

# Experimental investigation of ion-ion recombination at atmospheric conditions

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**Abstract.** We present the results of laboratory measurements of the ion-ion recombination coefficient at different temperatures, relative humidities and concentrations of ozone and sulfur dioxide. The experiments were carried out using the Cosmics Leaving OUtdoor Droplets (CLOUD) chamber at CERN, the walls of which are made of conductive material, making it possible to measure small ions. We produced ions in the chamber using a 3.5 GeV  $c^{-1}$  beam of positively-charged pions ( $\pi^+$ ) generated by the CERN Proton Synchrotron (PS). When the PS was switched off, galactic cosmic rays were the only ionization source in the chamber. The range of the ion production rate varied from 2 to 100

$cm^{-3} s^{-1}$ , covering the typical range of ionization throughout the troposphere. The temperature ranged from -55 to 20 °C, the relative humidity from 0 to 70%, the SO<sub>2</sub> concentration from 0 to 40 ppb, and the ozone concentration from 200 to 700 ppb. The best agreement of the retrieved ion-ion recombination coefficient with the commonly used literature value of  $1.6 \times 10^{-6} cm^3 s^{-1}$  was found at a temperature of 5 °C and a RH of 40 % ( $1.5 \pm 0.6$ )  $\times 10^{-6} cm^3 s^{-1}$ . At 20 °C and 40 % RH, the retrieved ion-ion recombination coefficient was instead ( $2.3 \pm 0.7$ )  $\times 10^{-6} cm^3 s^{-1}$ . We observed no dependency of the ion-ion recombination coefficient on ozone concentration and a weak variation with sulfur dioxide con-

centration. However, we observed a more than fourfold increase in the ion-ion recombination coefficient with decreasing temperature. We compared our results with three different models and found an overall agreement for temperatures above 0 °C, but a disagreement at lower temperatures. We observed a strong increase in the recombination coefficient for decreasing relative humidities, which has not been reported previously.

## 1 Introduction

Air ions are fundamental to atmospheric electricity and play a central role in the proposed connection between solar activity, Galactic Cosmic Rays (GCRs) and climate (Israël, 1970; Carslaw et al., 2002; Usoskin and Kovaltsov, 2008). Ions are known to enhance nucleation rates in atmospherically relevant vapor mixtures (Kirkby et al., 2011). In particular, ion-ion recombination has been proposed and studied as the driving force behind atmospheric nucleation (Yu and Turco, 2008; Yu, 2010; Nagato and Nakauchi, 2014). However, the overall effect of ions on atmospheric new particle formation (NPF), and subsequent production of cloud condensation nuclei, has remained a controversial issue (Gagné et al., 2008; Kazil et al., 2010; Manninen et al., 2010; Yu et al., 2010; Hirsikko et al., 2011; Kontkanen et al., 2013; Kulmala et al., 2013). Air ions accumulating near cloud edges may affect cloud microphysics and ultimately climate via several mechanisms that are currently poorly quantified (Tinsley, 2000; Harrison and Ambaum, 2008).

Atmospheric ions are usually classified in three groups depending on their diameter (here, all diameters are reported as Millikan-Fuchs equivalent mobility diameters, Mäkelä et al., 1996): small ions (<1.9 nm), intermediate ions (1.9 - 7.7 nm) and large ions (>7.7 nm). In terms of mobility, they are classified as: small ions ( $>0.57 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ ), intermediate ions ( $4.3 \times 10^{-2} - 0.57 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ ) and large ions ( $<4.3 \times 10^{-2} \text{ cm}^2 \text{ V}^{-1}$ ) (Hörrak et al., 2000). Small ions have a high mobility, therefore they are the main responsible for the transfer of charge in the atmosphere (Chalmers, 1949; Ogawa, 1985) and play a role in NPF and in aerosol charging processes (Adachi and Kousaka, 1985; Manninen et al., 2011). This work focuses on small ions, from here on simply referred to as “ions” unless specified otherwise.

Air ions are continuously produced in the atmosphere from GCRs and terrestrial sources, such as radon decay and gamma radiation from the soil (Laakso et al., 2004). Within the planetary boundary layer, terrestrial sources play an important role in ionization processes, whereas at altitudes greater than 2 km, GCRs are the dominant source of ions (Fig.A.1 Harrison and Carslaw, 2003; Kazil and Lovejoy, 2004; Usoskin et al., 2004; Arnold, 2008; Zhang et al., 2011; Williams et al., 2011; Harrison et al., 2014).

When ionizing radiation hits an air molecule, an ion-electron pair is formed. The primary positive ion can be a molecular ion (e.g.,  $\text{N}_2^+$ ,  $\text{O}_2^+$ ), but it can also be  $\text{N}^+$  or  $\text{O}^+$ , as the ionizing radiation is very energetic (Smith and Spanel, 1995; Volland, 1995; Leblanc et al., 2008). After the interaction between an air molecule and ionizing radiation, the newly formed positive ion stays in Brownian motion around its initial position. Its diffusion coefficient is similar to the one of the surrounding molecules and its change in momentum is negligible.

The electron, however, gains a finite momentum in the interaction. If the energy gained is sufficiently high, the electron starts to interact with atmospheric molecules, exciting and ionizing them, until its energy decreases to thermal levels and it binds to an electronegative molecule (for example an electron and an  $\text{O}_2$  molecule, combining to form  $\text{O}_2^-$ ). This process takes place in a few nanoseconds.

Once molecular ions of opposite polarity are created, they can interact with molecules that have higher electronegativity or proton affinity (e.g.,  $\text{H}_2\text{O}$ ) and transfer their charge. Alternatively, they can establish hydrogen or other chemical bonds with other molecules and cluster (Smith and Spanel, 1995). Positive ions are transformed into hydronium-water clusters and ammonium-water clusters, reaching their equilibrium concentrations after about 10  $\mu\text{s}$  (Luts and Salm, 1994; Luts, 1998). After 1 s, less abundant trace gases (such as pyridine and dimethylamine) start to be part of the clusters.

Negative primary ions (mainly  $\text{O}_2^-$ ) cluster with water molecules in 0.1  $\mu\text{s}$  and reach their equilibrium around 0.1 s after the interaction between air and ionizing radiation. Once at equilibrium reactions with  $\text{NO}$ ,  $\text{NO}_2$  and  $\text{O}_3$  start to create nitric acid-water clusters. If present, iodine is known to cluster at longer time scales (several seconds) (Luts and Salm, 1994; Luts et al., 2011).

