INTRODUCTION

The emission of geogenic methane and carbon dioxide is regarded as one of the reasons of global climate changes. The results of studies run worldwide indicate that gas emissions from such sources strongly contribute to the increasing concentrations of greenhouse gases in the atmosphere, among which methane and carbon dioxide play crucial roles [Mazzini et al., 2009; Hong et al., 2009].

The emission of methane sourced by hydrocarbon generation and expulsion in petroleum basins is quoted as one of the two types of methane release to the atmosphere controlled by geological factors. Recently, it shares about 7–14% of global annual methane emission (from ~40 to 60 Tg CH\textsubscript{4}/year). In Europe, total methane emission from seepages reaches about 3 Tg/year [Etiopie, 2008]. The soil-to-atmosphere emission of carbon dioxide may result from bacterial oxidation of CH\textsubscript{4} or from migration from endogenic sources or from CO\textsubscript{2} generation under anoxic conditions, simultaneously with the methane [Waleńczak, 1987]. Other examples of geogenic sources of gas emissions can be mud volcanoes and/or geothermal systems.

EMISSION MEASUREMENTS OF GEOGENIC GREENHOUSE GASES IN THE AREA OF “PUSTY LAS” ABANDONED OILFIELD (POLISH OUTER CARPATHIANS)

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ABSTRACT

The Carpathians may play a significant role as a supplier of greenhouse gases to the atmosphere. Unfortunately, most of the discovered oil and gas deposits are recently only historical objects. An example is the Sękowa-Ropica Górna-Siary oil deposit located in the marginal part of the Magura Nappe where oil had been extracted in dug wells until the mid XX century. One of such extraction sites is the “Pusty Las” oilfield. In that area, 10 methane and carbon dioxide emission measurement sites were located, among which 4 in dried dug wells and 6 in dig wells still filled with oil and/or water. Dynamics of methane and carbon dioxide concentration changes were measured with the modified static chambers method. Gas samples were collected immediately after the installation of the chamber and again, after 5 and 10 minutes. In the case of reclaimed or dry dug wells, static chamber was installed directly at the ground surface. In wells still filled with oil and/or water the chamber was equipped with an “apron” mounted on special sticks. The dynamics of concentrations changes varied from -0.871 to 119.924 ppm\textsuperscript{-1}min\textsuperscript{-1} for methane and from -0.005 to 0.053% obj\textsuperscript{-1}min\textsuperscript{-1} for carbon dioxide. Average methane emission was 1.9 g m\textsuperscript{-2}d\textsuperscript{-1} and that of carbon dioxide was 26.95 g m\textsuperscript{-2}d\textsuperscript{-1}. The measurements revealed that an abandoned oil field supplies significant amounts of greenhouse gases to the atmosphere although the emission of methane is lower than that measured e.g. in mud volcanoes located in various parts of the world.

Keywords: greenhouse gas, methane, carbon dioxide, Outer Carpathians
Hydrocarbon seepages are the most typical evidences of hydrocarbon potential of petroleum basins. The Outer Carpathians are the area of significant petroleum potential, as revealed by both the macro- and microseepages, and gas exhalations [Kuśmierek et al., 2007; Kuśmierek and Machowski, 2008; Lipińska 2010]. The hydrocarbon macroseeps are typical of petroleum basins of complicated tectonics and are observed mostly in the elevated, axial zones of fold structures, and in dislocation zones. Hydrocarbon migration from deep accumulations to the near-surface zone is predominantly controlled by diffusion and effusion [e.g.: Jones and Drozd, 1983; Matthews, 1996]. Hence, the Carpathians may be a regional source of supply of greenhouse gases to the atmosphere.

In the early years of modern petroleum exploration, hydrocarbon seeps were the first prospection premises in many world-famous basins, including the Carpathians. Even before the World War II most of petroleum wells from which commercial inflows of oils and gas were obtained, were localized within the zones of natural hydrocarbon seeps [Link, 1952].

