

# Influence of the ambient humidity on the concentration of natural deposition mode ice nucleating particles

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## Abstract.

This study reports measurements of deposition mode ice nucleating particle (INP) concentrations at ground level during the period July-December 2014 in Córdoba, Argentina. Ambient air was sampled into a cloud chamber where the INP concentration was measured at a temperature of  $-25^{\circ}\text{C}$  and a 15 % supersaturation over ice. They were performed on days with different thermodynamic conditions, including rainy days. The effect of the relative humidity at ground level ( $\text{RH}_{amb}$ ) on the INP concentration was analyzed. The number of INP activated varied from  $1\text{ L}^{-1}$  at  $\text{RH}_{amb}$  of 25% to  $30\text{ L}^{-1}$  at  $\text{RH}_{amb}$  of 90%. In general, a linear trend between the INP concentration and the  $\text{RH}_{amb}$  was found, suggesting that this variability must be related to the effectiveness of the aerosols acting as INP. From the backward trajectories analysis, it was found that the link between INP concentration and  $\text{RH}_{amb}$  is independent of the origin of the air masses. The role of biological-INP and nucleation occurring in pores and cavities were discussed as a possible mechanism to explain the increase of the INP concentration during high ambient relative humidity events. This work provides valuable measurements of deposition mode INP concentrations from the Southern Hemisphere where INP data is sparse so far.

## 1 Introduction

Natural ice nucleating particles (INP) are considered important in atmospheric processes since they induce freezing in clouds, thus initiating an efficient mechanism for cloud particles to reach a precipitating size. Consequently, many studies have been undertaken to determine their absolute concentrations, their origin and their chemical composition (DeMott et al. (2011); Hoose and Möhler (2012)).

Homogeneous freezing of supercooled droplets occurs at temperatures near  $-40^{\circ}\text{C}$ , but at temperatures warmer than approximately  $-36^{\circ}\text{C}$  heterogeneous nucleation occurs (e.g. Cantrell and Heymsfield (2005)), which involves four different modes: condensation freezing, immersion freezing, contact freezing and deposition nucleation (Vali, 1985). Cloud condensation nuclei (CCN) can serve as INP at temperatures below  $0^{\circ}\text{C}$  during the condensation freezing mode. During the immersion-freezing mode, a particle suspended in the droplet initiates freezing. In the contact-freezing mode, ice is formed by the collision of cloud droplets with interstitial aerosols within the cloud. When the environment is supersaturated with respect to ice and subsaturated with respect to liquid water, ice deposits on the INP directly from the vapor phase. This last mode is known as deposition nucleation. Measurements performed in the regime below water saturation are important in the understanding of the deposition nucleation (DeMott et al., 2011).

Temperature and supersaturation ratio are the two main parameters determining the deposition nucleation. Several reports have provided measurements of INP concentration for different temperature and supersaturation conditions. Studies linking INP concentration to temperature have shown an exponential trend in the dependence of INP on temperature (e.g. Fletcher (1962); Hussain and Saunders (1984); Meyers et al. (1992); López and Ávila (2013)). This same correlation was found when studying INP concentration regarding supersaturation (e.g. Rogers (1982); Al-Naimi and Saunders (1985); Hussain and Saunders (1984); Cooper (1980); López and Ávila (2013)).

Although temperature and supersaturation are recognized as the two main parameters in the activation of the INP, a significant divergence among the results of INP quantification is found in the literature when INP measurements are carried out at around the same temperature and supersaturation conditions (DeMott et al. (2011); Hoose and Möhler (2012)). In general, the results show that the ice-nucleating particles content of the atmosphere varies considerably from day to day and from place to place. For instance, important anomalies of atmospheric ice nucleating particles were observed under thunderclouds (Isono and Tanaka (1966); Isaac and Douglas (1973)). Increments at ground level ice nucleating particles concentrations by a factor of 10 or higher have been measured during precipitation from convective storm precipitation. These anomalies were associated with downdrafts in and under the thunderclouds. Prenni et al. (2013) also studied the impact of precipitation at a forested site on the concentration and composition of INP. They showed that at ground level INP-concentrations increase during rain events.