The number concentration of ions depends on the balance between ion sources and sinks. The production rate is proportional to the amount of ionizing radiation present, whereas the loss mechanisms depend on the attachment rate of ions onto aerosol or macroscopic surfaces and on the ion-ion recombination rate (Tammet et al., 2006). The ion-ion recombination rate is the rate at which one ion collides with another ion of opposite charge and get neutralized. It depends on the mobility of ions and possibly on their chemical composition and ambient conditions.

Ion-ion recombination becomes dominant in extremely clean environments, where the integral of the aerosol surface distribution is negligible with respect to the ion-ion recombination rate, and the probability for ion-aerosol attachment is low (Volland, 1995). The attachment coefficient for an ion attaching to a neutral aerosol particle ranges from 0.1 to  $2.0 \times 10^{-6} \text{ cm}^3 \text{ s}^{-1}$ , for an ion of 1.1 nm and an aerosol particle ranging from 10 to 100 nm mobility equivalent diameter (Tammet and Kulmala, 2005).

Recombination is also important at high ionization rates, when the production of ion pairs is so high that the proba-

bility of colliding with an aerosol particle is minimal with respect to the probability of colliding with an ion of opposite charge.

In the past, Bates and Flannery (1969), defined the equations to describe ion-ion recombination as a sink for air ions, continuing the theoretical work initiated by Thomson (1924) and continued by Natanson (1960). Hoppel and Frick (1986) studied the theory of ion-aerosol attachment, and its limiting case, ion-ion recombination, with the aim to enable the use of differential mobility analysis to measure the aerosol population. McGowan (1965) studied ion-ion recombination in laboratory air to improve dosimetry techniques for ion chambers. Gringel et al. (1978) measured vertical profiles of air conductivity and showed good agreement up to 20 km altitude, using a theoretical recombination coefficient of small ions based on a three-body recombination process. Lee and Johnsen (1989) investigated ion-ion recombination in helium and argon at atmospheric densities and found that, in their setup, along with three-body recombination, stabilization by mutual neutralization played an important role. Heptner et al. (2012) conducted experiments to study relative changes in ion-ion recombination in dry filtered air at atmospheric pressure. In most of these cases, the ions were studied under conditions that were either far from those found in the atmosphere, or allowed little control over trace gas concentrations and ion production rates. In this work, we present the results of laboratory experiments performed at conditions close to those found in the Earth's atmosphere, providing quantitative results about the ion-ion recombination coefficient at varying temperature, relative humidity and trace gas concentrations.

## 2 Experimental methods

The typical experiment carried out consisted of measuring the ion concentrations at several beam intensities, each time for long enough (about 30 min) to reach steady state conditions while all the other variables were kept constant (Figure A.3). The ion concentrations were averaged over the last 10-15 minutes, before changing the beam intensity. The averaging time allowed us to compensate for occasional fluctuations in the ion concentrations that were caused by beam pulses, or by convective transport due to the two fans stirring the air inside the chamber. The beam intensity was varied from 0 to  $1.5 \times 10^5$  pions  $s^{-1}$ , corresponding to an ion production rate ranging from about 2 to 100 ion pairs  $cm^{-3}s^{-1}$ . The concentration of aerosol particles was constantly monitored with a CPC (TSI 3776) and it remained below  $30 cm^{-3}$  in all experiments. The concentration of ions with  $D_p > 1.9$  nm was negligible as the experiments were carried out in an aerosol-free chamber. Usually, the beam intensity was increased consecutively, generating a series of steps of ion production rate. Each time, at the end of the last step, the beam shutter was closed, and we observed the ion concentration decay (Figure

A.4). The mobility range of the ions considered in this study was from  $3.2$  to  $0.57 cm^2V^{-1}s^{-1}$ , that corresponds to a size range of  $0.8 - 1.9$  nm in mobility equivalent diameter ( $D_p$ ). In this analysis the signal of the NAIS in ion mode was integrated over all the channels corresponding to  $D_p < 1.9$  nm. The rest of the ion number size distribution was not considered, as it was negligible. From this dataset, we retrieved the ion-ion recombination coefficient under different conditions.

### 2.1 The CLOUD Chamber

The Cosmics Leaving Outdoor Droplets (CLOUD) chamber (Kirkby et al., 2011) is a cylindrical vessel with a diameter of 3 m and a volume of  $26.1 m^3$  made of electro-polished 316L stainless steel (Figure A.2.c). Its walls are conductive, which makes it possible to measure small ions that, in a traditional aerosol chamber made of polytetrafluoroethylene (PTFE), would be removed in less than one second by the parasitic electric fields created by the dielectric material (McMurry and Rader, 1985). In the interior of the chamber, all plastic components are avoided using copper o-rings and metallic coating on electrically insulating components. When it is necessary to work in an ion-free environment, two circular grids, one located at the top and one at the bottom of the chamber, are provided with a potential difference of up to 60 kV, generating an axial electric field able to remove the small ions in less than 0.2 s. When the two circular grids are not in use, they are grounded to avoid ground loops or parasitic electric fields.

The total in-flow to the chamber varies between 100 and  $150 L min^{-1}$ . Most of the flow is taken by the instruments connected to the chamber and a portion of it is purged through a valve that controls the pressure inside the chamber. The chamber is kept at 5 mbar above the atmospheric pressure to avoid contamination from the outside. The chamber is equipped with an ultraviolet fiber-optic system that is installed at the top plate of the chamber, allowing OH production (Kupc et al., 2011) without heating up the chamber. The temperature is controlled by air circulating between the chamber and the insulation surrounding it. The temperature ranges from  $-80$  to  $100 ^\circ C$  with a stability inside the chamber of  $\pm 0.1 ^\circ C$  for each experiment. The chamber is filled with ultra-pure synthetic air, consisting of  $N_2$  and  $O_2$ , obtained from the evaporation of liquid samples. The air is humidified with a Nafion system, using water purified by recirculation through Millipore Super-Q filters and irradiated with UV radiation. Ozone is produced by illuminating a portion of the incoming dry air with UV light. The other trace gases, such as  $SO_2$  and  $NH_3$ , are added from gas cylinder reservoirs. These measures aim for the cleanest possible laboratory conditions (Schnitzhofer et al., 2014).

