

Original Research

Air Pollution Origins Using PM₁₀ Data and CO₂ Isotopic Analysis

Anna Zwoździak^{1*}, Maciej Górka², Izabela Sówka¹, Dominika Lewicka-Szczebak²,
Jerzy Zwoździak¹, Mariusz Orion Jędrysek²

¹Ecologistics Division, Institute of Environmental Protection Engineering, Wrocław University of Technology,
Pl. Grunwaldzki 9, 50-377 Wrocław, Poland

²Laboratory of Isotope Geology and Geoecology, Department of Applied Geology and Geochemistry,
Institute of Geological Sciences, University of Wrocław,
Cybulskiego 30, 50-205 Wrocław, Poland

Received: 11 February 2010

Accepted: 30 April 2010

Abstract

PM₁₀ concentrations (in relation to main gaseous pollutant levels and prevailing meteorological conditions) and carbon isotopic composition of atmospheric CO₂ have been analysed in two health resorts in SW Poland (Cieplice and Czerniawa) to determine the main air pollution sources. In Cieplice, PM₁₀ concentrations followed the variations in NO₂ and CO, as well as SO₂, suggesting traffic and domestic heating to be the main sources of aerosol particles. Conversely, in Czerniawa periodically pollutant rich air masses originating from regional sources contribute to increases in SO₂ and NO₂ levels, whereas the PM₁₀ concentrations are at the background level. The analyzed $\delta^{13}\text{C}(\text{CO}_2)$ values reinforced the conclusions drawn on the basis of traditional methods and indicated the different impact of pollution sources between the investigated regions. Czerniawa was controlled by regional sources of CO₂, whereas Cieplice was dominated by multiple local sources of pollutants (mainly traffic pollutants).

Keywords: PM₁₀, SO₂, NO_x, episode, carbon isotope

Introduction

Lower Silesia is a province in southwestern Poland located at the contact point of Polish, German, and Czech Republic borders. The presence of old and diverse geological structures makes this region rich in various natural resources, like copper ores, coal, brown coal, and mineral waters of proved healing properties. But simultaneously, the exploration and processing of natural resources, which are fundamental for the local economy, appeared to be – indirectly – the source of environmental deterioration. In the 1980s and '90s the region, where the borders of Poland,

Germany, and the Czech Republic adjoin, was called the “Black Triangle.” It was considered one of the most polluted regions in Europe because of numerous power plants without efficient gas cleaning techniques, wide open coal mining areas, and domestic heating based on brown coal. Since the early 1970s deforestation has been the major concern due to the extent and severity of the observed damage. Massive areas of forest suffered serious deterioration presenting strong visual symptoms of ecological damage [1, 2]. After the political changes great effort was made to reduce the emissions from the highest industrial companies. Permanent air monitoring carried out since 1997 proved the evident decrease of gaseous and particulate pollution [3, 4].

The overall results of our field studies in two health resorts located in the Polish part of the former “Black

*e-mail: anna.zwozdziak@pwr.wroc.pl

Triangle" area [5] indicated that the monthly average concentrations of SO₂, NO_x, and PM₁₀ (particulate matter diameter below 10 μm) have been decreasing since 1996, but episodically both local and regional influences are still evident. Air quality in these two health resorts has improved significantly compared to the 1990s. Nevertheless, pollution by PM₁₀ is believed to remain a concern.

Based on the above-mentioned survey, we have undertaken a more detailed evaluation of air pollution sources. We have analyzed PM₁₀ concentrations in relation to main gaseous pollutant levels and prevailing meteorological conditions. Moreover, we report the results of the atmospheric carbon dioxide analyses concerning its concentration and carbon isotopic signatures, comparing them with the above described investigations. The geochemical and isotopic analyses of CO₂ have been carried out as a part of complex investigations of atmospheric pollutants in that area [5, 6]. The major goals of our investigations presented here are:

- (i) to indicate the processes that probably influence the pollutant concentrations and carbon isotopic composition of atmospheric CO₂;
- (ii) to assess air quality in the investigated area.

Isotopic analyses enable tracing of the pollution origin. The total concentration and carbon isotopic composition of atmospheric carbon dioxide consists of background gases and additional inputs of anthropogenic and/or biogenic gases [7, 8]. The biogenic gases are derived from plant leaf respiration or from two pathways for production of CO₂ in soil. The biogenic isotopic fractionation factor depletes atmospheric CO₂ through absorption by terrestrial photosynthesis. This results in the shift of δ¹³C value of C3 plants from -16 to -20‰ and of C4 plants from -2 to -8‰ [9]. Therefore, the δ¹³C(CO₂) values derived from C3 plant respiration processes at night (biogenic gases) can reduce to -20‰ [10, 11]. Carbon dioxide produced in the inter soil processes can originate from two main sources: root respiration and decomposition of organic matter by soil microorganisms [12]. The δ¹³C(CO₂) values originated from the first inter soil process reach values close to the whole plant respiration, i.e. ca. -20‰. On the contrary, the second inter soil process generates the CO₂ enriched in ¹³C [12]. In contrast, the δ¹³C(CO₂) values derived from burning fossil fuels (anthropogenic sources) obtained by Widory and Javoy [13] varied in the wide range from -40.5‰ (natural gas burning fumes) to -24.6‰ (coal burning fumes). In winter the photosynthesis input is almost negligible (non-vegetation period), hence the CO₂ is derived from anthropogenic sources (heating period) or soil respiration [14, 15]. Conversely, in summer season the anthropogenic input is negligible and the biological input is dominant [8]. Obviously, the natural system is more complicated; usually a combination of all the above processes can be observed [15]. Therefore, our winter observations, when biogenic input did not complicate the whole system, are the most useful in order to correctly access the processes controlling air quality in the region under study.