Thus, our knowledge of ice formation through the nucleation modes is still scarce and limited. As a consequence, the ice nucleation processes are currently very poorly represented in global climate models (e.g. Hoose et al. (2010)). Furthermore, the study of ice cloud formation over a wide range of atmospherically relevant temperatures and humidity values is challenging (e.g. DeMott et al. (2011); Murray et al. (2012)). Particularly, studies on anomalies in atmospheric ice nucleating particles concentration are important, since they will give us clues for solving problems on the origin and the nature of atmospheric ice nucleating particles. The major goal of this work is the study of the factors

that regulate the effectiveness of aerosols on deposition ice nucleation. Regarding the possible effect  
60 of precipitation on the INP-concentration, this study reports measurements of INP-concentration  
at ground level during the period July-December 2014 in Córdoba, Argentina. The experiments  
were performed on days with different thermodynamic conditions, including precipitation days, and  
the effect of the relative humidity at ground level on the INP concentration was analyzed. INP-  
65 concentrations in deposition nucleation mode were determined at a temperature of approximately  
–25°C and supersaturation of approximately 15 % over ice. The ambient relative humidity at ground  
level was measured during all the experiments, in order to link this variable to the INP concentration.

## 2 Experimental

Measurements were carried out in Córdoba, a mediterranean city of Argentina, located in the center  
of the country, in a semiarid region (latitude –31.4°C; longitude –64.18°C; 470 m.a.s.l.). Córdoba  
70 is the second largest city in the country, with approximately 1.3 million inhabitants. The climate  
is sub-humid and the prevailing wind direction is NE. The annual average precipitation is close to  
700 mm. However, the area is affected by severe and persistent dry periods that occur cyclically.  
Snowfalls are infrequent; thus, precipitations consist mostly of rain. Rainfall is highly seasonal,  
concentrated mainly in summer (December-February) having accumulated values higher than 400  
75 mm, while the dry winter season (June-August) has a cumulative precipitation depth close to 20 mm.

In a previous work we have quantified the INP-concentration (López and Ávila, 2013) in depo-  
sition nucleation mode at ground level in Córdoba, for temperatures between –15°C and –30°C,  
and ice supersaturation ranging between 2 and 20 %. In the present work the experiments were  
carried out following the same procedure as described by López and Ávila (2013) and a total of 63  
80 measurements of INP concentration were carried out during 45 days, since 28 July 2014 until 10  
December 2014 (winter and spring seasons). This procedure involves a cloud chamber of 46 L in  
volume placed inside a cold room. Controlled volumes of humid air were injected from the exterior  
(at ambient temperature) in order to increase the relative humidity in the cloud chamber. Since the  
same volume of air is removed while this air is injected, the number of aerosols and the pressure  
85 inside the chamber are kept around constant. The increasing of the relative humidity over ice inside  
the cloud chamber ( $RH_i$ ) is explained according the adiabatic isobaric mixing of two masses of air  
with different relative humidity and temperature (Curry and Webster, 1999). The supersaturation is  
achieved starting with a  $RH_i$  slightly lower than 100% in the cloud chamber and introducing con-  
trolled air injections from outside. Since the increase of  $RH_i$  for each air injection is known, it is  
90 possible to estimate the increase produced by a known number of injections, even once the vapor is  
supersaturated over ice. Once the supersaturation is reached, the INP are activated and then grown at  
the expense of vapor until ice crystals fall down onto a dish placed on the floor of the cold chamber.  
This dish contains a supercooled aqueous solution of cane sugar. Once in the sugar solution, the

ice crystals grow and reach a size large enough to be counted with the naked eye, as can be seen in  
95 Figure 1. For a more detailed description of the procedure, see López and Ávila (2013).