## 2.2 The particle beam

The particle beam is produced at the CERN Proton Synchrotron. The CLOUD chamber can be exposed to a 3.5 GeV  $c^{-1}$  positively-charged pion ( $\pi^+$ ) beam produced by a secondary target of aluminium or copper (Duplissy et al., 2010). The intensity of the particle beam can be varied to cover the full range of ionization in the troposphere, from an ion production rate of about  $2 \text{ cm}^{-3} \text{ s}^{-1}$ , equivalent to boundary layer levels, to  $80 \text{ cm}^{-3} \text{ s}^{-1}$ , equivalent to the levels in the free troposphere (Kazil and Lovejoy, 2004; Zhang et al., 2011). The beam consists of pions and muons that have approximately the same energy of about 3.5 GeV and come in spills. Each spill lasts for a few microseconds and delivers  $10^8$  pions at a time, at intervals of 20 s.

We varied the beam intensity, and thus the ion production rate in the chamber, by varying the aperture of the beam collimator, which consists of two mechanical jaws made of concrete that can slide vertically and horizontally. At full aperture (60 mm  $\times$  60 mm) we have the maximum flux of pions through the chamber. When the collimator was partially closed we reduced the flux of pions and we illuminated a smaller part of the chamber. When the collimator was closed, almost no pion could reach the chamber. In this case, only GCRs, which are passing through the chamber vertically, and some residual pions that pass the blocker, hitting the chamber horizontally, were responsible for the ionization. For this reason, the ion production rate in the chamber at GCR conditions was about 30% higher than it would be if the accelerator had been shut down completely.

The pion beam is deliberately defocused to maximize the area where the ionization takes place (Figure A.2 .c, shaded area). About 70% of the volume of the chamber is directly ionized and the two mixing fans produce a uniform distribution of ions inside the chamber, the mixing time inside the chamber is estimated to be between 1.7 and 3.6 min (Voigtländer et al., 2012).

## 2.3 The hodoscope and the B1.2 counter

The hodoscope is an array of scintillation detectors. It measures the pion flux through the chamber and gives accurate positional information on the particle beam. It consists of 9 plastic scintillator slabs adjacent to one another in a vertical orientation and 9 plastic scintillator slabs oriented horizontally (Mizin et al., 2011). The vertical and horizontal slabs form a grid orthogonal to the particle beam, covering an area of about 2 m  $\times$  2 m. An ionizing particle, hitting a point on this grid, is located by measuring the coincidence of the signal of two detectors (one vertical and one horizontal). The B1.2 counter consists of two scintillation detectors placed right in front of the beam aperture. It also measures the coincidence of the signal of two detectors in order to count only the pions from the beam. From the number of particles measured per unit time and from the location of the scintillators

crossed, we can independently retrieve the beam intensity (Hz) and its horizontal and vertical profile. This allows us to discriminate between high energy particles coming from the beam, which travel almost horizontally, and the GCRs.

## 2.4 Neutral cluster and Air Ion Spectrometer (NAIS)

The Neutral cluster and Air Ion Spectrometer (NAIS, Gagné et al., 2011; Mirme and Mirme, 2013, SER NAIS12) is manufactured by Airel Ltd in Estonia and measures atmospheric ions in the range [0.8 - 42] nm and total aerosol particle population in the range [2.5 - 42] nm. It consists of two cylindrical Differential Mobility Analysers (DMAs) working in parallel, classifying negative and positive ions at the same time (Manninen et al., 2009). The ions are simultaneously classified according to their mobility and detected by a stack of 21 electrometer rings for each analyser. The device is equipped with a unipolar charging unit for each analyser that can be switched on and off.

During the CLOUD experiments, the instrument operated in three modes: particle, ion and offset. In particle mode the main charging unit is turned on. In ion mode the main charging unit is turned off. In offset mode the main charger unit stays off and a filter unit is turned on to make a zero measurement. This zero measurement is used to determine the net signal due to ions and particles. To maximize the flow in the tube and reduce the diffusion losses, the NAIS and a mass spectrometer shared part of the 2.54 cm outer diameter sampling line. Outside the chamber, the flow was split (20 L  $\text{min}^{-1}$  to the NAIS, 10 L  $\text{min}^{-1}$  to the APi-TOF) using a Y union (Figure A.2.b).

The NAIS operates at a 54 L  $\text{min}^{-1}$  total inlet flow. The high intake of the NAIS is a challenge in aerosol chamber experiments, where it is required to minimize the amount of air withdrawn. For this reason, the NAIS was operated with a recirculation system, which diluted the inlet flow with filtered air coming from the exhaust of the instrument. The filtered air formed an annulus around the sample flow (Figure A.2.a). The use of the dilution system allowed us to reduce the withdrawn flow from the chamber from 54 to 20-30 L  $\text{min}^{-1}$ . In this study, we only present the data recorded in ion mode, in particular the data from of the first 9 electrometers that collect ions with mobility diameter smaller than 1.9 nm (small ions). The detailed setup used for this work is presented in Figure A.2.

## 3 Theoretical methods

### 3.1 Calculation of the ion-ion recombination rate

Using the balance equation at steady state, the ion-ion recombination rate and a linear loss term were determined using the measured ion concentration and the ion production rate. We can describe the ionization processes inside the chamber us-

ing the following balance equation:

$$\frac{dn_{\pm}}{dt} = q - \alpha n_+ n_- - \beta_{\pm} n_{\pm} - k_{CS}(N, D_p) n_{\pm}, \quad (1)$$

where  $n_{\pm}$  is the absolute concentration of positive or negative small ions ( $\text{cm}^{-3}$ ),  $q$  is the ion production rate (IPR,  $\text{cm}^{-3} \text{s}^{-1}$ ),  $\alpha$  is the recombination coefficient ( $\text{cm}^3 \text{s}^{-1}$ ),  $\beta$  is a first order loss term ( $\text{s}^{-1}$ ) that describes the ion-wall interactions in the aerosol chamber, and other loss mechanisms of the first order,  $k_{CS}$  is the coagulation sink, i.e., the rate at which ions are lost by diffusion onto aerosol particles ( $\text{s}^{-1}$ ) (Kulmala et al., 2001; Leppä et al., 2011). At a constant  $q$ , the steady state that is eventually reached depends on the values of  $\alpha$ ,  $\beta$  and  $k_{CS}$ .

