The creation of pollution mapping and measurement of ambient concentration of sulfur dioxide and nitrogen dioxide with passive sampler

Andaç Akdemir

Abstract

Background: Measurements of nitrogen and sulfur dioxide using passive sampler over 12 months in Samsun, Turkey, are compared with SO2 and NO2 concentrations obtained from a co-located chemiluminescence analyzer. The concentrations of Sulfur and nitrogen dioxide in the ambient air during the period from November 2009 to September 2010 are analyzed.

Results: The highest value for annual NO2 and SO2 averages of passive sampler was 29.65 μg/m³ and 21.01 μg/m³ for exposures of 2-weeks at an industrial site. The maximum monthly concentration for SO2 was observed at the 10th measurement station with 44.19 μg/m³ for August. The maximum monthly concentration for NO2 was observed on the 3rd measurement station with 42.83 μg/m³ for November. A negative correlation between nitrogen dioxide concentrations and temperature (R² = −0.5489) was estimated. A positive correlation between nitrogen dioxide measurement with passive sampler and continuous measurement (R² = 0.6571) was estimated.

Keywords: Passive sampler, Sulfur dioxide, Nitrogen dioxide

Background

Industrial establishments operating in the Samsun Industrial Zone for years, activities of the mobile power plants opened but then closed down and natural gas combined cycle plant scheduled to be reactivated require detailed studies of the region in terms of air pollution. The readings of the measurements made with uninterrupted system measurement devices at one single location revealed that the limit values of the European Union were exceeded seasonally in terms of nitrogen dioxide. However sulfur dioxide values did not exceed national and international legislations in any period (See Table 1) [1].

In the first instance the passive sampler was developed in America as an on-person air sampler by Palmes et al. (1976), it ensures that gas or vapor on the atmosphere is carried by molecular diffusion with a physical process and kept in a membrane. Here, the effective power is concentration change between the absorption surface and the air surrounding it and the amount of diffusion occurred is given on 1st Fick rule on equation (1) [6].

\[ F = -D \frac{dC}{dL} \]  

Where; F is the molar flux (μg.cm⁻².min⁻¹), D is the diffusion coefficient (cm²/min), C is the concentration (μg.cm⁻³), and L is the diffusion path (cm). The contaminant concentration kept in the passive sampler depends on the period of atmosphere exposure, quantity of total gas transfer collected in the passive sampler tube, geometric structure of the tube and diffusion coefficient, ambient temperature, ambient humidity, sorbent strength and face velocity. Passive sampler is a preferred method since it does not require power supply, has cost-efficient investment, light, simple analysis procedure as well as not requiring calibration, and suitable for simultaneous multipoint measurement of ambient air pollution. Passive samplers are extremely useful to assess long-term concentration trends (e.g., yearly) and are small, light, re-usable, and...
Nitrogen dioxide has an irritant effect on the respiratory organs, and long exposures can increasingly lead to airway disorders, such as chronic bronchitis. The major health impact of sulfur dioxide include effects on breathing, respiratory illness, weakness of lung defenses, increase in the effects of existing respiratory and cardiovascular disease, and death. Explanations of effects of pressure, temperature, and wind speed on the samplers for NO2 and SO2 have been reported. Diffusive sampling can be used if the average, instead of the real-time, pollutant concentration is adequate for the purpose of monitoring.

### Methods

The measurements took place at an industrial zone located in Samsun Tekkekokoy district, Selyeri locality. This district is located on the north-east coast of Turkey. The area is enclosed in the north by the Black sea. The area is also crossed by one of the main Samsun and Turkey highways, the D010 (Figure 1, yellow line). The Industrial zone is nearly 15 km away from the city center.