In all the measurements carried out in the present work the conditions of temperature (T) and  
supersaturation over ice ( $S_i$ ) were settled at around  $-25^\circ\text{C}$  and 15%, respectively. These values  
were chosen because, under these conditions, the number of activated INP are significant and can be  
easily determined (López and Ávila, 2013). Furthermore, these T and  $S_i$  values are experimentally  
100 quick to reach, which facilitate carrying out the experiments. In order to determine  $RH_i$  and T in  
the cloud chamber the Testo 435-4 multifunction measuring instrument was used. This instrument  
measures relative humidity in the temperature range  $[-50, 150]^\circ\text{C}$ . When  $RH_i$  is measured over ice  
at temperatures below  $0^\circ\text{C}$ , the resolution is 0.1%. The experimental uncertainty associated with the  
determination of T and  $S_i$  was  $\pm 2^\circ\text{C}$  and  $\pm 2\%$ , respectively. These errors arise from variations  
105 in the experimental conditions involved in each measurement, such as difference in the temperature  
between the beginning and the end of the air injections, variability in the  $RH_i$  increase between the  
injections, number of injections until reaching the particular  $S_i$ , and accuracy of the humidity sensor.

Thus, considering that T and  $S_i$  were the same for all the experiments (apart from the inherent  
errors), the INP concentration was studied as a function of the ambient relative humidity at ground  
110 level ( $RH_{amb}$ ); this is the relative humidity of the air mass injected from the exterior.  $RH_{amb}$  was  
measured during all the experiments by using the instrument EE31 Series Model D (E+E Elektronik),  
which had a remote sensing probe for relative humidity measurements in the temperature range  
 $[-40, 180]^\circ\text{C}$ . The uncertainty in the determination of  $RH_{amb}$  was  $\pm 1\%$ .

### 3 Results and discussions

115 The experiments were all performed for a supersaturated atmosphere with respect to ice and sub-  
saturated with respect to liquid water; thus, only the deposition mode INP concentrations were  
quantified. The INP concentration was determined at a temperature of approximately  $-25^\circ\text{C}$  and  
supersaturation of approximately 15% over ice. Figure 2 shows the temporal variations of the INP  
concentration values during the period under study. The grey points in the graph correspond to mea-  
120 surements performed during a rain event or immediately after rain had stopped. The error associated  
in the estimation of INP concentration did not exceed 10%. There are some time periods without  
experimental data mainly because the cold chamber was unable to achieve the expected temperature.  
The period of study is too short to perform a reliable statistical test showing a seasonal behavior of  
INP concentration; nevertheless, the results do not seem to indicate that there is any particular trend  
125 during this period.

In order to study the effect of the  $RH_{amb}$  in the INP concentration, Figure 3 displays the INP  
concentration as a function of the  $RH_{amb}$ . Again, the grey points indicate measurements performed  
on rainy days. INP concentration values can be seen to increase with  $RH_{amb}$ . The best linear fitting

of the data yield values of 0.39 and  $-9.2$  for the slope and ordinate, respectively, being the square  
130 of the correlation coefficient ( $R^2$ ) equal to 0.71. The linear trend between the INP concentration and  
the  $RH_{amb}$  is remarkable and the dispersion of the experimental points is likely related to the natural  
variability in the INP concentration and to the inherent errors associated with the determination of  
T,  $S_i$ ,  $RH_{amb}$ , and INP-number, which are present in each experiment.

The number of INP activated varies from  $\sim 1 \text{ L}^{-1}$  at  $RH_{amb} \sim 25\%$  to  $\sim 30 \text{ L}^{-1}$  at  $RH_{amb} \sim$   
135  $90\%$ . This variation is consistent with the results reported in López and Ávila (2013) on the same  
site, who found that INP concentration values varied between 2 and  $35 \text{ L}^{-1}$  at temperature of  $\sim$   
 $-25^\circ\text{C}$  and  $\sim 15\%$  of supersaturation over ice. Thus, the variability in the INP concentration in the  
previous work could be related to the variability in  $RH_{amb}$  at the time the measurements were carried  
out (this variable was not recorded in such experiments). Current results are also in agreement with  
140 those reported by Prenni et al. (2013), who showed that, at ground level, the INP concentration rises  
by a factor of up to 40 during rain. Actually, here we observe that not only rainy days are required  
for increasing INP concentrations but also high relative humidity is needed, as shown in Figure 3  
during rainfall events with low humidity.