In this study  $\alpha$  does not include the effect of initial recombination (Gryzinski et al., 2007). Initial recombination would mainly affect primary ions, which are not considered in our study that focuses on cluster ion-ion recombination, as the NAIS measures cluster ions with electrical mobility smaller than  $3.2 \times 10^{-4} \text{ m}^2 \text{ V}^{-1} \text{ s}^{-1}$  (larger than 0.8 nm in mobility equivalent diameter).

We can make some assumptions to simplify Eq.1. By assuming that  $n_- \simeq n_+$ , and that the ion loss by coagulation is negligible compared with the other sink terms, we get:

$$\frac{dn}{dt} = q - \alpha n^2 - \beta n, \quad (2)$$

These assumptions were well posed in our case, since the average difference between the number concentration of positive and negative ions was 10 %. The 25<sup>th</sup> and 75<sup>th</sup> percentile are 5 % and 14 %, respectively (see Fig. A.3). This variation in the concentration of positive and negative polarities is compatible with the ratio of the square root of the mean ion electrical mobilities ( $Z$ ), which is proportional to the loss rate of ions to the walls of the chamber  $\sqrt{\frac{Z_{pos}}{Z_{neg}}} = 0.9$ . During our experiments  $k_{CS}$  was constantly below  $3 \times 10^{-7} \text{ s}^{-1}$ , due to negligible aerosol concentration in the chamber. The solution of Eq. 2 can be found analytically (Israël, 1970):

$$n(t) = \frac{n_1(n_0 - n_2) - n_2(n_0 - n_1) \exp(-t\sqrt{\beta^2 + 4\alpha q})}{(n_0 - n_2) - (n_0 - n_1) \exp(-t\sqrt{\beta^2 + 4\alpha q})}, \quad (3)$$

where  $n_0$  is the concentration of ions at  $t = 0$ ,  $n_1 = \frac{-\beta + \sqrt{\beta^2 + 4\alpha q}}{2\alpha}$  and  $n_2 = \frac{-\beta - \sqrt{\beta^2 + 4\alpha q}}{2\alpha}$ .

If we consider steady state conditions,  $\frac{dn}{dt} = 0$ , Eq.2. becomes a second order polynomial. The recombination coefficient  $\alpha$ , and the linear loss term  $\beta$ , can be retrieved by fitting a second order polynomial function to the data, treating them as free parameters. The ion concentration was set to zero at  $q$  equal to zero. The linear loss term  $\beta$ , equal to  $(8.3 \pm 1.6) \times$

$10^{-3} \text{ s}^{-1}$ , was calculated based on the dataset with the best statistics ( $T = 20 \text{ }^\circ\text{C}$ ,  $\text{RH} = 38 \%$ ). We assumed that the value of  $\beta$  did not change in other cases. To check our assumption, we performed a sensitivity analysis of the ion-ion recombination coefficient  $\alpha$ , shown in Fig. 5. The variation in  $\alpha$  for different assumed values of  $\beta$  is linear and small compared with the observed variation of  $\alpha$  due to changing conditions in the chamber.

Equation 3 was used to check the values of  $\alpha$  and  $\beta$  retrieved at steady state by comparing the resulting  $n(t)$  to the ion decay data (Figure A.4). The ion production rate from the beam ( $q_b$ ) was calculated using the following equation (Duplissy et al., 2010):

$$q_b = N_b I L / V, \quad (4)$$

where  $N_b$  is the number of pions per unit time that hit the chamber.  $N_b$  can vary between 0 and  $1.5 \times 10^6 \text{ s}^{-1}$ .  $I = 61 \text{ i.p. cm}^{-1}$  is the mean ionization per cm for a 3.5 GeV/c  $\pi^+$  in air at s.t.p. (Smirnov, 2005),  $L = 284 \text{ cm}$  is the path length of a beam particle in the chamber, and  $V = 26.1 \times 10^6 \text{ cm}^3$  is the chamber volume. The ion production rate was scaled for different air density at different temperatures.

We used a value of  $1.84 \text{ cm}^{-3} \text{ s}^{-1}$  for the intensity of the GCRs. The total ion production rate  $q$  is given by the sum of the GCR contribution  $q_0$  and the beam contribution  $q_b$ ,  $q = q_0 + q_b$ . The ion concentration was corrected for sample dilution, due to the dilution system described in section 2.4 and shown in Fig. A.2. The dilution system was tested in the laboratory and, for the flow range that was used in the experiments (20 - 25 LPM), it agreed within 28 % with the ideal system used for our calculations.

### 3.2 Modelled ion-ion recombination coefficient

The temperature dependency of the calculated ion-ion recombination coefficient was compared to the values computed using three different model approaches. The first model derived by J. J. Thomson (Gardner, 1938; Loeb, 1955; Thomson and Thomson, 2013) considers recombination governed mainly by Brownian motion of ions and molecules and computes the recombination coefficient as:

$$\alpha = \sqrt{2} \pi d^2 C \epsilon. \quad (5)$$

Where  $C = \sqrt{3k_b T / m}$  is the root mean square thermal velocity of the ions,  $d = e^2 / [(3/2)k_B T]$  is the distance where the Coulomb potential energy between the two ions of opposite polarity is equal to their thermal kinetic energy, and  $\epsilon$  is the probability of the two ions to recombine once they are at distance  $d$ . As derived by Thomson,  $\epsilon = 2w - w^2$ ,  $w = 1 - 2[1 - \exp(-x)(x + 1)]/x^2$ ,  $x = 2d/L$  and  $L$  is the mean free path of the ions. The equation for  $\alpha$  then becomes:

$$\alpha (\text{cm}^3 \text{ s}^{-1}) = 1.90 \times 10^{-5} (273/T)^{\frac{3}{2}} 3/2 \sqrt{1/M} \epsilon, \quad (6)$$

when  $T$  is the temperature in Kelvin and  $M$  is the mass of the ion in Da.

The second model taken into consideration and compared with our experimental data is described by López-Yglesias and Flagan (2013). Based on Hoppel and Frick (1986), this model was developed for ion-aerosol attachment, but it can be used to compute the recombination coefficient, if we use an aerosol that is as small as the ion and with opposite charge. The model accounts for Brownian motion, Coulombic interaction, image charging, polarization of the molecules and three-body trapping.

The third model, by Brasseur and Chatel (1983), is a parametrization used for describing the ions in the stratosphere based on earlier work by Bates (1982) and Smith and Adams (1982) that has the expression

$$\alpha(\text{cm}^3 \text{s}^{-1}) = 6 \times 10^{-8} \sqrt{300/T} + 6 \times 10^{-26} [M_{\text{air}}] (300/T)^4, \quad (7)$$

where  $[M_{\text{air}}]$  is the concentration of air molecules in  $\text{cm}^{-3}$ .