| Pollutants          | Period  | RG [2] | EU [3] | WHO [4] | NAAQS [5] |
|---------------------|---------|--------|--------|---------|-----------|
| **Sulfur Dioxide**  | 1-hour  | 57.25  | 350    | -       | 75        |
|                     | 24-hour | 47.71  | 125    | 7.63    | -         |
|                     | 3-hour  | -      | -      | -       | 500       |
|                     | Annual  | 7.63   | -      | -       | -         |
| **Nitrogen Dioxide**| 1-hour  | 53.14  | 200    | 10.63   | 100       |
|                     | 24-hour | -      | -      | -       | -         |
|                     | Annual  | 10.63  | 74.40  | 21.26   | 53        |

The Industrial Zone digital map of Samsun Tekkekooy District, scale 1:50000, is used at the GIS-software ArcGIS (version 9.2, ESRI, CA, USA). The measurement results were introduced in the ESRI Geographic Information System software ArcGIS and the extension geostatistical Analyst was used to obtain distribution maps. The geostatistical Analyst, an extension to ArcGIS, a product of the Environmental Systems Research Institute (ESRI), California, USA, can be applied to analyze the data from air quality monitoring networks and to generate maps of spatial distribution of the monitored air pollutants. Several spatial interpolation methods available were tested. The selected method was the geostatistical interpolation method-linear interpolation for commonly used.

For the pilot study, there were 10 samplers available for field sampling and the sampling area was approximately 20 km²; therefore; samplers were distributed through background locations only using a systematic grid of 1×1 Km, as can be seen in Figure 1. The distances between sampling points are approximately 1 – 1.5 kilometers. The samplers of site were identified and tracked using a Global Positioning System (Magellan Mobile Mapper TM6). The data on measurement locations are given on Table 2.

The passive samplers used for the measurements are from English Gradko Company and the technical specifications are explained in detail on Table 3. The analysis of nitrogen dioxide and sulfur dioxide passive samplers were conducted at Gradko’s laboratory in England. The analysis is carried out according with Gradko International Ltd Internal Laboratory Quality Procedure. All air analyses have been all conducted by UKAS Accredited Methods, working to ISO 17025. The calibration of sampling tubes was made with Gradko Environmental Laboratory accredited by United Kingdom Accreditation
Table 2 Measurement points and coordinates

| Station no | Coordinates | N   | E   |
|------------|-------------|-----|-----|
| 1          | 41°14'22.05" | 36°26'24.4" |
| 2          | 41°13'46.7"  | 36°26'43.7" |
| 3          | 41°13'24.5"  | 36°27'18.2" |
| 4          | 41°12'51.4"  | 36°28'7.7"  |
| 5          | 41°14'04.9"  | 36°26'40.52" |
| 6          | 41°15'22.2"  | 36°29'31.1" |
| 7          | 41°13'50.9"  | 36°28'51.9" |
| 8          | 41°13'57.1"  | 36°29'57.1" |
| 9          | 41°13'48.9"  | 36°28'58.9" |
| 10         | 41°14'13.8"  | 36°27'46.3" |

Table 3 Properties of passive samplers

| Properties                | Nitrogen Dioxide | Sulfur Dioxide |
|---------------------------|------------------|----------------|
| Model Number              | DIF 100 RTU      | DIF 600 RTU    |
| Absorbent                 | % 20 Triethanolamine/Deionized Water | Potassium Hydroxide |
| Method of Analysis        | UV/VIS Spectrophotometric | Ion Chromatography |
| Tube Size                 | 71 mm diameter   | 71 mm diameter |
| Duration of Exposure      | 14 day           | 14 day         |
| Amount of Air Intake      | 72 cm³ hour⁻¹    | 0.95 cm³ hour⁻¹ |
| Average Measurement       | +/-% 3.69        | +/-% 12.38     |
| Accuracy                  |                  |                |
| Lower Detecting Limit     | 0.34 ppb         | 0.5 ppb        |
| Reference                  | TS EN 13528/1-3   | TS EN 13528/1-3 |

Service (UKAS). The concentrations of Nitrite ions and hence NO2 chemically adsorbed (the grid made of stainless steel impregnated with 20% triethanolamine) are quantitatively determined by UV/Visible Spectrophotometry (Camspec M550 Spectrophotometer) with reference to a calibration curve derived from the analysis of standard nitrite solutions (UKAS Accredited Methods) which is measured colorimetrically. Likewise, for the sulfur dioxide analysis, the grid made of stainless steel impregnated with potassium hydroxide was extracted and then measured with Dionex IC 3000 ion chromatography. Determined by an overall m.o.u of 14.081% +/- and a limit of detection of 0.026 μg S for sulfur dioxide. Determined by an overall m.o.u of 10.907% +/- and a limit of detection of 0.021 μg NO2 for nitrogen dioxide.