In the Carpathians, hydrocarbons have been commercially exploited since the mid XIXth century. The peak oil production from the Carpathian fields was obtained in 1909 (over 2 Mt) [Karnkowski, 1999]. An example of such a field where oil was worked from shallow dug wells is the Sękowa-Ropica Górna-Siary deposit, which operated until the mid XXth century. The deposit is located at the marginal zone of the Magura Unit. The name of the exploitation site is the "Pusty Las" ("Empty Forest") [Pietrzycki, 2013].

The present paper aims to estimate the quantity of methane and carbon dioxide emission to the atmosphere from old, inactive dug wells located in the area of the "Pusty Las" oil site.

**STUDY AREA**

The study area is located in the Sękowa Commune, in the Gorlice County of the Małopolska District. From geological point of view, this area belongs to the Outer Carpathians, precisely, to the marginal zone of the Magura Unit (the Siary Sub-unit) (Fig. 1).

**Geological setting**

The Magura Unit is the structurally highest and southwardmost tectonic unit of the Outer Carpathians. From the south, it borders the Pieniny Klippen Belt and from the north it is thrust over the Fore-Magura and the Silesian units [Oszczypko, 2004].

![Figure. 1. Geological sketch of research area (Żytko et al., 1989; Świerczewska, 2005; modified)](attachment:image)
The Magura Unit comprises four sub-units (from the south): Krynica, Bystrica, Raca and Siary [Birkenmajer, 1977]. The rocks observed at the surface are mostly the Paleogene sediments. In the Siary Sub-unit, the oldest member is named the Ropianka Beds (=Inoceramus Beds) followed by the Labowa Shales Formation [Ryłko, 2004], in which the two members were distinguished: the Skawce Sandstones and the Pasierbiec Sandstones [Oszczypko-Clowes, 2001, Ryłko, 2004]. Within the Siary zone, the Magura Beds directly cover the variegated shales of the Labowa Formation. Lithologically, these are sandstones and shales [Książkiewicz, 1972; Oszczypko-Clowes, 1999]. The Magura beds are overlain by the Gładyszów beds (uppermost Lower Oligocene) [Kopciowski and Garecka, 1996].

Generally, the Magura Unit is less hydrocarbon-prone unit in the whole Outer Carpathians [Karnkowski, 1999].

The Sękowa-Ropica Górna-Siary oil deposit

In the area of the Sękowa-Ropica Górna-Siary deposit, three oil exploitation sites (traditionally named "oil mines") are known: "Pusty Las", "Sękowa" and "Ropica Górna". Totally, 785 wells were completed here. Hydrocarbon exploration had continued until 1939. The oil was reservoired in sandstones of the Inoceramus Beds, at depth interval from 11 to 700 meters. Estimated total production was about 208000 metric tons from assessed area of 1.77 km$^2$ [Karnkowski, 1999].

METHODOLOGY OF FIELD AND LABORATORY WORKS

Field works

The studies were carried out in the area of Pusty Las “oil mine”. In this area some oil dug-wells have already been closed, i.e., entirely filled up with rubble, some others were still empty (or “open”) and, thus, recognizable as hollows of various size.

In the field studies, the modified static chamber method was applied [Leventhal, 1992; Dzieńiewicz et al., 2002; Korus et al., 2005; Sechman et al., 2006], in which gas is collected in predefined time intervals from the inner space of closed chamber installed at the terrain surface. For sampling of dug wells filled with oil and/or water, the chamber rested over the open space on two transversal levers. The chamber itself is a vessel of the volume of 48 dm$^3$, made of stainless steel and it is provided with a sealed opening to allow a sample of gas from the interior volume of the chamber. The area directly covered by the chamber is 28.27 dm$^2$.

