have been conducted. Opti-

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mally, the corona-generated ions hindering the measurement signal of the NAIS can be accurately determined and subtracted for any given operation condition.

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Table 1. Listing of some major negative and positive compounds in the ion spectra recorded using the APi-TOF. Names are given for the corresponding not deprotonated (negative ions) or protonated (positive ions) compounds. Elemental formulas in italics are not unambiguously identified.

Identified negative ions		Identified positive ions	
Integer <i>m/z</i>	Molecular formula	Integer <i>m/z</i>	Molecular formula
62	NO ₃ ⁻ (nitric acid)	19	H ₃ O ⁺ (water)
87	C ₃ H ₃ O ₃ ⁻	37	H ₂ O·H ₃ O ⁺
89	C ₃ H ₅ O ₃ ⁻ (lactic acid)	55	(H ₂ O)·H ₃ O ⁺
97	HSO ₄ ⁻ (sulphuric acid)	100	<i>C₅H₁₀ NO</i> ⁺
101	C ₄ H ₅ O ₃ ⁻	114	<i>C₆H₁₂ NO</i> ⁺
112	SO ₅ ⁻	130	<i>C₈H₂₀ N</i> ⁺
115	C ₅ H ₇ O ₃ ⁻	171	<i>C₉H₁₉ N₂ O</i> ⁺
121	C ₇ H ₅ O ₂ ⁻	199	C ₅ H ₉ NO·C ₅ H ₁₀ NO ⁺
124	H ₂ CO ₃ ·NO ₃ ⁻	213	C ₆ H ₁₁ NO·C ₅ H ₁₀ NO ⁺
125	HNO ₃ ·NO ₃ ⁻	610	(SiOC ₂ H ₆) ₈ OH ₂ ⁺
127	I ⁻	684	(SiOC ₂ H ₆) ₉ OH ₂ ⁺
129	C ₆ H ₉ O ₃ ⁻ & C ₅ H ₅ O ₄ ⁻	758	(SiOC ₂ H ₆) ₁₀ OH ₂ ⁺
133	C ₄ H ₅ O ₅ ⁻	832	(SiOC ₂ H ₆) ₁₁ OH ₂ ⁺
135	<i>C₄H₇O₅</i> ⁻	906	(SiOC ₂ H ₆) ₁₂ OH ₂ ⁺
149	<i>C₄H₅O₆</i> ⁻	980	(SiOC ₂ H ₆) ₁₃ OH ₂ ⁺

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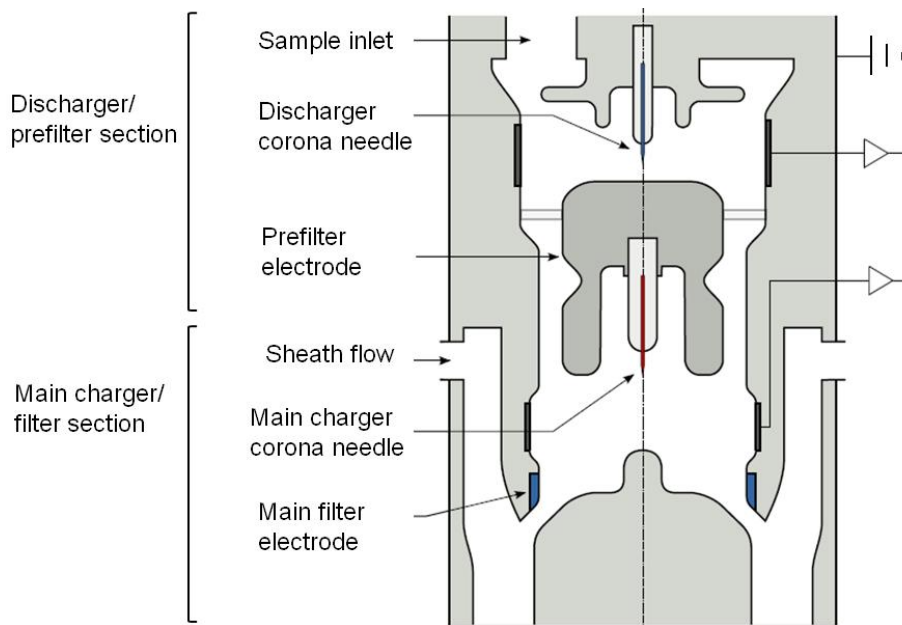


Fig. 1. A schematic cross section diagram of the NAIS aerosol processing unit including two pairs of corona-needle charger and electrical filter sections.