A variety of insoluble particles such as volcanic ash, mineral dust, soot, metallic particles, or  
145 primary biological particles have been suggested as INP. However, the requirements needed for a  
surface to be an efficient INP are not completely understood (Pruppacher and Klett (1997); DeMott  
(2010); Hoose et al. (2010)). Prenni et al. (2013), Hader et al. (2014) and Bigg et al. (2015) presented  
evidence that the rising of INP concentration during rain events can be related to the increase of  
biological particle concentration. It has been shown that biological particles can be released during  
150 precipitation through mechanical mechanisms. For instance, winds produced by storms can release  
biological particles from leaf surfaces and soil. The release of these biological particles include  
biological INP (bacteria, fungal spores, and pollen) which can impact the INP population. This  
assumption was supported by different studies reporting both: the release of biological particles  
during rain events and the role of these particles as sources of INP (Huffman et al. (2012); Huffman  
155 et al. (2013)). Furthermore, Wright et al. (2014) provided observational evidence that the high  
relative humidity, coordinated with the arrival of a cold-frontal boundary on an observation site,  
triggered the active release of biological particles. Since mixing along the frontal boundary allows  
for the biological material to be entrained into the warm air that is being lifted over the cold front,  
they suggest that these biological particles can serve as immersion mode INP while droplets cool  
160 during ascent.

It is important to remark that unlike the present study, where the deposition nucleation mode  
was examined, the previous studies on a relation between INP concentrations with relative humidity  
or rain events mainly focused on condensation and immersion freezing modes. We have no direct  
evidence, so far, that the increase of the INP concentration with increasing humidity observed in the  
165 present work can be explained by biological particles.

Mineral dust aerosols seem to play a relevant role in deposition nucleation. They have active sites which are able to start ice nucleation. These active sites are linked to surface defects, like cracks, capillaries, pores or cavities which allow water vapor to condense in them under vapor pressures below saturation (Fukuta (1966); Fletcher (1969)). Water vapor condenses in a capillary of a wettable particle under a pressure below water saturation ( $p$ ), according to Kelvin equation

$$RT r \ln \left( \frac{p}{p_b} \right) = -2\gamma \nu_l \quad (1)$$

$R$  is the gas constant,  $T$  the absolute temperature,  $r$  the radius of the curved water surface,  $p_b$  the water vapor pressure over a flat water surface,  $\gamma$  the surface tension of water and  $\nu_l$  the molar volume of liquid water. Recently, Marcolli (2014) hypothesized that ice nucleation below water saturation occurs predominantly in water confined between closely spaced surfaces, such as cavities and pores that may form in aerosol particles. In this sense, it is plausible to expect that as the  $RH_{amb}$  increases, more water molecules from the surrounding environment can be adsorbed in cavities and pores. As a result, the condensation and the subsequent freezing can occur in the cavity when the relative humidity is lower than 100% outside. Considering this fact, a significant enhancement in ice nucleation efficiency should be expected at higher ambient relative humidity, which is consistent with the results reported here. Even though further work is needed to test and corroborate this hypothesis, the results presented in this work may support the idea that the deposition nucleation can be a pore condensation and freezing which occur in the defects on the surface of some aerosol particles as proposed by Marcolli (2014).