For gas sampling, 10 sites were selected (Table 1) from which 4 sites were filled-up (closed) dug wells (sites Nos. 1, 2, 3 and 4) and next 6 were open dug wells (sites Nos. 5, 6, 7, 8, 9 and 10). The earlier experiments [Sechman and Dzieńiewicz, 2009] demonstrated that the highest dynamics of changes in gas concentrations occur in time intervals from 0 to 10 minutes since chamber installation. Hence, at each site, gas was sampled immediately after chamber setting, then after 5 and after 10 minutes. Overall, during the tests 30 samples of gas from chambers were collected. During the sampling, the atmospheric pressure was measured together with soil temperature at 10 cm depth. Gas samples were pumped to special glass vessels filled with brine [Dzieńiewicz and Sechman, 2002]. Totally, 30 gas samples were collected from 10 sampling sites. Additionally, atmospheric air sample was taken from the area of the "oil mine". The field works were run in the area of the Pusty Las "oil mine” in April, 2013.

In the case of closed dug wells, the rubbish and outer, 10-cm-thick soil layer were removed, and the chamber was pressed into the ground. The chamber walls contacting the ground were sealed (Fig. 3A). As the chamber could not be installed directly in the dug wells filled with oil and/or water, it was equipped with a special "apron” and placed over the well space onto the two transversal bars (Fig. 3B).
Laboratory tests

The molecular composition of gas samples were analyzed at the Laboratory of Gas Chromatography of the Department of Fossil Fuels. We used FISSONS Instruments GC 8160 and CARLO ERBA Instruments GC 6300 gas chromatographs equipped with FID and TCD detectors. In each sample, methane, ethane, propane, i-butane, n-butane, neo-pentane, i-pentane, n-pentane, ethylene, propylene, 1-butene and carbon dioxide were determined. Detection limit for FID is 0.01 ppm for hydrocarbons. Analytical precision is 2% of measured value and 10% at detection limit. TCD detection limits for carbon dioxide is 100 ppm and estimated precision is 2% of measured value and 10% at detection limit.

Table 1. Principal statistical parameters of alkanes and carbon dioxide concentrations calculated for all-30 measurements

<table>
<thead>
<tr>
<th>Stat. parm. (ppm)</th>
<th>CH₄</th>
<th>C₂H₆</th>
<th>C₃H₈</th>
<th>i-C₄H₁₀</th>
<th>n-C₄H₁₀</th>
<th>neo-C₅H₁₂</th>
<th>i-C₅H₁₂</th>
<th>n-C₅H₁₂</th>
<th>CO₂ **</th>
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<tbody>
<tr>
<td>Min</td>
<td>0.8</td>
<td>b.d.l.</td>
<td>b.d.l.</td>
<td>b.d.l.</td>
<td>b.d.l.</td>
<td>b.d.l.</td>
<td>b.d.l.</td>
<td>b.d.l.</td>
<td>0.07**</td>
</tr>
<tr>
<td>Max</td>
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<td>3.62</td>
<td>3.50</td>
<td>0.77</td>
<td>0.65</td>
<td>0.01</td>
<td>0.41</td>
<td>0.05</td>
<td>0.22**</td>
</tr>
<tr>
<td>Mean</td>
<td>65.4</td>
<td>0.25</td>
<td>0.16</td>
<td>0.04</td>
<td>0.03</td>
<td>0.000</td>
<td>0.02</td>
<td>b.d.l.</td>
<td>0.12**</td>
</tr>
<tr>
<td>Stand. dev.</td>
<td>221.0</td>
<td>0.69</td>
<td>0.62</td>
<td>0.14</td>
<td>0.16</td>
<td>0.08</td>
<td>0.01</td>
<td>0.04</td>
<td>0.04**</td>
</tr>
<tr>
<td>Median</td>
<td>3.1</td>
<td>b.d.l.</td>
<td>b.d.l.</td>
<td>b.d.l.</td>
<td>b.d.l.</td>
<td>b.d.l.</td>
<td>b.d.l.</td>
<td>b.d.l.</td>
<td>0.12**</td>
</tr>
<tr>
<td>Percentage of samples*</td>
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<td>38.7</td>
<td>19.4</td>
<td>19.4</td>
<td>16.1</td>
<td>3.2</td>
<td>16.1</td>
<td>16.1</td>
<td>100</td>
</tr>
</tbody>
</table>

* – percentage of samples with concentration of given component over detection limit; ** – minimum, maximum, mean, median and standard deviation in vol.%, b.d.l. – below detection limit (detection limit for hydrocarbons is 0.01 ppm and 100 ppm for carbon dioxide).