Experimental Procedures

Study Area

Field sampling campaigns were performed in July 2006 and 2007, and in February 2007 and 2008 in order to represent two different seasons: summer and winter. Two sites were selected to represent health resort areas: Cieplice (50°54'N, 15°44'E, 330 m asl) and Czerniawa (50°54'N, 15°20'E, 645 m asl) situated in SW Poland (Lower Silesia region) in the Sudety Mts. Cieplice is part of Jelenia-Góra city (about 90,000 inhabitants), whereas Czerniawa (30 km west of Cieplice) is part of Świeradów (about 5,000 inhabitants). Świeradów health resort with the Spa House is situated 200 m below the Czerniawa site. Moreover, data from other two monitoring stations representing regional rural background: Wlen (51°00'N, 15°40'E, 303 m asl) and Jeleniów (51°13'N, 15°15'E, 244 m asl) were also used to compare the temporal and spatial variations in PM₁₀ concentrations in order to recognize the local and regional influences. The location of the sampling sites is shown in Fig. 1.

NO₂, O₃, CO, SO₂, and PM₁₀ Sampling Methodology

Air quality data (NO_x, O₃, CO, SO₂, and PM₁₀) and meteorological parameters were obtained from the Voivodeship Inspectorate for Environment Protection in Wrocław (www.wroclaw.pios.gov.pl). PM₁₀ measurements were conducted with the Environnement S.A. MP101M monitor (based on β-attenuation). Continuous trace gas measurements of NO_x (NO and NO₂), SO₂, CO, and ozone were made using chemiluminescence for NO_x (Environnement S.A., Model 142M), UV fluorescence for SO₂ (Environnement S.A., Model 22M), infrared gas filter correlation for CO (Environnement S.A., Model 12M), and UV absorption (Environnement S.A., Model 41M) for ozone, respectively.

Meteorological parameters measured during the period of field studies were wind speed and direction, temperature, relative humidity, pressure and solar radiation, which were obtained from an automatic station equipped with LSI-LASEM weather sensors and installed near the sampling point. The backward trajectories of the air masses were calculated by the HYSPLIT [16] model run by the U.S. National Oceanic and Atmospheric Administration (<http://www.arl.noaa.gov>).

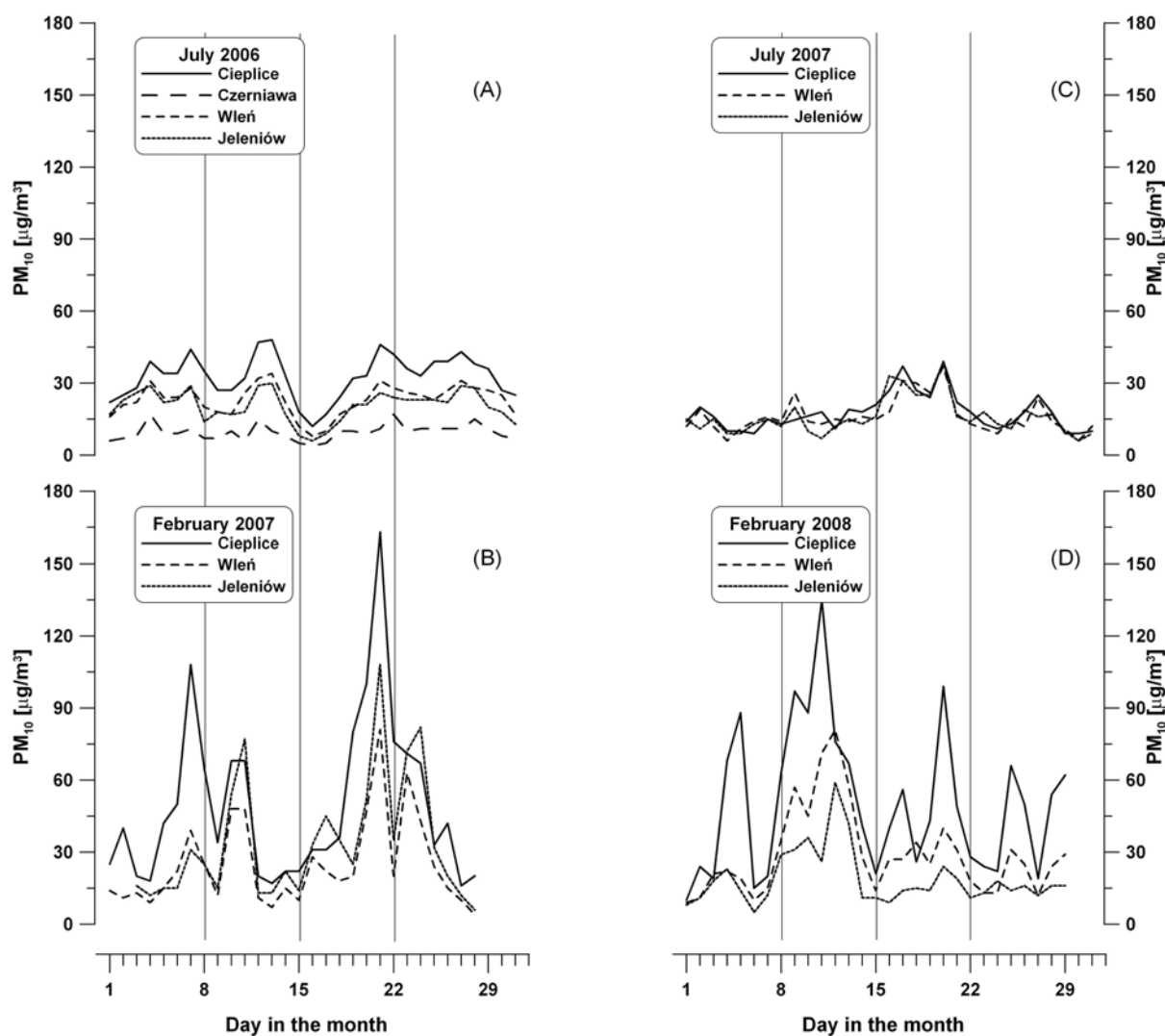
All data were quality-controlled in the data management network system operated by the Voivodeship Inspectorate for Environment Protection in Wrocław.

