



Ion Concentrations And Sources Of Sequential Sampling Of Wet Deposition In Istanbul, Turkey

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Abstract

This study presents the relationships between ion concentrations and sources of ions in the precipitation events which happened on a) September the 08th, 2007 b) November the 10th, 2007 and c) December the 05th, 2007. Fortyfour sequential rain samples were collected in three rain events that were sampled in Istanbul-Turkey making use of a sequential rain sampler. Concentration of the main cations (K⁺, Ca²⁺, Na⁺, and Mg²⁺) and main anions (Cl⁻, SO₄²⁻, and NO₃⁻) were analyzed. The ratio of the mean Ca²⁺ (0.85) non-sea salt concentration to the mean total concentration led to the conclusion that most of the Ca²⁺ came from soil. This ratio for SO₄²⁻ ranged between 0.49 and 0.98, indicates a possible conclusion that the domestic heating and industrial activities were the main source of this ion.

Key Words

“Aerosol sources, sequential sampling, wet deposition.”

1. INTRODUCTION

Atmospheric particles have many sources which are natural and anthropogenic. Natural sources include fine soil blown into the air, coarse particles from soil by wind-blown, sea salts from breaking waves, reactions between natural gaseous emissions, biological material release, and volcanic eruptions. Dust from roads, wind erosion of tilled land and cropland, biomass burning, fossil-fuel combustion, construction, transportation sources and industrial emissions are main anthropogenic sources. Wide processes, which are used to find the sources ions in the precipitation, are non-sea salt (NSS) and enrichment factor (EF) estimations. There are many studies on investigation for researching the sources of the atmospheric constituents. Xu and Gao (2015) proved that sea salt and nss-SO₄²⁻ were the main components of marine aerosols over sampling region, of which high sea salt concentrations were associated with high wind speeds, while high nss-SO₄²⁻ could mainly come from anthropogenic emissions, especially over Asian marginal seas. Huang et al. (2009) claimed that in their study, the influence on precipitation chemistry of sea salts is strong since Guangzhou is located adjacent to the South China Sea. The NSS calculations of ions in the precipitations, in their study, are summarized as follows: The monthly VWM amount of the SSF of Cl⁻ is presented to be 37.8-100%; equivalent values are 26.6-68.9% for Mg²⁺, < 10% for K⁺, and just < 2% for F⁻, SO₄²⁻, and Ca²⁺. According to these data, Cl⁻ in the rain water primarily comes from the sea. Marine source can also be accepted to be the source of the half of the Mg²⁺ which demonstrates that for the explanation of Mg²⁺ in the precipitation, an additional source is needed. The SSFs of Ca²⁺, K⁺, F⁻, and SO₄²⁻, seems to be very low (<10%) which means that there exists another origin besides a marine source. Previous monitoring studies carried out to find the sources of constituents of atmospheric pollutants have profited by NSS estimations (Okay et al., 2002; Sakihama et al., 2008; Budhavant et al., 2010).

The present study is focused on the investigation of sources of the chemical components in the sequential rain events sampled in Istanbul, Turkey using the method of NSS and EF estimations. 44 sequential precipitation samples were collected in the aggregate for the three rain events. It was found Mg²⁺ came partly from NSS sources (37%) and SO₄²⁻ almost completely from industrial activities (87%). Ca²⁺ and K⁺ are found to have a high mean EF value. This finding leads to the result that the main source of these ions to be the soil.

2. MATERIAL AND METHODS

Goztepe that is the sampling area is on the Asian side of Istanbul (Figure 1). Istanbul that is populated by nearly 15 million people stands as the largest urban settlement in Turkey.