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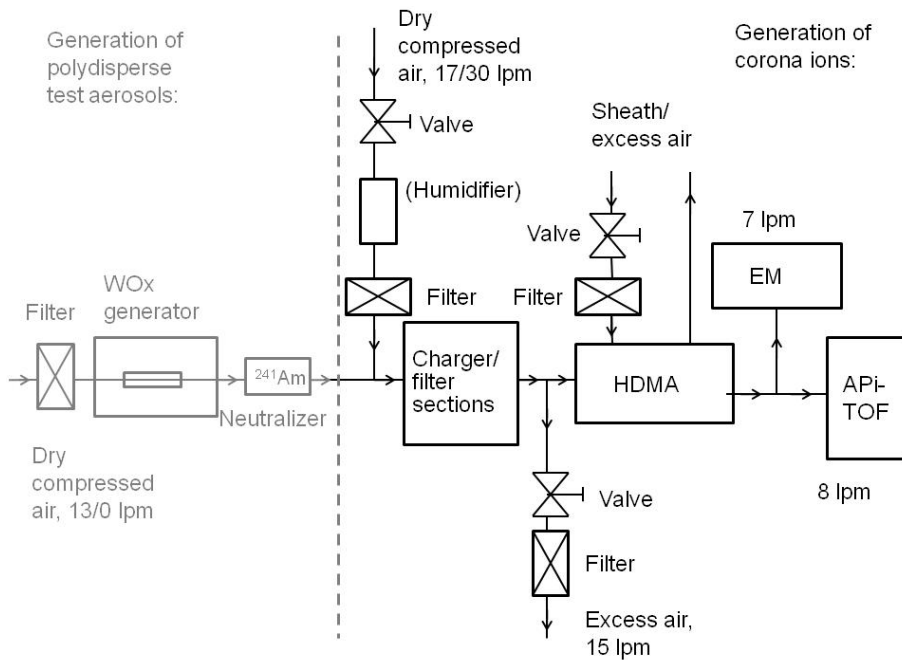


Fig. 2. A schematic layout of the experimental set-up. The layout printed in black was used to investigate the corona-generated ions and the part printed in gray was used when the WO_x particles were introduced to the system to be charged by the corona ions.

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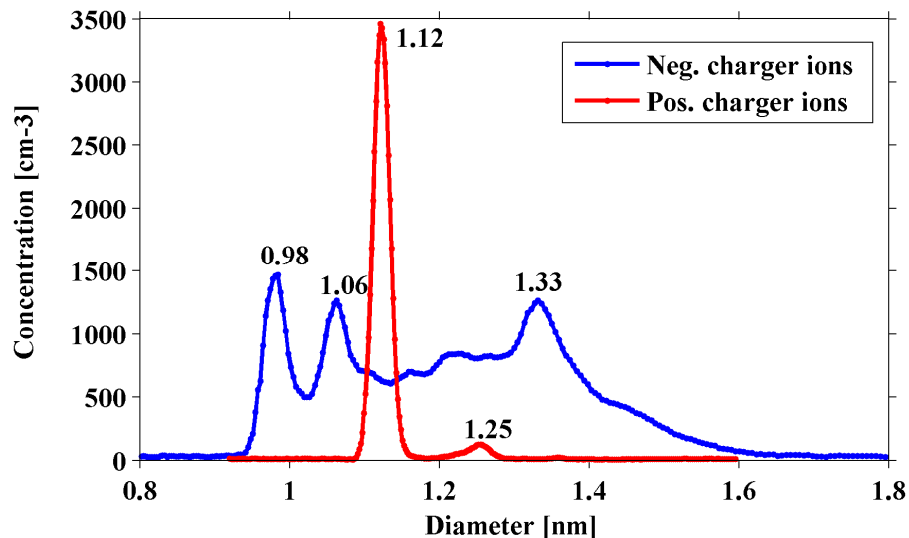


Fig. 3. The mobility diameter distribution of negative and positive ions produced by corona discharge when ± 3 kV was applied to the corona-needle and the carrier gas was filtered compressed air.