In order to compare passive samplers with the results of uninterrupted system measurement method operated with chemical optical spectroscopic methods, the data of the measurement station owned by the Ministry of Environment and Forestry was used (Station No.3). The Station No.3 where the reference method chemiluminescence and UV Fluorescence method for automatic measurement of NO2 and SO2 concentrations and are in use. An automatic ambient SO2 analyzer (AF22M, Environment S.A, France) and a ambient NO2 analyzer (AC32M, Environnement S.A, France) were used to measure the average concentrations of sulfur dioxide and nitrogen dioxide. The fixed measurement station data was obtained from Air Quality Monitoring Network database. The data were compiled as 14-day average data and then compared with the passive sampler readings.

The measurements started on 18.10.2009 and were completed on 19.09.2010. The measurements were made at total 10 measurement stations. Measurements were performed twice each month to account for the seasonally changing climatic conditions. The passive samplers were located far from walls and 2.0 m above the ground. While selecting the locations of passive samplers used for the measurements, several particulars were taken into consideration such as; exactly determining the industrial contamination in the region, taking into consideration the impact areas by bearing in mind the dominant wind direction, taking into consideration the impacts of significant point contaminants, preventing the impact of pollution caused by vehicles, minimizing the intervention on parks and of wind, comparing the results with uninterrupted measurement system and ensuring ease and safety of passive sampler installation.

Results and discussion

The measurement readings obtained for nitrogen dioxide and sulfur dioxide from all measurement stations are given separately on Figures 2 and 3, and seasonal variation of sulfur dioxide and nitrogen dioxide for all stations on Figures 4 and 5. The highest pollutant concentrations were in the 1., 2., and 10 number of stations. The highest average concentrations were in winter for nitrogen dioxide and sulphur dioxide (Figures 4 and 5).

When a digital map of the readings regarding monthly averages of sulfur dioxide measurements between October 2009 and October 2010 was formed by using ArcGIS software, it was observed that the values obtained in August 2009 were the highest and reached up to 154.49 μg/m³ (Figure 6). As for the sulfur dioxide; the overall result observation the digital maps prepared by using ArcGIS software was that the level of pollution at measurement locations number 1, 2, 3, 4, 5 and 10, in other words on the south-west and north-west of the region, is higher.

According to the measurement readings calculated for nitrogen dioxide, the highest NO2 values reached up to 43.27 μg/m in November 2009 (Figure 7).
Figure 2 The values of SO₂ and NO₂ for 1 – 5 stations.

Figure 3 The values of SO₂ and NO₂ for 6 – 10 stations.
Figure 4 Average values of SO\textsubscript{2} concentrations for seasonal variation during 2009–2010.

Figure 5 Average values of NO\textsubscript{2} concentrations for seasonal variation during 2009–2010.
Figure 6 Results of measurement of average sulfur dioxide for August 2010 (as the highest mount).

Figure 7 Results of measurement of average nitrogen dioxide for November 2009 (as the highest mount).

Figure 8 Dispersion of annual average NO₂ and SO₂ pollutants in Samsun in 2009–2010.
The highest value for annual NO\textsubscript{2} averages was 29.65 \(\mu\text{g/m}^3\) and 21.01 \(\mu\text{g/m}^3\) for SO\textsubscript{2} (Figure 8).