## 4 Results and discussion

The ion-ion recombination was measured in the CLOUD chamber at different temperatures, relative humidities and concentrations of ozone and sulfur dioxide. We obtained a value of  $2.3 \pm 0.7 \times 10^{-6} \text{cm}^3 \text{s}^{-1}$ , at 298 K and RH = 40%, higher than the constant value of  $1.6 \times 10^{-6} \text{cm}^3 \text{s}^{-1}$  usually reported in the literature (Laakso et al., 2004). Interestingly, we found that  $\alpha$  depends on both temperature and relative humidity (Table A2 and A1). Figure A.5 shows the results of four sets of experiments, where the ion concentration was measured as a function of  $q$ . For each set of experiments, we kept all the parameters constant except the one under investigation: the concentration of ozone (200 – 700 ppb), sulfur dioxide (0 – 50 ppb), temperature (218, 248, 278 and 293 K) and relative humidity (0, 7, 40 and 70%). The measured recombination rate showed a strong dependency (about a factor of 5 change) on temperature and relative humidity and, to some extent (a factor of 2), on the concentration of sulfur dioxide. The variability in the ozone concentration appeared to have little influence, if any, on the measured recombination rate.

The retrieved loss rate for ions ( $8.3 \times 10^{-3} \text{s}^{-1}$ , Fig. A.6), differs markedly from the linear loss rate retrieved for the sulfuric acid neutral monomer ( $1.7 \times 10^{-3} \text{s}^{-1}$ , Almeida et al., 2013). This difference between the loss rates might be partially explained by the image charge effect of the ions with the chamber walls, even though this remains a controversial topic (McMurry and Rader, 1985; Mayya and Sapra, 2002; Vauge, 2002; Chang et al., 2012). Losses might be due to some non-ideal behavior, e.g., retention of surface charge, of some insulator in the clearing field cage region or in proximity to the mixing fans. Yet another possibility is that the mix-

ing in the chamber influences the ion concentrations, creating a higher linear loss rate. Finally, it is possible that sulfuric acid is not lost to the walls with unit efficiency.

### 4.1 Temperature and relative humidity dependency of the recombination rate

The strongest effect we observed was an increase in the ion-ion recombination coefficient with decreasing temperature (Fig A.5). The ion-ion recombination coefficient increased by approximately a factor of 4 (from 2.3 to  $9.9 \times 10^{-6} \text{cm}^3 \text{s}^{-1}$ ) as temperature decreased from 293 to 218 K. This behavior seems not to be predicted by two of the three models and only partially by the third model, as shown in Figure A.7. Interestingly, all the models agree with each other and with the experimental results over a range of temperatures from 273 to 298 K, but there are big discrepancies in the absolute values as well as in the functional form of the temperature dependency at temperatures below 273 K. The model that seems to agree best with our data is the one by Brasseur and Chatel (1983), linked more directly to atmospheric data. The model by López-Yglesias and Flagan (2013) that accounts accurately for all the possible physical processes does not agree with our data. Another option could be related to the evaporation of ions in the line. If the number of ions reaching the instrument was lower because of the losses for evaporation, the resulting recombination would be overestimated. The effect of the mixing fan and the chemical composition of the ions could vary with temperature, which would affect the recombination coefficient.

We observed a strong dependency of the recombination coefficient on relative humidity (RH). The ion-ion recombination coefficient decreases with decreasing RH from  $9.4 \times 10^{-6}$  to  $2.0 \times 10^{-6} \text{cm}^3 \text{s}^{-1}$  as RH drops from 70% to 0%, at a constant temperature of 298 K (Figure A.8). The decrease of the ion-ion recombination coefficient at increasing RH values could be related to an increase in size of the small ions: higher values of RH would form larger hydrated ions that would be less mobile, thus decreasing the recombination rate. The change in ion mobility is plausible, as we know that aerosol particles with a diameter larger than 10 nm tend to be hygroscopic, changing their diameter according to the RH (Onasch et al., 1999; Keskinen et al., 2013). We also know, from quantum chemistry calculations, that ions form clusters with water and that the amount of water attached is dependent on RH (Kurtén et al., 2007; Husar et al., 2012; Henschel et al., 2014; Olenius et al., 2014). This explanation matches the observed data qualitatively. In fact, according to quantum calculations shown in Kurtén et al. (2007) for sulfuric acid ions at 20% RH there are mostly 1 or 2 molecules of water in the cluster, whereas at 80% RH there are 3 or 4. We get the equation below if we assume that the ions in the chamber: 1) are mainly sulfuric acid molecules, or behave the same way as sulfuric acid molecules, 2) are perfect spheres, and 3) that their mass is the sum of the masses of the

molecules that compose the ions.

$$D_p = \sqrt[3]{\frac{6 \cdot (M_{H_2SO_4} + M_{H_2O} \cdot n_w)}{\pi \rho}} + 0.3 \text{ nm} \quad (8)$$

with

$$\rho = \rho(n_w) = \frac{M_{H_2SO_4} \cdot \rho_{H_2SO_4} + M_{H_2O} \cdot \rho_{H_2O} \cdot n_w}{M_{H_2SO_4} + n_w \cdot M_{H_2SO_4}} \quad (9)$$

Where  $D_p$  is the mobility diameter of the ions,  $M_{H_2SO_4}$  and  $M_{H_2O}$  are the masses in Da of a sulfuric acid molecule and of a water molecule, respectively,  $n_w$  is the number of water molecules in a cluster and 0.3 nm is the difference between mobility diameter and mass diameter (Mäkelä et al., 1996; Ku and de la Mora, 2009). This calculation gives a  $D_p$  of 0.91 - 0.94 nm at 20% RH and of 0.94 - 1.01 nm at 80% RH. Unfortunately, the effect of RH on the ion diameter could not be observed directly, because this change in diameter is too small to be detected by the NAIS, given its low size resolution (Mirme and Mirme, 2013). However, the RH dependency of the recombination coefficient as a change in ion mobility is unclear, and cannot be predicted satisfactorily by any of the models found in the literature. Only the model described in Gardner (1938) predicts an increase of the recombination coefficient for smaller ions, although it underestimates the absolute values.