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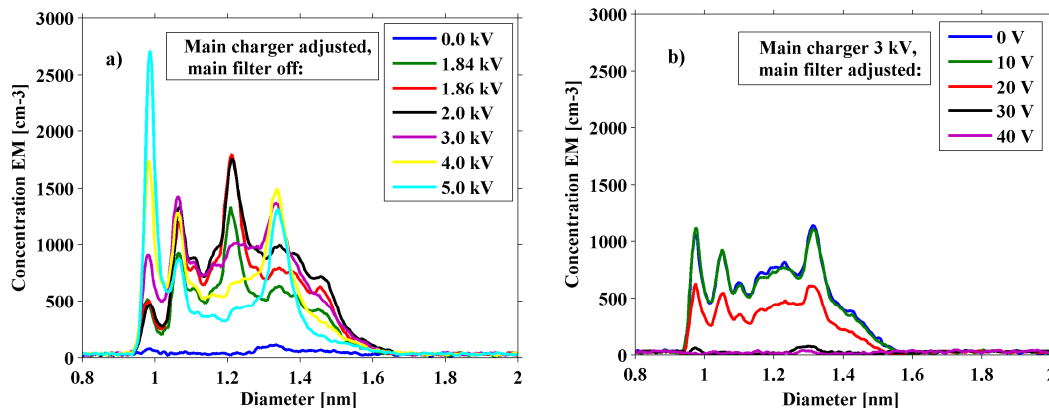


Fig. 4. Size distribution of negative corona-generated ions as a function of (a) negative corona voltage applied to the needle and (b) negative voltage applied to the main filter when corona voltage was kept constant at -3 kV.

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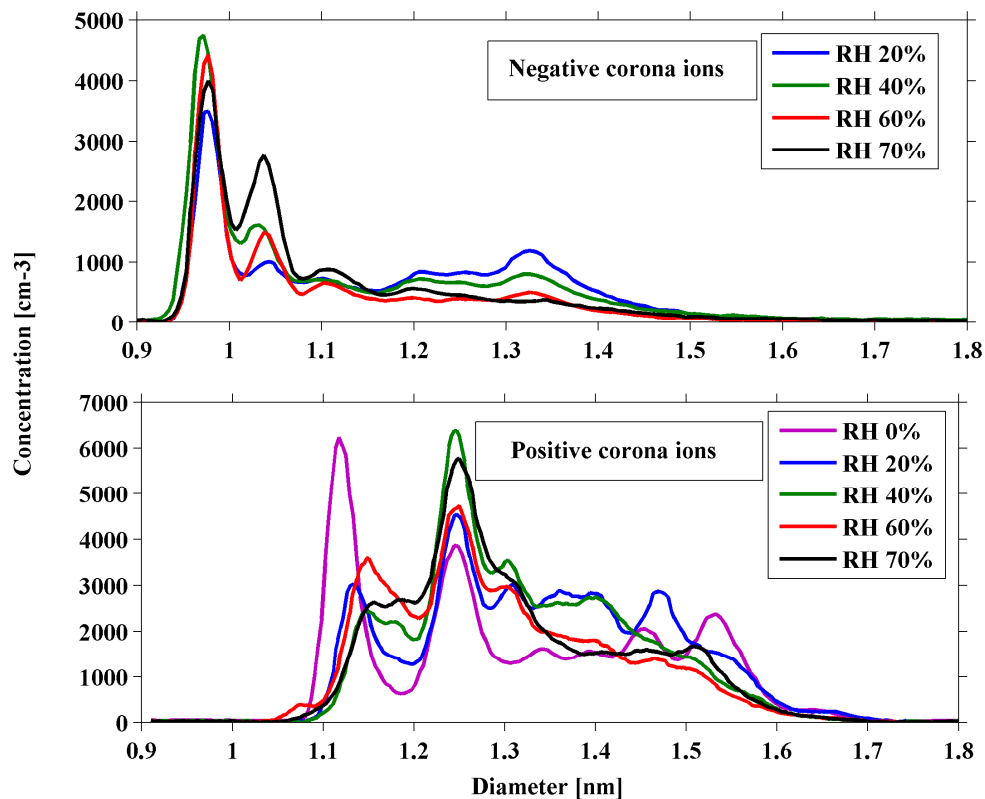