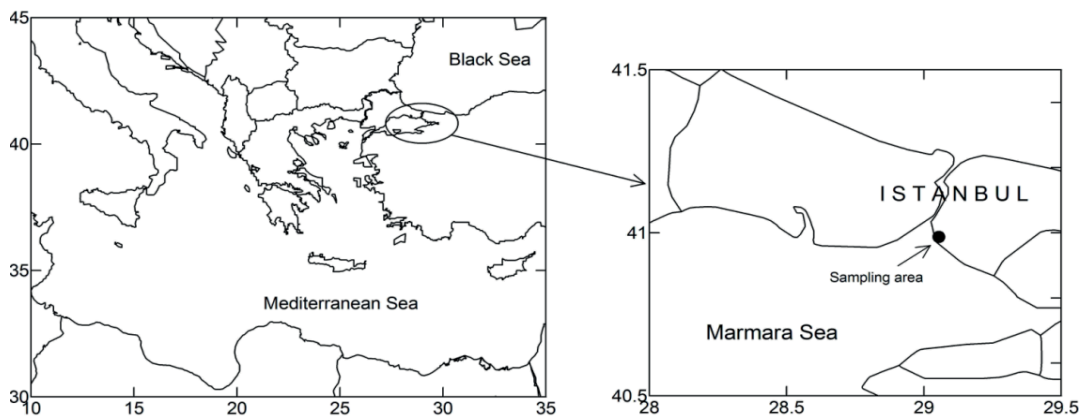


Figure 1. Sampling Site: Goztepe

The EF is a ratio that is usually used to distinguish aerosol sources (Vermette et al., 1988; Singer et al., 1993). This calculation relies on the elemental ratio which is found between ions that are collected in the precipitation or in the atmosphere. When the elemental ratio is compared with a ratio, which is similar, for a reference material, it gives significant information about the elements's source. In this study, the following equation is used in order to calculated EFs for sea water:

$$EF = \frac{(x/c)_{precipitation}}{(x/c)_{reference\ material}} \tag{1}$$

NSS ion concentration formula given by Vermette et al. (1988) was used in order to found the contribution of non natural sources of ions by scavenging processes.

$$NSSx_p = Totalx_p - c_p(x_r / c_r) \tag{2}$$

In this equation, x is the concentration of the ion of interest while c is the concentration of the reference ion. The subscript p and r stand for the ions in the precipitation and reference material, respectively.

To calculate the sea salt ratio, Brewer’s (1975) sea surface water concentrations were used as references. We chose Na⁺ as the reference element for sea to calculate the EFs of ions in the rain water.

An automatic sequential rain sampler that was designed and produced in the project carried out with the support of The Scientific and Technological Research Council of Turkey (TUBITAK) was used to collect samples (Akkoyunlu et al., 2008; Akkoyunlu et al., 2011; Akkoyunlu et al., 2013). Sequential sampler device has a polyethylene funnel of 0.28 m² with circular area and 12 polyethylene cups of 200 cm³ volume mounted on a sampler body. Precipitation sampling caps was taken from the sequential rain sampler body that they were attached to and brought to the laboratory prior to filtering of the samples and storing in pre-cleaned polyethylene bottles in the refrigerator at 4 °C before the process of chemical analysis.

K⁺, Ca²⁺, Na⁺, and Mg²⁺ concentrations were measured using an Hitachi-180-80 atomic absorption spectrophotometer. Cl⁻, SO₄²⁻, and NO₃⁻ concentrations were analyzed by a high performance Dionex model 4001 ion chromatography.

3. SOURCES OF THE IONS

Equation 1 has been used to calculate the mean EFs of the ions and equation 2 was for obtaining the mean NSS concentrations for all rain events, and these results were depicted in Figure 2. It’s been predicted that the rain event with the largest mean EF values in the rain water was on the December the 05th, 2007 due to K⁺, Ca²⁺, and Mg²⁺ were more enriched by soil compared to the other days of the rain event. Higher anthropogenic contribution for SO₄²⁻ during this day of the rain event which leads to higher mean EF values could be probably mentioned to detect the reason. The ratio for SO₄²⁻ ranged between 0.49 and 0.98. The result that the main source of SO₄²⁻ was the domestic heating and industrial activities might be concluded from these values.

The precipitation events occurred in Istanbul generally have high SO₄²⁻ concentrations mainly originated from the industrial processes and domestic heating. The authors found that SO₄²⁻ was enriched between 1.1 and 150 with respect to the sea (Akkoyunlu and Tayanç, 2003).