Figure 3. Measurements within the filled-up dug wells (A) and open dug wells (B) (Fot. D. Pietrzycki)

METHODOLOGY OF STATISTICAL ANALYSIS AND RESULTS PRESENTATION

The populations of measured concentrations of gaseous hydrocarbons were initially characterized by determination of basic statistical parameters, calculated for all the obtained results and separately, for samples collected over the closed and the open dug wells. The relationships between measured methane and carbon dioxide concentrations were evaluated using the correlation plots based upon the cartesian coordinates XY system. Such plots were constructed for the whole results and separately, for the results for closed and open dug wells. In order to evaluate the dynamics of changes in concentrations of methane and carbon dioxide, linear plots were drawn, which illustrated...
the changes of gas concentrations in time. Then, computer-generated, linear trend of concentration changes was overprinted on the plots. For the trend lines, the coefficient of determination $R^2$ was calculated. For both the methane and carbon dioxide, results were noted in ppm and in volume percents per minute. The emission was expressed in milligrams per square meter per day for methane emission and in grams per square meter per day for carbon dioxide emission.

**METHODOLOGY OF EMISSION CALCULATION**

The quantities of methane and carbon dioxide emissions were calculated using the formula established during the previous geochemical surveys [Korus et al., 2002; Dzieniewicz et al., 2006; Sechman and Dzieniewicz, 2009]. If the collection chamber was installed at the ground surface the area covered by the chamber was constant. If the collector was mounted over the open dug wells the chamber parameter depended on the changing surface covered by the "apron".

**RESULTS AND DISCUSSION**

Statistical characterization of population of all measured gas concentrations.

In the analyzed gas samples, saturated hydrocarbons $C_1$-$C_5$ and carbon dioxide were detected. Methane concentrations varied from 0.81 to 1220 ppm. Maximum ethane concentration was 3.62 ppm and this gas was found in almost 40% of the analyzed samples. Propane occurred in 19.4% of the analyzed samples and its values reached up to 3.5 ppm with the mean value 0.16 ppm. The same percentage of samples contained i-butane whereas i-pentane and n-pentane were detected in over 16% of the analyzed samples. Maximum butane and pentanes concentrations did not exceed 1 ppm. Trace amounts of ethylene were measured in only two samples whereas the remaining gaseous alkenes were absent in the analyzed material. The concentrations of carbon dioxide varied from 0.07 to 0.22 vol.% with the mean value 0.12 vol.% (Table 1).

Concentrations of methane and carbon dioxide measured in the atmospheric air from the area of the Pusty Las site were 3.68 ppm and 0.09 vol.%, respectively. For comparison, the average concentration of atmospheric methane is about 2 ppm and that of carbon dioxide is about 0.03 vol.%. Moreover, concentrations of methane and carbon dioxide measured over the closed bituminous coal mines in the Wałbrzych mining district (SW Poland) were 5 ppm and 0.05 vol.% respectively [Korus et al. 2002], and methane concentration measured in the atmospheric air sampled immediately over the mud volcano from Azerbaijan was 99.65 vol.% [Mazzini et al., 2009].