## 4.2 Atmospheric implications

It is important to account for the ion-ion recombination when modelling ion concentrations. As described by equation (1), the term representing the ion-ion recombination is always present and becomes dominant for large ion production rates and when coagulation sinks are small, e.g. in the free troposphere (Volland, 1995).

The possibility of a strong dependency of the ion-ion recombination rate on temperature must be taken into account when interpreting and modelling data of ion concentration or ion production in environments or model domains with a large temperature variability. In fact, a change in temperature from -54 °C to 20 °C, a typical temperature change from the free troposphere down to the boundary layer, can cause a change in the recombination rate as high as one order of magnitude.

The RH seems to influence the rate of recombination as well, by decreasing the mobility of ions. In this study, the measurements at different RH were taken at constant temperature and, therefore, changes in RH reflect those in the absolute water vapor concentration. The lower the RH, the faster the ion loss via recombination was. This might be analogous to what happens at high altitudes, where water molecules are scarce.

It should be noted that, in the present study, the pressure was kept constant at +5 mbar above the atmospheric level, during the experiments. In the atmosphere, however, a decrease in temperature with increasing altitude is associated

with a decrease in pressure as well. In the CLOUD chamber it is not possible to lower the pressure below typical sea level values, as the chamber is not designed to withstand under-pressure with respect to the outside pressure. So it was not possible to experimentally measure the variation of the ion-ion recombination rate at pressures lower than 900 hPa. It has been shown, however, that the ion-to-particle coefficients are considerably different in ambient conditions at 20 km altitude than at ground level, with the recombination coefficient being an order of magnitude lower at 20 km altitude than at ground level López-Yglesias and Flagan (2013). When moving from ground level to 20 km altitude, the decrease in recombination coefficient due to decrease in pressure is only partly counterbalanced by the increase in recombination coefficient due to decrease in temperature. Consequently, the pressure effect on recombination coefficient needs to be taken into account when interpreting data measured at substantially different pressure than in this work, but investigating the pressure effect is beyond the scope of this study.

## 5 Conclusions

We carried out an experimental determination of the ion-ion recombination coefficient in the CLOUD chamber at CERN. The retrieved recombination coefficient at 20 °C and 40% RH agrees with the literature values and with the models. We see no clear dependency of the ion-ion recombination rate coefficient for different ozone concentrations and only a weak variation at varying sulfur dioxide concentrations. Instead, we notice a strong dependency of the ion-ion recombination coefficient on temperature and relative humidity that has not been reported in previous studies. The ion-ion recombination coefficient varied between  $9.7 \times 10^{-6}$  and  $2.3 \times 10^{-6} \text{ cm}^3 \text{ s}^{-1}$  over the temperature range 220 to 293 K and between  $9.3 \times 10^{-6}$  and  $1.5 \times 10^{-6} \text{ cm}^3 \text{ s}^{-1}$  over the range of relative humidities from 0 % to 70 %. The temperature dependency is not well described by any of the models found in the literature, only the model by Broussier and Chatel (1983) seems to give results following the temperature dependence of our experimental data within 50% uncertainty. The RH dependency of the ion-ion recombination coefficient is not well understood or described theoretically, but can be interpreted as an effect of ion hydration: ions clustering with water molecules decreases their mobility, and therefore they recombine at a lower rate. Only the model by Gardner (1938) shows a functional dependency that supports this explanation, but it fails in reproducing the absolute values. It is also important to note that our results for the RH dependence of the ion-ion recombination coefficient were studied for only one temperature, 20 °C, and that the extent of the dependency might be different at different temperatures.

This was the first study to investigate experimentally the ion-ion recombination in a highly controlled environment at atmospheric relevant temperature, humidity and trace gas concentrations. Our main finding was that the recombination coefficient depends strongly on temperature and relative humidity, quantities that have a large variability in the troposphere. This work can be considered as a starting point for future studies in which pressure, chamber inhomogeneities and ion chemical composition could be taken into account.

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## Appendix A

### Solution of the balance equation

$$\frac{dn}{dt} = Q - \alpha n^2 - \beta n. \quad (1A)$$

The right hand term of Eq.(1A) is a second degree equation, therefore

$$\frac{dn}{dt} = -\alpha(n - n_1)(n - n_2), \quad (2A)$$

where:

$$n_1 = \frac{-\beta + \sqrt{\beta^2 + 4\alpha Q}}{2\alpha}$$

and

$$n_2 = \frac{-\beta - \sqrt{\beta^2 + 4\alpha Q}}{2\alpha}$$

are the solutions. Using the method of separation of variables (2A) becomes:

$$\frac{dn}{(n - n_1)(n - n_2)} = -\alpha dt \quad (3A)$$

Integrating, (3A) can be written as

$$\int_{n_0}^n \left( \frac{A}{n - n_1} + \frac{B}{n - n_2} \right) dn = \int_0^t -\alpha dt \quad (4A)$$

where  $n_0$  is the ion concentration at  $t = 0$  and  $A(n - n_2) + B(n - n_1) = 1$ .  $A(n - n_2) + B(n - n_1) = 1$  can be written as  $n(A + B) - (An_2 + Bn_1) = 1$  and its solution is:

$$A = -B = \frac{1}{(n_1 - n_2)} = \frac{\alpha}{\sqrt{\beta^2 + 4\alpha Q}}$$

As

By solving (4A) we obtain:

$$A \ln \left( \frac{|n - n_1|}{|n_0 - n_1|} \right) - A \ln \left( \frac{|n - n_2|}{|n_0 - n_2|} \right) = -\alpha t$$

We can drop the absolute value because these values are always greater than zero:

$$A \ln \left( \frac{n - n_1}{n_0 - n_1} \right) - A \ln \left( \frac{n - n_2}{n_0 - n_2} \right) = -\alpha t$$

$$A \ln \left\{ \frac{(n - n_1)(n_0 - n_2)}{(n - n_2)(n_0 - n_1)} \right\} = -\alpha t$$

$$\frac{(n - n_1)}{(n - n_2)} = \frac{(n_0 - n_1)}{(n_0 - n_2)} \exp \left( -\sqrt{\beta^2 + 4\alpha Q} t \right)$$

from this we can solve  $n(t)$ :