CO₂ Sampling Methodology

The atmospheric samples were collected using glass (July 2006 and February 2007) or steel (July 2007 and February 2008) 2-litre vacuum containers [17]. Each container was evacuated to about 10⁻² hPa in the laboratory before field usage. In each sampling action the flask stopcock was opened into the wind direction and the air was



Fig. 1. Location of sampling points.

Fig. 2. Daily concentrations of PM_{10} at the selected monitoring stations in July 2006 and 2007, and February 2007 and 2008.

sucked in. After filling, the flask was immediately closed and transported to the laboratory within a few days (max. 3 days) for further carbon isotopic analysis of atmospheric CO₂.

The atmospheric CO₂ concentration [ppm] and temperature [°C] were measured using a Telaire 7001 portable instrument.

Carbon Isotopic Analysis

From air samples for carbon isotopic analyses CO₂ was obtained by cryogenic purification and collected in hermetic glass ampoules [10, 11, 18]. Carbon isotopic ratios were measured using the Finnigan Mat Delta E mass spectrometer (Laboratory of Isotope Geology and Geoecology at University of Wrocław, Poland). The standard uncertainty of the δ¹³C determination was less than 0.1‰. IAEA NBS-22 and USGS-24 standards were used as the carbon stable isotope standards. The additive correction of +0.22‰ was included for N₂O contribution in each final calculated δ¹³C(CO₂) value [19, 20].

Results and Discussion

General Weather Conditions during Field Studies

The summers of 2006 and 2007 (campaigns I and III) were characterized by sunny weather and high air temperature. In winter 2006/07 and 2007/08 (campaigns II and IV) the weather conditions were rather cloudy, with foggy and snowy episodes. However, days with the rebuilding baric field and the growth of Eastern European anticyclone with the centre over Ukraine were also observed. Air temperature in campaigns II and IV oscillated about 0°C. Minimum daily air temperature was reported on 13 February 2008 in Czerniawa (-0.4°C), while the maximum was 22 July 2006 in Cieplice (32.4°C).

Spatial and Temporal Variations of PM₁₀ Data

Diurnal time series of PM₁₀ (particulate matter, diameter below 10 μm) concentrations registered in July 2006 and 2007, and February 2007 and 2008 in the selected air monitoring stations are shown in Fig. 2. For Czerniawa these results were available only for July 2006 and have not been measured afterwards. Therefore, data from the other air monitoring stations were used to show the spatial and temporal variations in PM₁₀ concentration and to see how comparable the sites are within the area under study. Clearly, there is a great similarity in temporal variations of the PM₁₀ concentration between the sites, although the data show that during both summer and winter the PM₁₀ concentrations in Cieplice are higher than those in the other sites. The daily EU PM₁₀ standard of 50 μg/m³ [21] was exceeded several times a month (allowing 35 days per year).

However, the regional background of PM₁₀ levels registered in Wlen and Jeleniow was also very high during some episodes in winter. During wintertime, an extremely high PM₁₀ concentration (up to 175 μg/m³ in February 2007) was noticed in Cieplice, most likely due to the high load of local pollutants and local meteorology. During summertime, the PM₁₀ concentrations significantly decreased both in Cieplice (up to 50 μg/m³ in July 2006) and in Jeleniow and Wlen (up to 35 μg/m³ in July 2006).

Comparable temporal patterns in the PM₁₀ concentration for all sites can be explained as being strongly influenced by similar dust sources and/or similar meteorological characteristics of the area [22-24]. In order to understand these processes, we had to analyze the different trends of PM₁₀ levels, taking into account the gaseous pollutants and some meteorological factors. Figs. 3 and 4 show two examples of air pollution episodes registered in Cieplice and Czerniawa during the field studies. Similar patterns were also recorded in July 2007 and February 2008 (not presented here).