In order to relate the origin of the air masses to the behavior of INP measurements regarding RH, backward trajectories were calculated for all days when INP concentration measurements were carried out. Back trajectories were calculated by the NOAA HYSPLIT model (Hybrid Single-Particle Lagrangian Integrated Trajectory model) (Draxler and Rolph, 2003). Air mass back trajectories were calculated at 1000 m above mean sea level (AMSL), 48 h before the arrival time to the sampling site. NOAA trajectories were calculated every day of INP concentration measurements. The air mass back trajectories were subdivided into four groups, regarding the quadrants from the Compass Rose. As an example, Figure 4 shows of air mass back trajectories representative of the four studied sectors, which will be referred as North-West quadrant (example: Figure 4a), North-East quadrant (example: Figure 4b), South-West quadrant (example: Figure 4c) and South-East quadrant (example: Figure 4d). The lower sample number was obtained coming from NW quadrant (7.5%), while the highest numbers of INP measurements were obtained coming from NE quadrant (42.5%). Trajectories coming from SE and SW quadrants were 22.5% and 27.5%, respectively. Figure 5 shows the INP concentration measurements grouped following the previous classification of the origin of air masses. Observe that the number of INP measurements from NW quadrant is too scarce to make an analysis. However, the analysis from the three remaining quadrants show the increase of INP concentration with the increase in  $RH_{amb}$ , in the same way that it was observed in Figure

3. Results show that, from the studied period, variations in  $RH_{amb}$  are not explained because of the origin of the air masses. This is to say that, the relation between INP concentration and the ambient relative humidity is independent from the origin of the air masses. This fact discards the possibility  
205 that the increase in INP concentration is due to changes in the sources of the aerosols acting as ice nucleating particles. Thus, HYSPLIT results seem to be consistent with the previous assumption regarding to the importance of the ambient relative humidity on the ability of aerosols to act as INP.

#### 4 Summary and concluding remarks

Studies of heterogeneous ice nucleation with natural aerosols under a broad range of thermodynamic  
210 conditions are important, because they can provide clues for solving problems regarding the origin and the nature of atmospheric INP. This study reports measurements of deposition INP concentrations at ground level during the period July-December 2014 in Córdoba, Argentina. The experiments were carried out at temperature  $(-25 \pm 2)^\circ\text{C}$  and supersaturation  $(15 \pm 2)\%$  over ice. The effect of the relative humidity at ground level on the INP concentration was analyzed and a linear trend  
215 between the INP concentration and the  $RH_{amb}$  was found, suggesting that this variable must be related to the ability of the aerosols acting as INP. This assumption seems to be corroborated by the analysis of the backward trajectories of the air masses, which shows that the relation between INP concentration and the ambient relative humidity is independent from the origin of the air masses. Therefore, the variability observed on INP concentration cannot be explained regarding the changes  
220 in the sources of the aerosols acting as ice nucleating particles.

The role of biological-INP and nucleation occurring in pores and cavities (as suggested by Marcolli (2014)) was discussed as a possible mechanism to explain the increase on the INP concentration during high ambient relative humidity events. More ice nucleation studies should be carried out to address the differentiation between these mechanisms.

225 This work contributes to the study of the factors affecting the IN-concentration. Laboratory studies of heterogeneous ice nucleation with natural aerosols are important for the possibility to transfer the results into parameterizations for numerical models of clouds and climate. Furthermore, this work provides valuable measurements of deposition mode INP concentrations from the Southern Hemisphere where INP data is sparse so far.

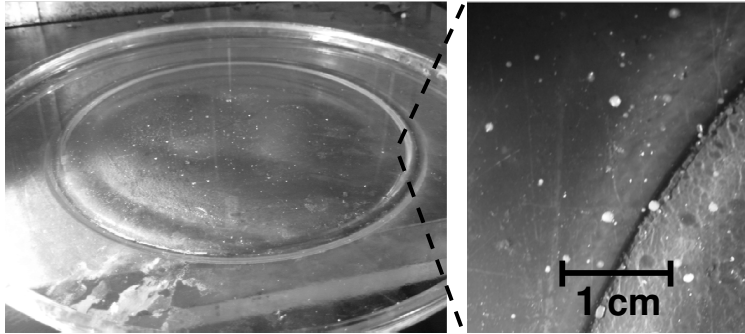
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**Fig. 1.** Images of crystals once they grown on the sugar solution.