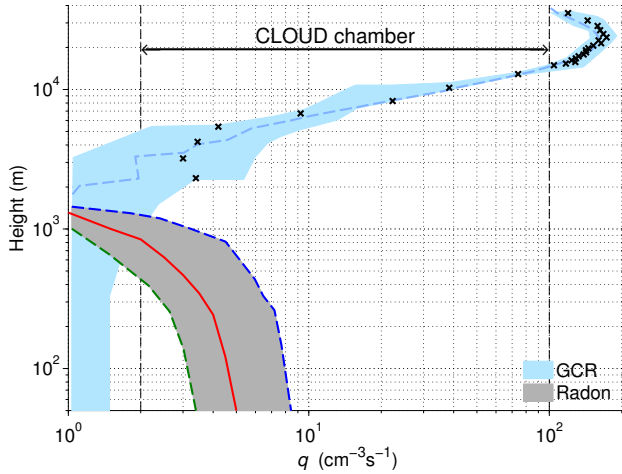
$$n(t) = \frac{(n_0 - n_2)n_1 - n_2(n_0 - n_1) \exp(-t\sqrt{\beta^2 + 4\alpha Q})}{(n_0 - n_2) - (n_0 - n_1) \exp(-t\sqrt{\beta^2 + 4\alpha Q})} \quad (5A)$$

T (°C)	$\alpha \times 10^{-6}$ ( $\text{cm}^3 \text{s}^{-1}$ )	$\sigma_\alpha \times 10^{-6}$ ( $\text{cm}^3 \text{s}^{-1}$ )
20	2.3	0.7
5	1.6	0.6
-25	7.6	1.0
-55	9.7	1.2

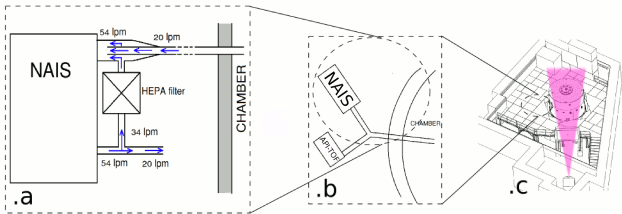
**Table A1.** Values of recombination coefficient and its uncertainty at different temperatures. The uncertainty reported is calculated with error propagation and includes the uncertainty in the fit.

RH (%)	$\alpha \times 10^{-6}$ ( $\text{cm}^3 \text{s}^{-1}$ )	$\sigma_\alpha \times 10^{-6}$ ( $\text{cm}^3 \text{s}^{-1}$ )
70	2.0	0.7
40	2.3	0.7
7	8.1	2.8
0	9.9	3.0

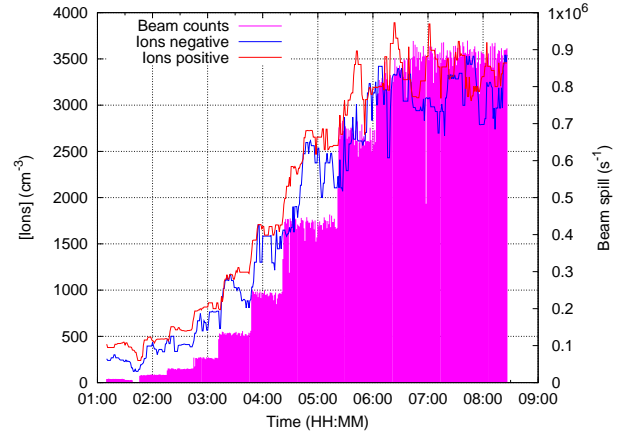
**Table A2.** Values of recombination coefficient and its uncertainty at different relative humidities. The uncertainty reported is calculated with error propagation and includes the uncertainty in the fit.



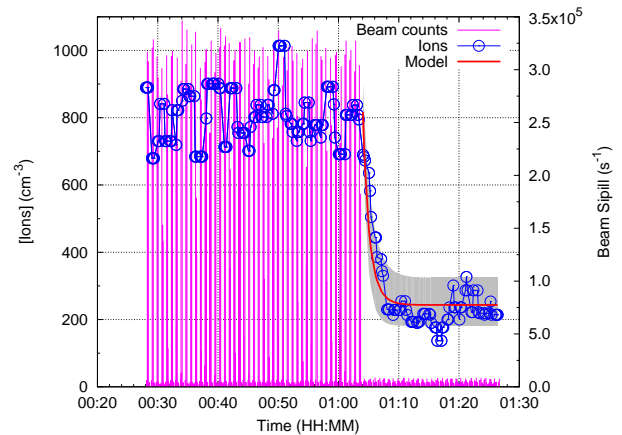
**Fig. 1.** A vertical profile of the ion production rate  $q$  based on literature data. The contribution of radon decay at mid latitudes is shown in gray, accounting for seasonal variation. The minimum is in summer and the maximum is in winter (Zhang et al., 2011). The contribution of galactic cosmic rays at mid latitudes, as presented by Usoskin et al. (2004), is shown in cyan. Black crosses indicate measurements by Harrison et al. (2014). The double arrow at the top shows the range of  $q$  explored in this study.



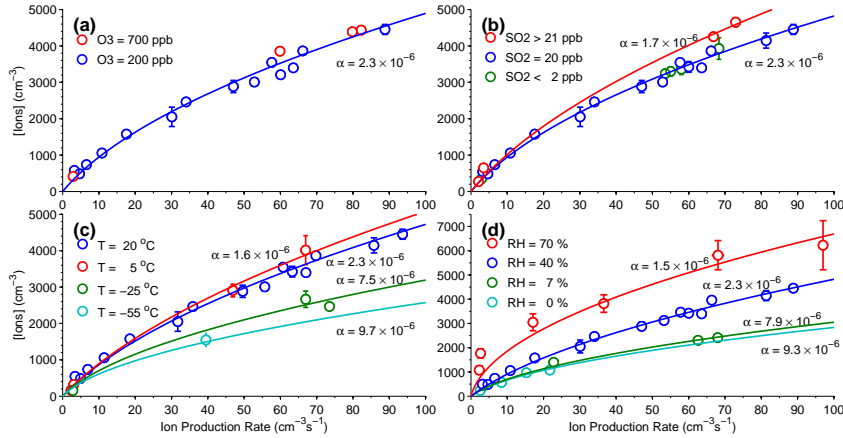
**Fig. 2.** a) Neutral cluster and Air Ion Spectrometer (NAIS) and its dilution system used during the CLOUD experiments. The sample air flow is withdrawn from the chamber at a flow rate that varies between 20 and 30 L min<sup>-1</sup> sample air from the chamber is diluted with a portion of the exhaust air of the instrument, which is filtered with a High Efficiency Particulate Air (HEPA) filter and mixed with the sample air. b) Configuration of the NAIS during the CLOUD experiments. c) Sketch of the chamber and the beam. The dashed circle represents the area where the NAIS was located, outside from the beam trajectory. The beam is deliberately defocused to maximize the volume in the chamber where ionization takes place.