On 20-21 July 2006 (Fig. 3) high PM₁₀ (up to 55 μg/m³) and very high ozone concentrations (up to 180 μg/m³) were noticed in Cieplice, a result of which was most likely due to stagnant conditions with weak winds combined with high temperatures and high solar radiation. The region was in a low pressure gradient between two high pressure areas, one centred east of Poland (Ukraine) and the

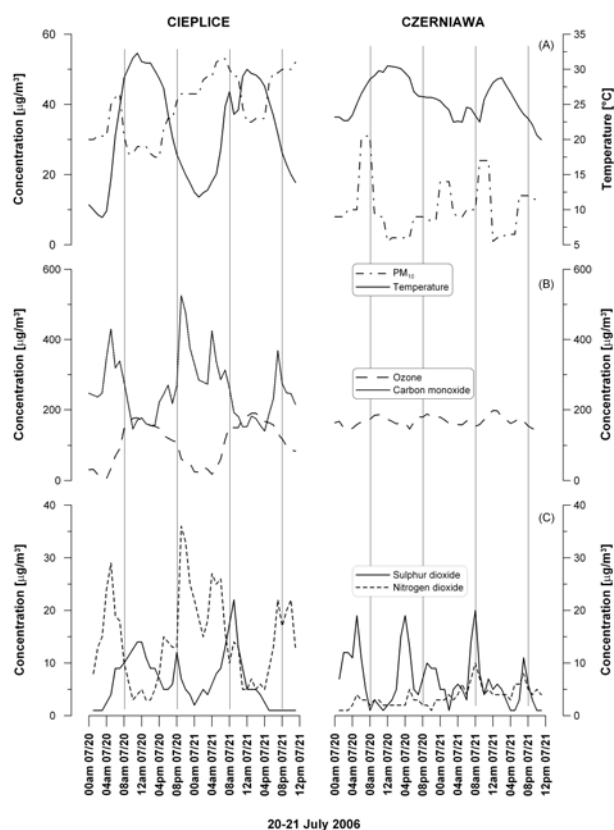


Fig. 3. Temporal variations of temperature and pollutant concentrations in Cieplice and Czerniawa on July 20-21, 2006. All hours are in local time.

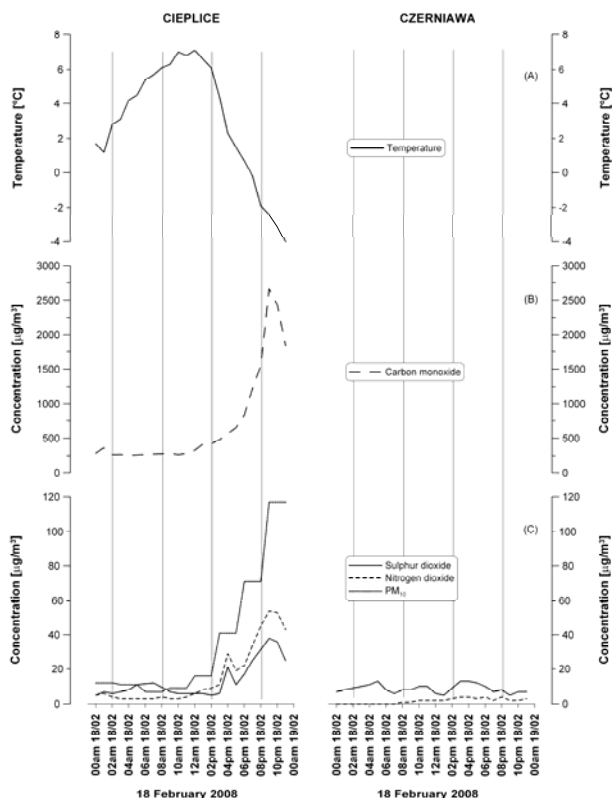


Fig. 4. Temporal variations of temperature and pollutant concentrations in Cieplice and Czerniawa (only SO₂ and NO₂) on February 18, 2007. All hours are in local time.

second west of Poland (the Alps). Large temperature differences occurred between day and night (up to 22°C) in Cieplice, which indicated intensive cooling processes during the night. In general, the radiational cooling of the surface at night generates a temperature inversion. The magnitude

of the inversion can be estimated based on simultaneous temperature measurements at stations located at different altitudes. The temperature difference (12°C) between Czerniawa and Cieplice can be deduced from data in Fig. 3. These data confirmed a substantial inversion over the city that reduced air turbulence and led to a well-known accumulation of pollutants as observed in Cieplice. This process seems to be responsible for early morning (5:00-6:00 LT) and late evening (20:00-21:00 LT) CO and NO₂ peaks, rather than rush hour traffic, which occurs later in the morning (8:00-9:00 LT) and earlier in the evening (17:00-18:00 LT). In addition to these local influences, regional pollutants sometimes can be recognized. The most prominent SO₂ event took place (simultaneously in Czerniawa) on the morning of 21 July, when the SO₂ levels reached 22 µg/m³, rising from 5 µg/m³. At the same time, NO₂ (but not CO) concentrations noted a slight peak. Air masses contained more SO₂ and NO₂ but were not distinctly depleted in ozone. This episode was associated with trajectory from W-SW (Fig. 5), and most probably originated from lignite power plants located 30 km in that direction.