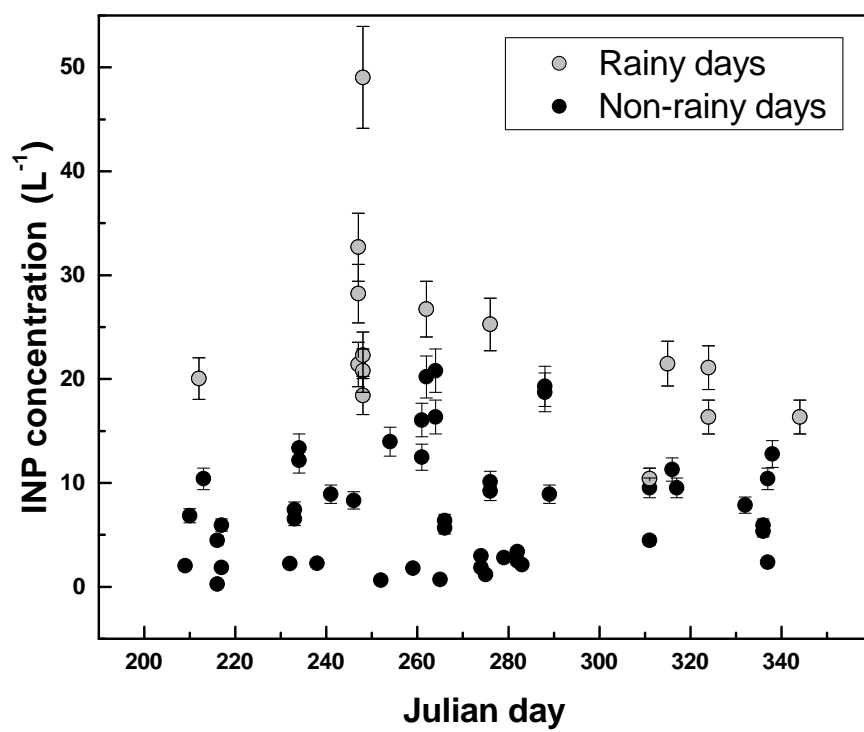


Fig. 2. Data of INP concentrations measured throughout the period studied.