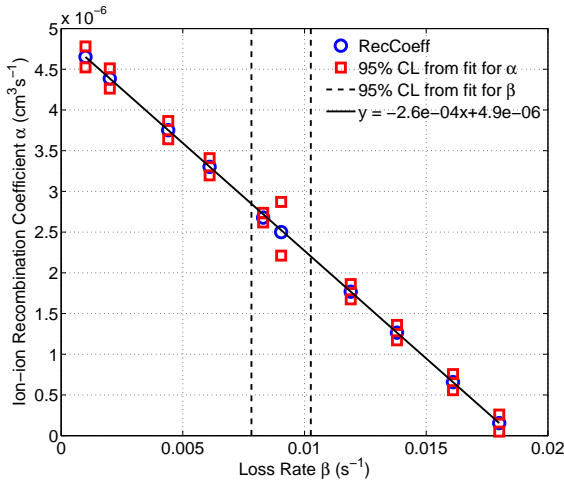
**Fig. 3.** In a typical experiment, the beam intensity is varied, changing the flux of pions crossing the chamber. The beam intensity is directly proportional to the Ion Production Rate (IPR) and is kept constant for a period of approximately 30 min at each step, to make sure that the ion concentration reaches the steady state. The mean ion concentration is related to the mean IPR at steady state via the balance equation (Eq.1). Blue and red colors correspond to negative and positive ions, respectively. Magenta is the beam intensity.



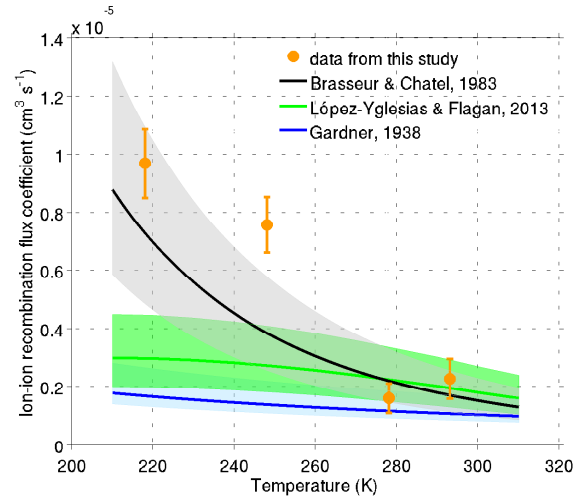
**Fig. 4.** Typical ion decay experiment. The ion concentration (blue line and circles), the solution to Eq. (1) (red line) and the beam counts (magenta line). The ion concentration is presented as the average between the concentration of negative and positive ions. When solving Eq. (1), the recombination coefficient and linear loss term, retrieved independently by fitting the steady state balance equation at the same conditions ( $T = 293$  K,  $RH = 0\%$ ), were  $9.3 \times 10^{-6} \text{ cm}^3 \text{ s}^{-1}$  and  $8.3 \times 10^{-3} \text{ s}^{-1}$ , respectively, and the ion production rate was  $8.3 \text{ cm}^{-3} \text{ s}^{-1}$ . The initial concentration of small ions,  $n_0$  ( $t=01:03$ ), was  $810 \text{ cm}^{-3}$  (average over time range from 00:24 to 01:03). The grey shaded area is the model uncertainty assuming uncertainty of  $\pm 30\%$  on  $n_0$ ,  $\alpha$  and  $\beta$ .



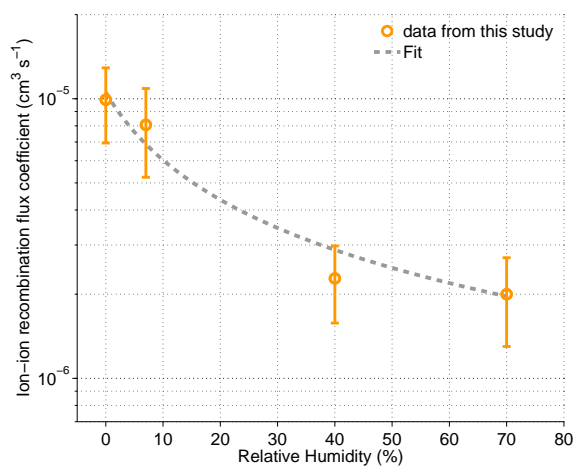
**Fig. 5.** Ion concentration as a function of ion production rate at (a) two different ozone concentrations (at 200 ppb and at 700 ppb); the other variables were kept constant ( $T = 20\text{ }^{\circ}\text{C}$ ,  $\text{RH} = 40\%$  and  $[\text{SO}_2] = 20\text{ ppb}$ ); (b) different  $\text{SO}_2$  concentrations between 2 and 30 ppb, temperature, RH and ozone concentration were kept constant ( $T = 20\text{ }^{\circ}\text{C}$ ,  $\text{RH} = 40\%$  and  $[\text{SO}_2] = 20\text{ ppb}$ ); (c) different temperatures (20, 5, -25 and -55  $^{\circ}\text{C}$ ); and (d) different relative humidities (0, 7, 40 and 70 %) at a constant temperature of 20  $^{\circ}\text{C}$ . The ion concentration is presented as the average between the concentration of negative and positive ions, the uncertainty is calculated as the standard deviation of the mean values at steady state.



**Fig. 6.** Variability of the retrieved recombination rate as a function of loss rate. The dataset used here includes experiments performed at  $T = 20\text{ }^{\circ}\text{C}$  and  $\text{RH} = 40\%$ . The point at the center presents the larger confidence level interval as the loss term was kept as free parameter. The other points were obtained forcing the loss term  $\beta$  to values varying from  $1 \times 10^{-3}$  to  $0.18\text{ s}^{-1}$ .



**Fig. 7.** A comparison between the ion-ion recombination coefficient, measured at different temperatures (points with error bars) and the recombination coefficient modelled using different models. The shaded area represents 50% uncertainty.



**Fig. 8.** The ion-ion recombination coefficient measured at different relative humidities, at a constant temperature of 20 °C. The dots are measured points, the dashed line is an exponential fit to guide the eye.