Fig. 5. Size distributions of negative and positive ions generated by the corona discharge as a function of relative humidity (RH). The corona voltage was ± 3 kV.

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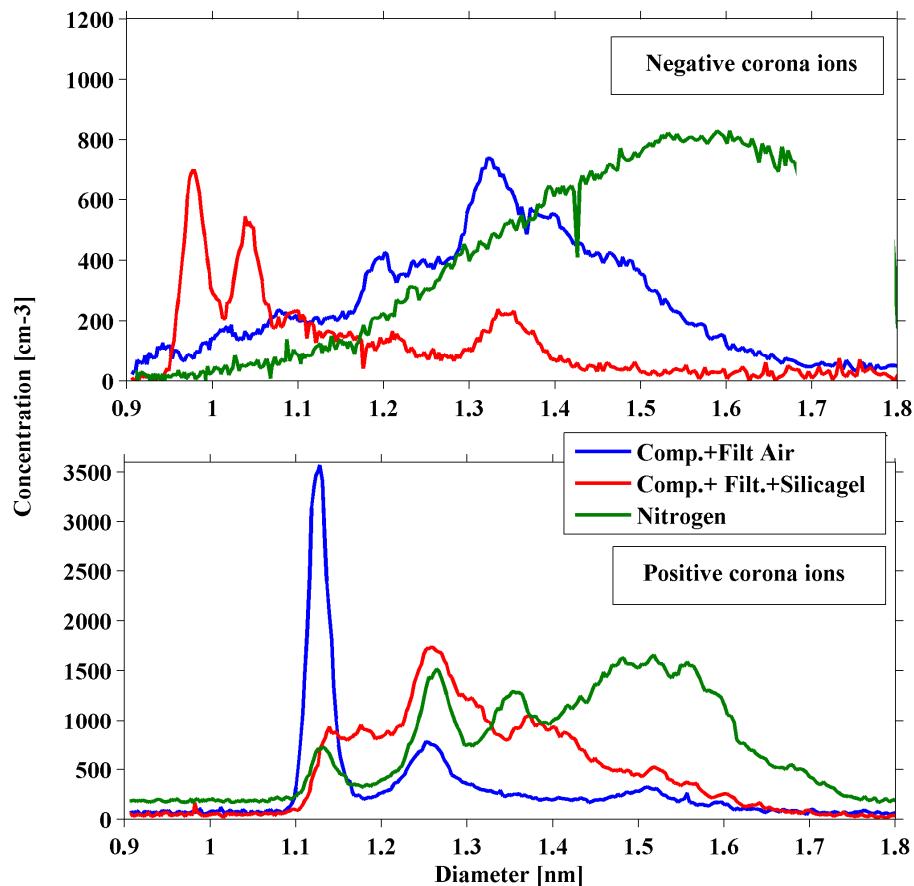


Fig. 6. Size distribution of negative and positive ions generated by the corona discharge with different sample (carrier) gas mixture. The corona voltage was ± 3 kV.