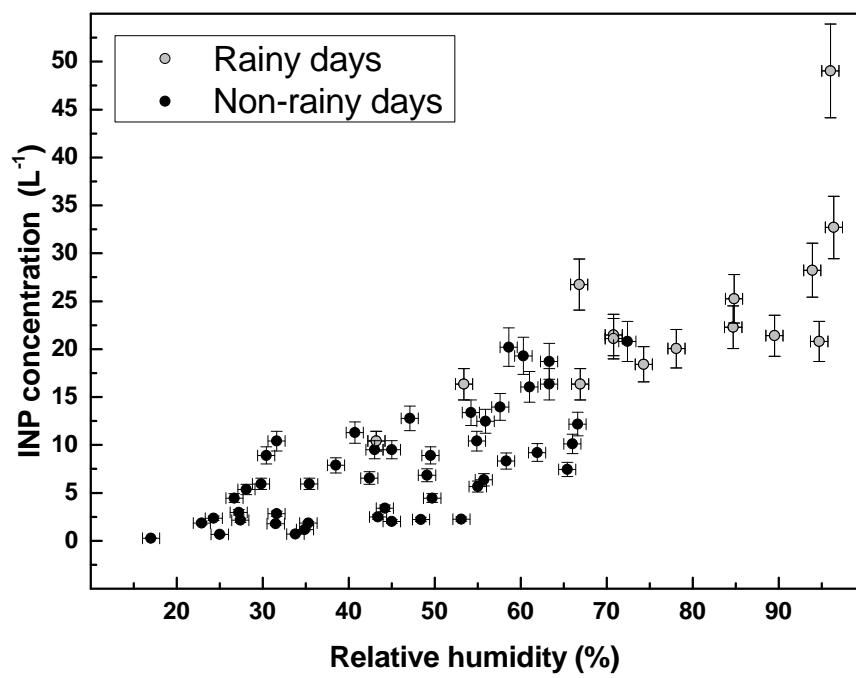
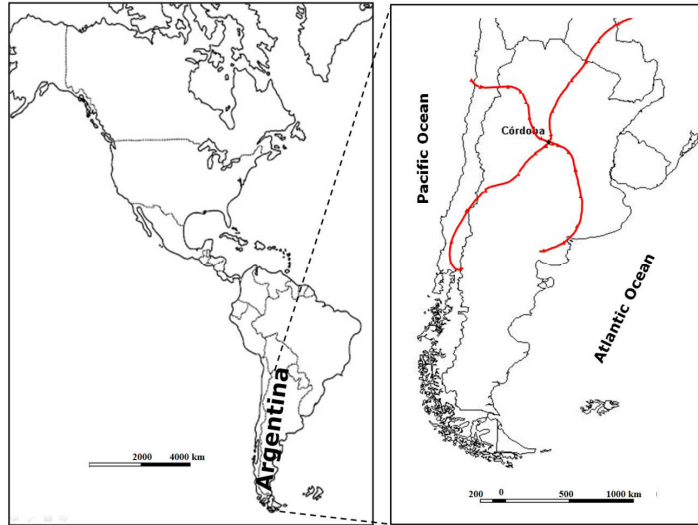
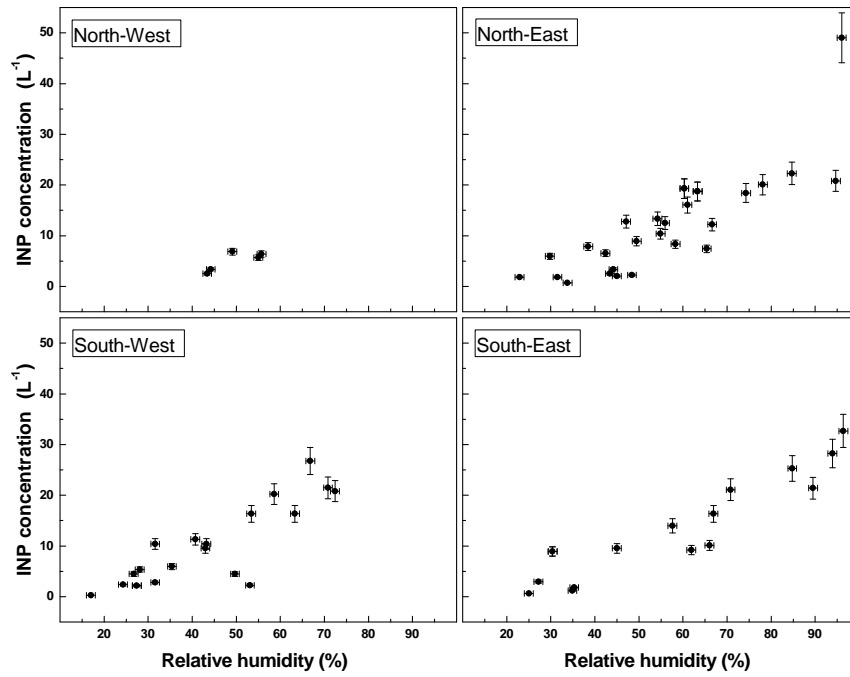


Fig. 3. Data of INP concentrations at different ambient relative humidity.



**Fig. 4.** Location of Argentina and air mass pathways of different quadrants: North-West, North-East, South-West and South-East.



**Fig. 5.** INP concentrations at different ambient relative humidity for the different origins of the air masses.