In Czerniawa the pollutants showed markedly different patterns. The variations in the ozone level proved that we dealt with different air masses in Cieplice and Czerniawa. At night, when temperature inversion occurred, the ozone concentration in the valley decreased more efficiently, but in Czerniawa (300 m higher) its concentration was still high. This behaviour indicates that Czerniawa (650 m asl) lies above the nocturnal boundary layer. The variations in the SO₂ and PM₁₀ concentrations over the 24-hour period show significant diurnal increases and decreases, which probably reflect the interplay between regional sources' impact and daily evolution of the boundary layer. The confirmation of the impact from regional sources may be the simultaneous significant increase in SO₂ and NO₂ levels,

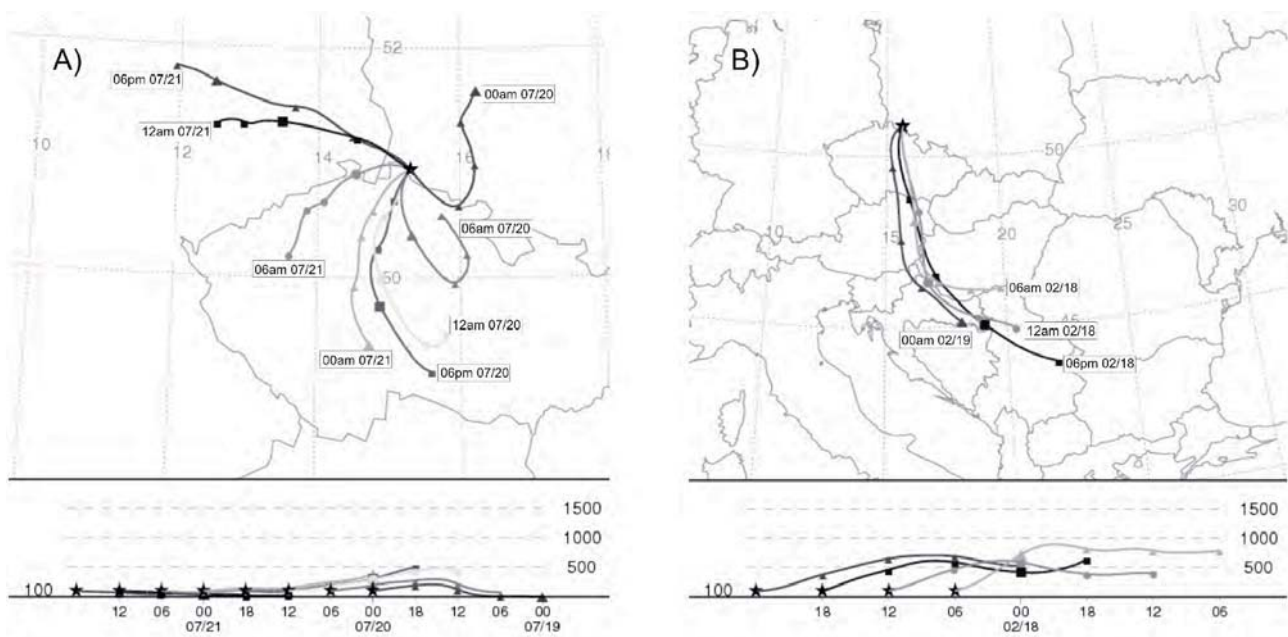


Fig. 5. Vertically modelled transport back-trajectories from (a) an altitude of 700 m asl (Czerniawa 650 m asl) ending at 18:00 UTC (+2 h – local time) 21 July 2006 and (b) an altitude of 330 m asl (Cieplice) ending at 00:00 UTC (+1 h – local time) 19 February 2007.

more evident on 21 July than 20 July. On 20 July air masses arrived from southern directions (Fig. 5a, the trajectory 06am07/21), and the next day from western directions. High SO_2 values were observed at both Czerniawa and Cieplice associated with winds from the west, where the lignite power plant is situated, being the main source of gaseous pollutants in that area.

Fig. 4 shows data for 18 February 2007, which gives an example of a local-emission-dominated winter-pollution episode with high concentration of PM_{10} (up to $117 \mu\text{g}/\text{m}^3$) and both traffic-generated pollutants (NO_2 , CO) and wintertime central heating (SO_2). An area of high pressure was centered over the Ukraine, leading to very low pressure gradients and low surface winds. The continuous process of sinking observed on February 18 (Fig. 5b) was attributed to the existence of large-scale anticyclonic subsidence. Because of the lower ventilation factor (mixing layer height (MLH) and wind velocity – weak air flow and a shallow mixing depth (MD)) the measured concentrations of all components were very high in Cieplice, most probably due to accumulation of local pollutants. On the contrary, we dealt with different air masses in Czerniawa, where regional SO_2 and NO_2 background concentrations have been noted.

Concentration and Carbon Isotopic Composition of Atmospheric CO_2

The obtained CO_2 concentration and carbon isotopic signatures of carbon dioxide ($\delta^{13}\text{C}(\text{CO}_2)$ values) show a seasonal variability (Fig. 6). Minimum value in atmospheric CO_2 concentration was noted on 13 February 2008 in Czerniawa (349 ppm) and a maximum on 17 July 2007 in Cieplice (566 ppm). Minimum $\delta^{13}\text{C}(\text{CO}_2)$ values were noted on 17 February 2007 in Cieplice (-11.51‰) and maximum in July 2006 in Cieplice (-9.12‰). The concentration and $\delta^{13}\text{C}(\text{CO}_2)$ values reported in this study are similar to those reported by other authors for southern Poland [7, 25–27].