In order to compare the results obtained with passive sampler method with the measurements made with uninterrupted measurement devices, passive samplers were located at the fixed measurement station of Ministry of Forestry and Water Works, namely Sampling Station Number 3. The measurement readings were obtained from Air Quality Monitoring Network web page as the reference method of chemiluminescence. The arithmetical averages of the 14-day exposure period of the passive samplers were calculated by taking into consideration the daily arithmetical averages of the values. The results of sulfur dioxide and nitrogen dioxide according to two measurement methods so compared are given on Figure 8.

Statistically, regression analysis carried out to show the degree of correlation between the continuous monitor and the diffusion tubes gave different regression coefficients. Correlation analysis between passive samplers and the continuous monitor (Figure 9) gave a correlation coefficient \((R^2)\) of value. According to the uninterrupted system and passive sampler methods used at measurement station number 3, there is a low-level relationship in terms of sulfur dioxide measurement methods \((R^2 = 0.1827)\) but the relation in terms of nitrogen dioxide measurements is confirmed to be \(R^2 = 0.6571\) (Figure 9).

Box-plots of yearly concentrations of SO\textsubscript{2} and NO\textsubscript{2} measured at the sampling points are given in Figure 10.

| Date          | Monthly average inputs |
|---------------|------------------------|
|               | Wind direction \(^{\circ}\) | Wind speed (m/s) | Temperature (\(^{\circ}\)C) | Relative humidity (%) |
| October 2009  | 170                     | 2.55             | 17.4                         | 84.2                  |
| November 2009 | 180                     | 3.06             | 11.7                         | 78.8                  |
| December 2009 | 170                     | 3.06             | 10.7                         | 71.0                  |
| January 2010  | 180                     | 3.57             | 8.9                          | 72.4                  |
| February 2010 | 190                     | 3.57             | 9.8                          | 73.6                  |
| March 2010    | 190                     | 3.06             | 8.0                          | 81.8                  |
| April 2010    | 190                     | 3.06             | 11.6                         | 82.8                  |
| May 2010      | 160                     | 2.55             | 17.1                         | 76.6                  |
| June 2010     | 200                     | 3.06             | 22.0                         | 80.7                  |
| July 2010     | 220                     | 3.06             | 24.9                         | 80.5                  |
| August 2010   | 190                     | 2.55             | 26.1                         | 75.9                  |
| September 2010| 180                     | 2.55             | 21.3                         | 80.4                  |
Conclusions

The conclusions of this study are listed below;

1. The uninterrupted measurement station values and passive sampler readings are compared at the measurement station of the Ministry of Forestry and Water Works. A correlation of 69.24% is determined during the comparisons. When the value is corrected according to this correlation, the annual nitrogen dioxide average is calculated as 42.36 μg/m³. This value for NO2 was below the national air quality standard. But, was above the air quality standard of the European Union. However, as mentioned on the literature, meteorological parameters have significant impact on the correlation and it was not possible to make meteorological measurements at the measurement location during this study.

2. When the digital map regarding the nitrogen dioxide measurements are reviewed, it is observed that the highest level of pollution is at measurement station of Station No.3 and expands around from that location. Besides, the nitrogen dioxide values of the measurement point Number 5 at Yesilyurt Port Administration vary between 26 to 29 μg/m³.

3. It is observed that the nitrogen dioxide is intensified at locations 1, 2, 5 and 10 on the west side, starting from measurement station number 3 and expanded towards the Black Sea from location 5 or towards the inner regions of Tekkeköy from location 3.

4. When the passive samplers are compared with uninterrupted measurement results at the measurement location number 3, in terms of sulfur dioxide, a correlation could not be established. Since the results obtained from both passive samplers and uninterrupted measurements are significantly low, we should consider the possibility of errors arising from factors having an impact on sulfur dioxide measurements and devices used. The reason for this, the sulfur dioxide concentration effected not only passive sampler measurements but also meteorological parameters.

5. It is observed that the sulfur dioxide distribution heads from south to north-west, towards the Black Sea from measurement station number 5.

Competing interests

The author declares that he has no competing interests.

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