Statistical characterization of gas concentrations in samples collected from over the closed oil dug wells

In gas samples collected from the closed oil dug wells, methane, ethane, propane, i-butane, i-pentane, n-pentane and carbon dioxide were detected. Methane was found in all of the analyzed samples, in concentrations from 0.8 to 1220 ppm with arithmetic mean 125.4 ppm and median value 3 ppm. Ethane was detected in almost 42% of analyzed samples in concentrations up to 3.62 ppm (arithmetic mean – 0.4 ppm). Propane was observed in almost 17% of the analyzed samples. Its concentrations reached up to 3.5 ppm with arithmetic mean 0.32 ppm. Maximum concentrations of i-butane and n-butane were 0.77 and 0.65 ppm, respectively, and were observed in almost 17% of the analyzed samples. Neo-pentane has not been detected whereas i-pentane and n-pentane were observed in about 17% of the analyzed samples, but in amounts below 0.5 ppm. The concentrations of carbon dioxide varied from 0.07 to 0.22 vol.% with the arithmetic mean 0.14 vol.% (Table 2).

Statistical characterization of gas concentrations in samples collected from over the open oil dug wells

In gas samples collected from the open oil dug wells, methane, ethane, propane, i-butane, neo-pentane, i-pentane, n-pentane and carbon dioxide were detected. Methane was observed in all of the analyzed samples, in concentrations from 1.41 to 288.5 ppm with arithmetic mean 28.7 ppm and median value 2.9 ppm. Ethane was found in almost 40% of the analyzed samples, in concentrations up to 1.23 ppm (arithmetic mean – 0.06 ppm). Propane was detected in over 22% of analyzed samples, in concentrations up to 0.44 ppm (arithmetic mean – 0.06 ppm). Neo-pentane was detected in a single sample (0.008 ppm).Maxi-
mum concentrations of i-butane and n-butane were 0.12 and 0.06 ppm respectively. Both i-pentane and n-pentane were found in about 16% of analyzed samples, in concentrations below 0.2 ppm. The concentrations of carbon dioxide varied from 0.07 to 0.15 vol.% with the arithmetic mean 0.11 vol.% (Table 3).

Evaluation of geochemical indicators

Generally, concentrations of alkanes in analyzed samples decreased with the increasing number of atoms in molecules, which suggests deep origin of these hydrocarbons (Table 1). Concentrations of methane and carbon dioxide were higher in samples taken from above the closed dug wells (Table 2, 3). The lower concentrations of both gases detected over the open dug wells can be related also to the fact that most of the wells were filled with water, which provides a barrier for both gases. Additionally, carbon dioxide, which is heavier than the air, may accumulate in local depressions. The concentrations of methane and carbon dioxide did not reveal correlation (Fig. 4A). However, the coefficient of determination $R^2$ calculated for samples from over the closed dug wells was 0.48 whereas that determined for open wells was only 0.03 (Fig. 4B, C). These values suggest that relationships between methane and carbon dioxide concentrations from these sites are much interrelated. It can be explained as an effect of microbial generation of these gases under anoxic conditions. Such a process may operate when the dispersed organic matter is contained in a rubble filling the dug wells. Hence, the newly generated microbial gases contribute to the flux of gases ascending from deep sources. In the open dug wells, carbon dioxide, which is heavier than the atmospheric air, presumably accumulates at the bottoms of the wells whereas methane, which is lighter than the air, readily raised towards the sampling chamber. Moreover, deep-sourced carbon dioxide more intensively dissolves in the water filling the wells that deep-sourced methane, which affects the relationships between both gases.

Table 2. Principal statistical parameters of alkanes and carbon dioxide concentrations calculated for closed dug wells (12 results)

<table>
<thead>
<tr>
<th>Stat. parm. (ppm)</th>
<th>Alkanes</th>
<th>CO$_2$**</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>CH$_4$</td>
<td>C$_2$H$_6$</td>
</tr>
<tr>
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<td>b.d.l.</td>
</tr>
<tr>
<td>Max</td>
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<td>3.62</td>
</tr>
<tr>
<td>Mean</td>
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<td>1.04</td>
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<tr>
<td>Median</td>
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<td>b.d.l.</td>
</tr>
<tr>
<td>Percentage of samples*</td>
<td>100</td>
<td>41.7</td>
</tr>
</tbody>
</table>

* – percentage of samples with concentration of given component over detection limit; ** – minimum, maximum, mean, median and standard deviation in vol.%, b.d.l. – below detection limit (detection limit for hydrocarbons is 0.01 ppm and 100 ppm for carbon dioxide).