The observation of the concentration and $\delta^{13}\text{C}(\text{CO}_2)$ values are summarized in Fig. 6. The data of CO_2 concentration and $\delta^{13}\text{C}(\text{CO}_2)$ values measured at Czerniawa in summer season (Fig. 6) reach similar values in both summer periods (2006 and 2007). In contrast, the data of CO_2 concentration and $\delta^{13}\text{C}(\text{CO}_2)$ values measured at Cieplice in summer 2007 show higher CO_2 concentration and lower $\delta^{13}\text{C}(\text{CO}_2)$ values than those obtained in summer 2006. Probably, we observed a larger input of air masses characterized by more anthropogenic character [8, 13] in summer 2007 in Cieplice. These theoretical considerations are confirmed by episodes and air mass trajectory observations for the summer period, which have indicated the regional sources (mainly lignite power plant) as the major impact controlling air quality in the region.

The $\delta^{13}\text{C}(\text{CO}_2)$ values obtained at Czerniawa in winter (Fig. 6) are characterized by similar values in both 2007 and 2008. This suggests the same level of anthropogenic impact

(biogenic is negligible in non-vegetative season) in both years. However, the winter data of CO_2 concentrations measured in Czerniawa show a radical decrease in CO_2 concentration (about 100 ppm) in 2008. The very similar carbon isotopic composition of CO_2 in both years, and the contemporary decrease of CO_2 concentration indicate a substitution of one anthropogenic source by another during the 2-year observations. In contrast, the data of CO_2 concentrations measured in Cieplice in winter (Fig. 6) suggest the improvement of air quality between 2007 and 2008. This hypothesis can be tested more precisely by the simultaneous analysis of concentrations and carbon isotopic composition of CO_2 . For the sample at Cieplice the lower CO_2 concentration (about 100 ppm) in 2008 is associated with higher $\delta^{13}\text{C}(\text{CO}_2)$ values (about 1.5‰). We postulate that it is caused by lower anthropogenic impact concerning the fuels (coal and wood) burning contribution in the total balance of CO_2 . These theoretical considerations are confirmed by episodes and air mass trajectory observation for the winter period, which indicate the regional background

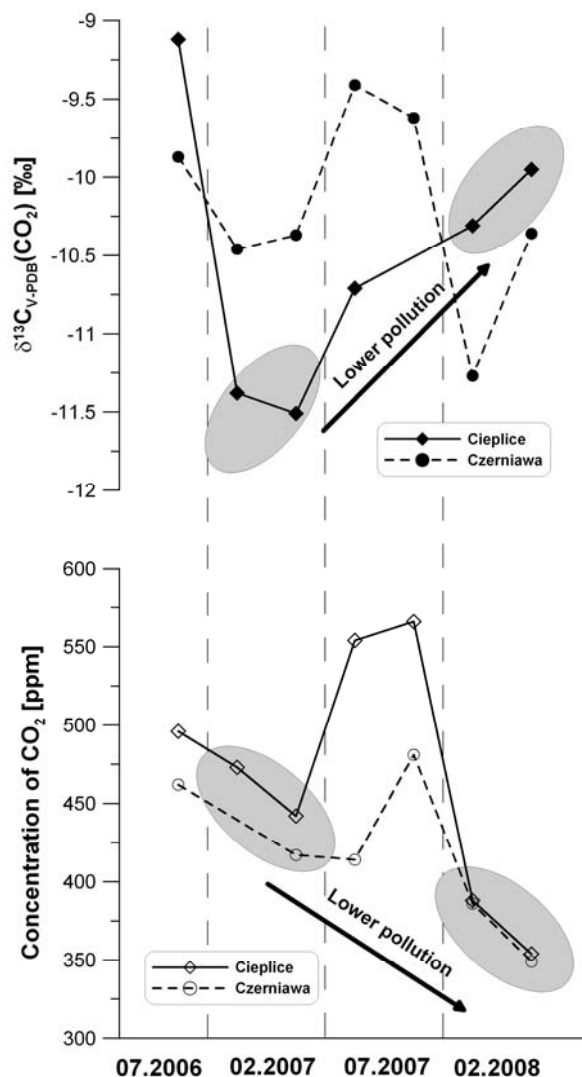


Fig. 6. Carbon dioxide concentrations and $\delta^{13}\text{C}(\text{CO}_2)$ values in Czerniawa and Cieplice between July 2006 and February 2008.

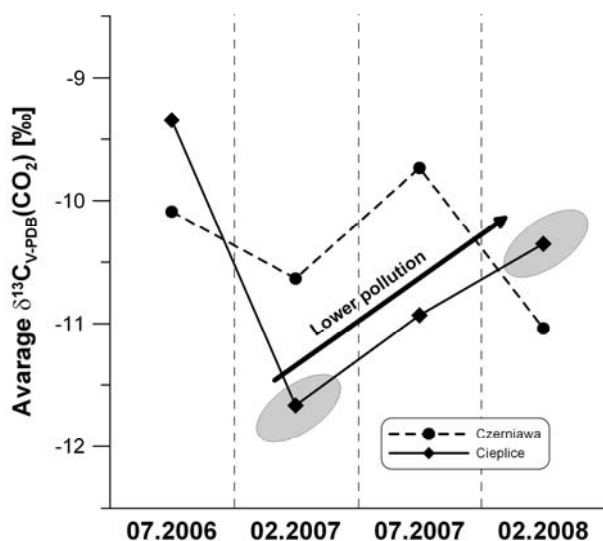


Fig. 7. Average $\delta^{13}\text{C}(\text{CO}_2)$ values in Czerniawa and Cieplice between July 2006 and February 2008.

sources influencing air quality in Czerniawa, whereas the air quality in Cieplice is controlled by local pollutants (central heating and traffic). Therefore, the obtained data of concentration and carbon isotopic composition of atmospheric CO₂ indicate the improvement of air quality only for Cieplice health resort during the 2-year monitoring period.