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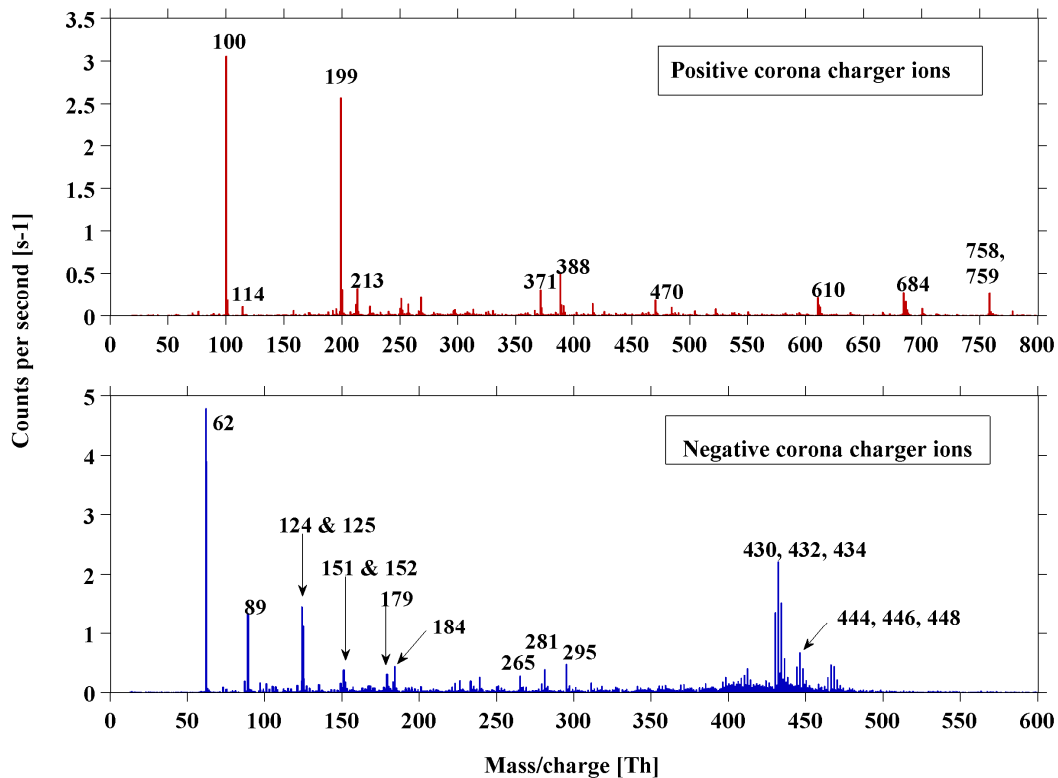


Fig. 7. The mass spectra of negative and positive ions generated by the corona discharge. The corona voltage was ± 3 kV.