Panyakapo and Onchang (2008) indicated that NH₄⁺ and Ca²⁺ were the significant alkaline species for neutralization process. The EFs analysis indicated that Na⁺ and greater part of Cl⁻ were from the sea in this study, while Ca²⁺, Mg²⁺, K⁺, and SO₄²⁻ originate from local sources. It was revealed that the crusting as well as anthropogenic activities play an important role in contributing ionic species in wet precipitation by the application of factor analysis. In our study, mean EF of Mg²⁺ was calculated as 2.63, which corresponds to 37% NSS fraction. It seems that Mg²⁺ was diluted by sea and concentrated by soil.

The values of mean Cl⁻ and NSS concentration with large difference indicate that NSS has very small amount of partition in the ion. Zhang et al. (2007) found similar values to those of this study. The authors calculated that anthropogenic source fraction of SO₄²⁻ was 98.8% and crust fraction of Ca²⁺, K⁺, and Mg²⁺ were 99.4%, 97.1%, 58.7%, respectively in the 82 precipitation samples collected in southeastern China.

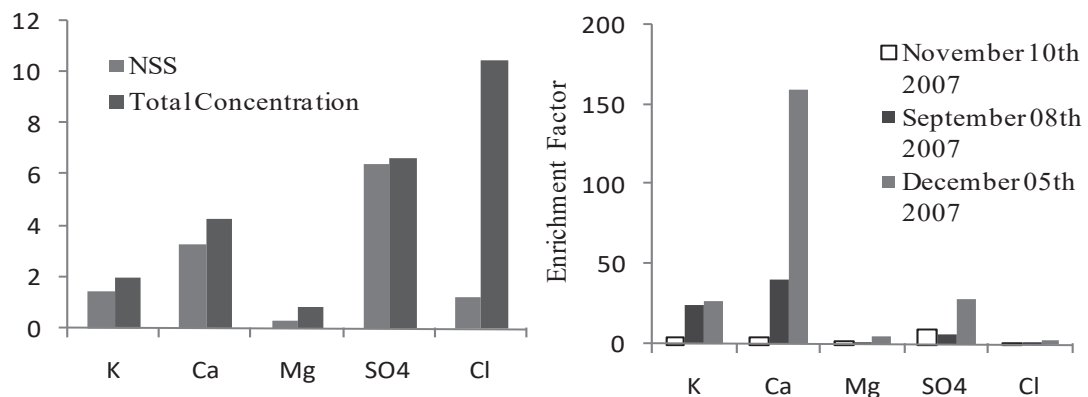


Figure 2. Mean NSS concentrations, total concentrations of ions (mg L⁻¹) and EF values.

Pearson correlation coefficients of the concentrations of ions for all three cases of rain days were given in Table 1. Maximum correlation coefficient was found 0.74 for the duration of sampling periods between SO_4^{2-} and NO_3^- that have industrial and domestic sources and in terms of concentration SO_4^{2-} and NO_3^- generally show similar variability. Na^+ and Cl^- is thought to be originated from sea as the same source. Correlation between them was calculated to be a second pick value as 0.71. SO_4^{2-} has high or moderate correlation with other ions such as K^+ , Ca^{2+} , Mg^{2+} which exist in the main soil wind-blown dust particles.

Nevertheless, no correlation between those of other ions and NSS concentration of SO_4^{2-} was found. It refers to the conclusion that SO_4^{2-} which was also thought as anthropogenic sources has different variability than the variability of other ions for duration of rain events.

Topçu et al. (2002) found similar results to the findings of this study. As an example, the authors calculated the correlation coefficient between SO_4^{2-} and NO_3^- as 0.92 and the correlation coefficients of Na^+ with Cl^- and Mg^{2+} as 0.77 and 0.54, respectively in the summer season, in Ankara, Turkey. Cheng et al. (2011) found that SO_4^{2-} and NO_3^- anions were highly correlated. It was found out that between these ions, the correlation coefficient was 0.7151.

Table 1. Pearson correlation coefficients between the overall concentrations of ions.