In order to better characterize the sampling environment and exclude accidental episodes, the average carbon isotopic signature of atmospheric carbon dioxide has been calculated. The arithmetic average values of all the samples obtained for each period are presented in Fig. 7.

The average carbon isotopic signatures for Czerniawa CO₂ samples most probably result from regional impacts (controlled by long-transport processes) and rather exclude the local influences. In contrast, the average isotopic composition of Cieplice samples has not indicated the clear seasonal repeatability (Fig. 7) that has been observed at Czerniawa. In our opinion, the irregular seasonal variations in Cieplice are caused by different mixing of local carbon dioxide sources and traffic pollutants in winter with occasional regional pollutant inputs in summer. The lack of typical seasonal variations suggests that the atmospheric pollution in Cieplice is governed mainly by local processes and the regional influences have minor influence on air quality. However, the average $\delta^{13}\text{C}(\text{CO}_2)$ data have confirmed the earlier conclusions concerning the improvement of air quality in Cieplice between 2007 and 2008.

Conclusions

In summary up, the data obtained from analyses of concentration and isotopic composition of the atmospheric CO₂ with comparison to traditional monitored pollutants (PM₁₀, NO_x, CO, SO₂, O₃) and meteorological conditions can provide comparable information about air quality.

The results about the seasonal data point out much higher PM₁₀ concentrations during the cold season than during the warm one, with the highest peaks observed at Cieplice in each sampling, exceeding the EU standards. Some meteorological (high pressure system, low winds, temperature inversion) and topographical (a relatively flat valley in Cieplice) factors were likely to be of significant importance during the episodes considered. In Cieplice the PM₁₀ concentrations followed the variations in NO₂ and CO in summer and in SO₂, NO₂, and CO in winter, suggesting that traffic could be the main source of aerosol particles in summer and domestic heating in winter. Conversely, Czerniawa was subjected to periodic pollutant-rich air originating from regional sources, which contributed to the increase in SO₂ and NO₂ levels, both in summer and winter months.

The data obtained from carbon isotopic analyses of CO₂ confirmed the different impact of pollution sources between the investigated regions. Czerniawa is mainly controlled by regional sources of CO₂. Conversely, Cieplice is dominated by multiple local sources of pollutants (mainly traffic pollutants). The seasonal characteristics of carbon dioxide pollutant indicate the improvement of air quality in Cieplice (in the 2-year monitoring period), and only a slight change of pollutants' impact in Czerniawa.

Stable isotopic analyses applied in this study constitute an innovative tool for tracing atmospheric pollution origin. Here we have demonstrated, coupling stable isotopic analyses with traditional monitoring of air quality, that the information obtained using both methods is consistent and complementary.

Acknowledgements

We would like to acknowledge Adriana Trojanowska PhD, Anna Worobiec de Maeyer PhD, Elżbieta Stefaniak PhD, and Anna Buczyńska MSc for their substantial help in our field work. The presented work was realized in the frame of Bilateral Project No. 4/1521, supported by the Flemish Administration for Innovation and Science (AWI) and State Committee for Scientific Research in Warsaw, and University of Wrocław grant Nos. 1017/S/ING/07-IX, 2022/W/ING/05-49, and 2022/W/ING/07-12.

References

- MATERNA J. Air pollution and forestry in Czechoslovakia, *Environ. Monit. Assess.* **12**, 227, **1989**.
- ZAWILA-NIEDZWIECKI T., ROCK B.N., GRECZYNSKY J., BOCHENEK Z. Landsat-based detection of change-over-time in forest conditions in the Sudety Mountains of Poland, In: *Global Change and Education*, ASPRS/ACSM/RT 92 Technical Papers, **1**, 281, **1992**.
- SPINDLER G., MULLER K., BRUGGEMANN E., GNAUK T., HERMANN H. Long-term size-segregated characterization of PM₁₀, PM_{2.5}, and PM₁ at the IFT research station Melpitz downwind of Leipzig (Germany) using high and low-volume filter samplers, *Atmos. Environ.* **38**, 5333, **2004**.