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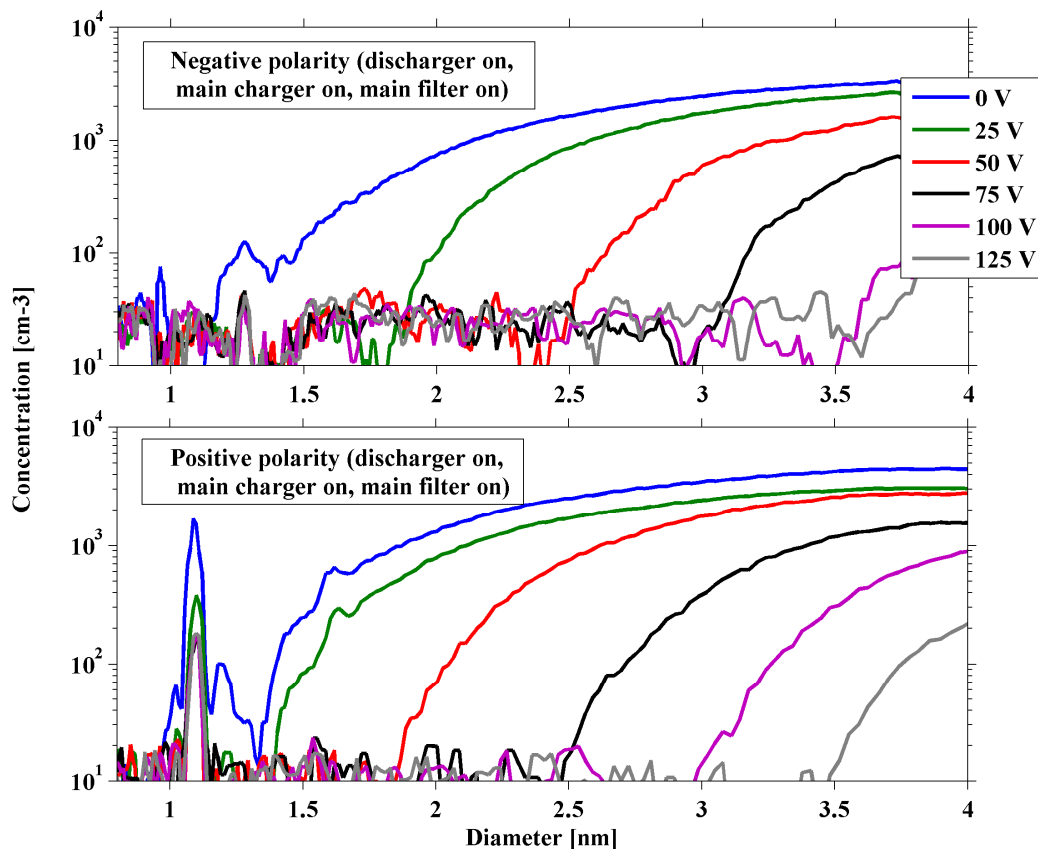


Fig. 8. Filtering of the charged particles with the electrical filter. The panels show the concentration of negative (top) and positive (bottom) corona ions and sample particles as a function of the voltage applied to the main filter electrode.

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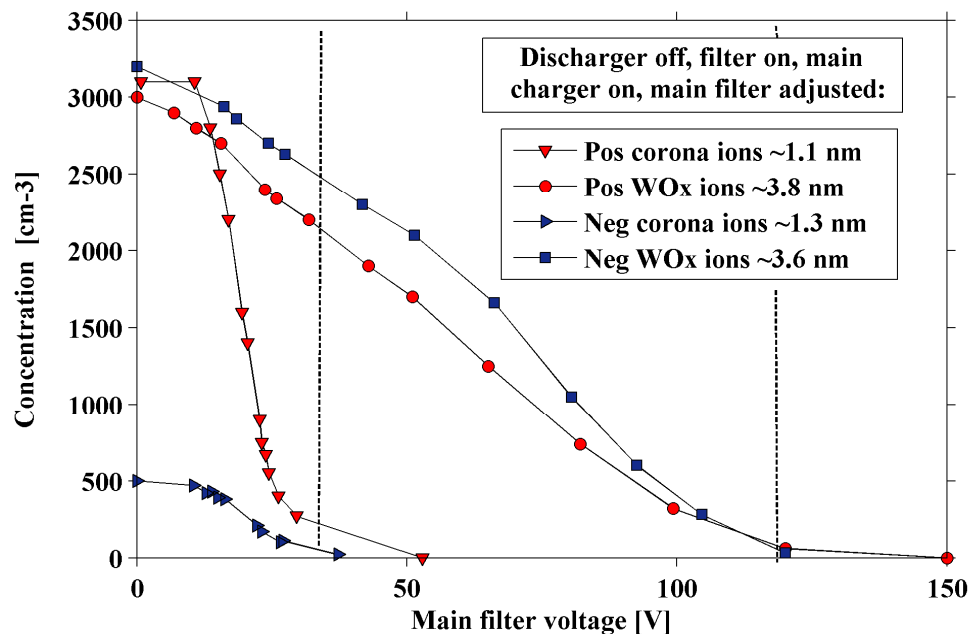


Fig. 9. Filtering of the charged particles with the main electrical filter. The concentration of corona ions and sample particles decreases as a function of the voltage applied to the filter electrode, with most of the charger ions being removed around 30 V. The corona voltage was ± 3 kV and pre-filter ± 50 V.

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