	Na^+	K^+	Ca^{2+}	Mg^{2+}	NO_3^-	SO_4^{2-}	Cl^-
Na^+	1	0.70	0.37	0.72	0.43	0.56	0.71
K^+		1	0.24	0.24	0.16	0.41	0.66
Ca^{2+}			1	0.36	- 0.14	0.04	0.72
Mg^{2+}				1	0.38	0.50	0.60
NO_3^-					1	0.74	0.23
SO_4^{2-}						1	0.23
Cl^-							1

The total concentrations which were described previously in the formula 2 and the reduction as a mean percentage of the NSS concentrations were indicated in Table 2. The decrease of concentrations during the initial stages as a percentage can be ordered in a descending order as $\text{K}^+ > \text{Cl}^- > \text{Na}^+ > \text{Mg}^{2+} > \text{Ca}^{2+} > \text{NO}_3^- > \text{SO}_4^{2-}$ and descending order of NSS concentrations as $\text{K}^+ > \text{Mg}^{2+} > \text{Ca}^{2+} > \text{SO}_4^{2-} > \text{Cl}^-$ in Table 2.

Fine particles of anthropogenic origin which contains SO_4^{2-} and NO_3^- are formed by gas-to-particle processes and their scavenging efficiency is much smaller than those of coarse particles. Due to the fact that rain drops are more efficient at scavenging the large particles and coarse mode of aerosols than fine particles. In comparison with the NO_3^- , SO_4^{2-} ions which exist as fine particles, more significant reduction of K^+ , Cl^- , Na^+ , Mg^{2+} , Ca^{2+} ions was found. It was found Mg^{2+} came partly from NSS sources (37%) and SO_4^{2-} almost completely from industrial activities (87%). There was no reduction of NSS concentrations of Mg^{2+} and SO_4^{2-} though other ions has different behaviors. This could be attributed to the possibility that the Mg^{2+} and SO_4^{2-} originated from NSS have undergone to similar scavenging effects during these precipitation events. Mean reduction of total and NSS concentration of all ions were calculated as 54.7% and 43.6%, respectively.

Table 2. Reduction as a percentage of the NSS and total concentrations.

	Na ⁺	K ⁺	Ca ²⁺	Mg ²⁺	NO ₃ ⁻	SO ₄ ²⁻	Cl ⁻
Total concentration	60.7	82.1	46.4	53.5	34.8	30.5	74.6
NSS concentration	-	63.6	41.0	55.2	-	35.5	22.8

4. SUMMARY AND CONCLUSIONS

Samples of sequential rains were collected in three rain events for the purpose of investigation of the ion concentrations and the sources of ions. During the beginning stages of each rain event, concentrations of nearly all of the ions decreased. The ratio of mean concentration for all ions sampled on the initial stages to that of the final stages of the rain events were calculated. The results of the rain events were found on September the 08th, 2007 as 1.30, on November the 10th, 2007 as 2.29, and on December the 05th, 2007 as 2. The decrease of concentrations during the initial stages was sorted in a descending order as K⁺> Cl⁻>Na⁺> Mg²⁺> Ca²⁺> NO₃⁻> SO₄²⁻. This indicated that K⁺, Ca²⁺, Cl⁻, Na⁺, Mg²⁺ generally in the coarse particle fraction were more efficiently scavenged than fine particles of anthropogenic origin which formed by gas-to-particle processes and contains SO₄²⁻ and NO₃⁻. Mean EFs and mean NSS concentrations of the ions were calculated for all rain events for the purpose of finding the sources of ions. It was found that the case of high EF for SO₄²⁻ and high correlation between NO₃⁻ and SO₄²⁻ indicates that large amounts of these ions originated in the combustion processes of industrial activities and domestic heating. Ca²⁺ and K⁺ are found to have a high mean EF value. This finding leads to the result that the main source of these ions to be the soil. EF of Mg²⁺ was calculated as 2.63. 63% of total amount of Mg²⁺ originated from sea. Sea might be the mutual source for the Na⁺, Cl⁻, and Mg²⁺ due to the high correlation coefficients among them.

5. ACKNOWLEDGEMENT

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