4. ABRAHAM J., BERGER F., CIECHANOWICZ-KUSZTAL R., JODŁOWSKA-OPYD G., KELLWOOD D., KEDER J., KULASZKA W., NOVÁK J. Joint report on air quality in the Tri-border region of the Czech Republic, Poland and Germany in 2004 (former Black Triangle region). ČHMÚ, WIOŚ, LfUG, UBA, (www.wroclaw.pios.gov.pl), **2005**.
5. WOROBIEC A., ZWOŹDZIAK A., SÓWKA I., ZWOŹDZIAK J., STEFANIAK E.A., BUCZYŃSKA A., KRATA A., VAN MEEL K., VAN GRIEKEN R., GÓRKA M., JEĐRYSEK M.O. Historical air pollution changes in the tri-border region of Poland, Czech Republic and Germany, *Environ. Prot. Eng.* **34**, (4), 81, **2008**.
6. GÓRKA M., JEĐRYSEK M.O., MAJ J., WOROBIEC A., BUCZYŃSKA A., STEFANIAK E., KRATA A., VAN GRIEKEN R., ZWOŹDZIAK A., SÓWKA I., ZWOŹDZIAK J., LEWICKA-SZCZEBAK D. Comparative assessment of air quality in two health resorts using carbon isotopes and palynological analyses, *Atmos. Environ.* **43**, 682, **2009**.
7. ZIMNOCH M., FLORKOWSKI T., NĘCKI J.M., NEUBERT R.E.M. Diurnal variability of $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ of atmospheric CO_2 in the urban atmosphere of Kraków, Poland, *Isotopes Environ. Health Stud.* **40**, (2), 129, **2004**.
8. LONGINELLI A. AND SELMO E. Seasonal and diurnal variations of $\delta^{13}\text{C}$ and concentration of atmospheric CO_2 at Parma, Italy, *Geol. Q.* **49**, (2), 127, **2005**.
9. MOOK W.G. ^{13}C in atmospheric CO_2 , *Neth. J. Sea Res.*, **20**, 211, **1986**.
10. SZARAN J. Seasonal variations of $\delta^{13}\text{C}$ values and CO_2 concentration in the air during vegetation growth, *Isotopes Environ. Health Stud.* **34**, 341, **1998**.
11. SZARAN J., NIEZGODA H., TREMBACZOWSKI A. Respiration and assimilation process reflected in the carbon isotopic composition of atmospheric carbon dioxide, *Nukleonika* **47**, (1), 59, **2002**.
12. AMUNDSON R., STERN L., BAISDEN T., WANG Y. The isotopic composition of soil and soil-respired CO_2 , *Geoderma* **82**, 83, **1998**.
13. WIDORY D., JAVOY M. The carbon isotope composition of atmospheric CO_2 in Paris, *Earth Planet. Sci. Lett.* **215**, 289, **2003**.
14. TAKAHASHI H.A., KONOHIRA E., HIYAMA T., MINAMI M., NAKAMURA T., YOSHIDA N. Diurnal variation of CO_2 concentration, $\Delta^{14}\text{C}$ and $\delta^{13}\text{C}$ in an urban forest: estimate of the anthropogenic and biogenic CO_2 contributions, *Tellus* **54B**, 97, **2002**.
15. DEMÉNY A., HASZPRA L. Stable isotope compositions of CO_2 in background air and at polluted sites in Hungary, *Rapid Commun. Mass Spectrom.* **16**, 797, **2002**.
16. DRAXLER R.R., ROLPH G.D. HYSPLIT (Hybrid Single Particle Lagrangian Integrated Trajectory) Model access via NOAA ARL READY Website (<http://www.arl.noaa.gov/ready/hysplit4.html>). NOAA Air Resources Laboratory, Silver Springs, MD., **2003**.
17. TROILER M., WHITE J.W.C., TANS P.P., MASARIE K.A., GEMERY P.A. Monitoring the isotopic composition of atmospheric CO_2 : Measurements from the NOAA Global Air Sampling Network, *J. Geophys. Res.* **101**, (D20), 25, 897, **1996**.
18. SZARAN J. The $\delta^{13}\text{C}$ and CO_2 concentration in the air, In: Course-book of Isotope Geology, (Ed. M. O. Jedrysek), Third School on Physics of Minerals, Part I – Isotopes., Wrocław University and Committee on Mineralogical Sciences, pp. 160-168, **1991**.
19. CRAIG H. The geochemistry of stable carbon isotopes. *Geochim. Cosmochim. Ac.*, **3**, 53, **1953**.
20. MOOK W.G., VAN DER HOEK S. The N_2O correction in the carbon and oxygen isotopic analysis of atmospheric CO_2 , *Isot. Geosci.* **1**, 237, **1983**.
21. EU-Commission, 1999. Council Directive 1999/30/EC of 22 April 1999 relating to limit values for sulphur dioxide, nitrogen dioxide and oxides of nitrogen, particulate matter and lead in ambient air. Official Journal of the European Communities L 163, 41-60, **1999**.
22. HARRISON R.M., DEACON A.R., JONES R. Sources and processes affecting concentrations of PM_{10} and $\text{PM}_{2.5}$ particulate matter in Birmingham (U.K.), *Atmos. Environ.* **31**, **1997**.
23. KUKKONEN, J., POHJOLA, M., SOKHI, R.S., LUHANA, L., KITWIROON, N., FRAGKOU L., RANTAMAKI M., BERGE E., ODEGAARD V., SLORDAL L.H., DENBY B., FINARDI S. Analysis and evaluation of selected local-scale PM_{10} air pollution episodes in four European cities: Helsinki, London, Milan and Oslo, *Atmos. Environ.* **39**, 2759, **2005**.
24. GRAMSCH E., CERECEDA-BALIC F., OYOLA P., VON BAER D. Examination of pollution trends in Santiago de Chile with cluster analysis of PM_{10} and ozone data, *Atmos. Environ.* **40**, 5464, **2006**.
25. KUC T. Concentration and carbon isotopic composition of atmospheric CO_2 in southern Poland, *Tellus* **43B**, 373, **1991**.
26. KUC T., ROZANSKI K., ZIMNOCH M., NECKI J.M., KORUS A. Anthropogenic emissions of CO_2 and CH_4 in an urban environment. *Appl. Energy* **75**, (2003), 193, **2002**.
27. SZARAN J., DUDZIAK A., TREMBACZOWSKI A., NIEZGODA H., HAŁAS S. Diurnal variations and vertical distribution of $\delta^{13}\text{C}$, and concentration of atmospheric and soil CO_2 in a meadow site, SE Poland., *Geol. Q.* **49**, (2), 135, **2005**.