Table 3. Principal statistical parameters of alkanes and carbon dioxide concentrations calculated for open dug wells (18 results)

<table>
<thead>
<tr>
<th>Stat. parm. (ppm)</th>
<th>Alkanes</th>
<th>CO$_2$**</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>CH$_4$</td>
<td>C$_2$H$_6$</td>
</tr>
<tr>
<td>Min</td>
<td>1.1</td>
<td>b.d.l.</td>
</tr>
<tr>
<td>Max</td>
<td>288.5</td>
<td>1.23</td>
</tr>
<tr>
<td>Mean</td>
<td>28.7</td>
<td>0.16</td>
</tr>
<tr>
<td>Stand. dev.</td>
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<tr>
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<td>2.9</td>
<td>b.d.l.</td>
</tr>
<tr>
<td>Percentage of samples*</td>
<td>100</td>
<td>38.9</td>
</tr>
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</table>

* – percentage of samples with concentration of given component over detection limit; ** – minimum, maximum, mean, median and standard deviation in vol.%, b.d.l. – below detection limit (detection limit for hydrocarbons is 0.01 ppm and 100 ppm for carbon dioxide).
Dynamics of changes in concentrations of methane and carbon dioxide, and quantities of their emission

The dynamics of concentrations changes of methane released to the atmosphere from oil dug wells is expressed by the range of migration rates from -0.87 to 119 ppm min\(^{-1}\). In 60% of sampling sites, the increasing trends were observed, in the remaining 40% the decreasing trend was found. Dynamics of carbon dioxide concentration changes is illustrated by values from -0.005 to 0.053 vol.% min\(^{-1}\). The trends of CO\(_2\) concentration changes were identical as those for methane (Fig. 4).

The emission values of methane varied from -7.8 to 18660 mg m\(^{-2}\) d\(^{-1}\) for closed dug wells and from -62.2 to 42.4 mg m\(^{-2}\) d\(^{-1}\) for open wells, negative emissions are the result of bacterial destruction of methane [e.g.: Klusman, 1993]. Carbon dioxide showed emission values from about -10 to about 47 g m\(^{-2}\) d\(^{-1}\) for closed wells and from -9.9 to about 226 g m\(^{-2}\) d\(^{-1}\) for open wells (Table 4).

Maximum emission observed at No. 2 sampling site was several orders of magnitude lower than that measured in the areas of mud volcanoes from various locations in the world [e.g. Dimitrov, 2002; Etiope et al., 2002].

Considering all of the analyzed samples, the average emission of methane expressed by median value was 4.75 mg m\(^{-2}\) d\(^{-1}\) and by mean value was 1876 mg m\(^{-2}\) d\(^{-1}\). Median value is one order of magnitude higher than the values obtained for various oil/gas fields in the Polish Outer Carpathians [Sechman and Dzieniewicz, 2009]. It must be emphasized that in 2009 research the measurements were run outside the hydrocarbon exploitation areas.

CONCLUSIONS

The sampling sites of methane and carbon dioxide were located in the area of historical, abandoned Pusty Las "oil mine". Totally, 10 oil dug wells were selected for sampling, from which 4 were closed (i.e., completely filled with rubble) and 6 were still recognizable (i.e. hollows of various size). Dynamics of concentration changes of gaseous components: alkanes and carbon dioxide was determined using the static chambers methodology. The results revealed that:

- gaseous hydrocarbons detected in samples collected with the static chambers originate from deep-seated sources,
- gaseous alkanes are dominated by methane; its concentrations over the closed dug wells varied from 0.81 to 1220 ppm and those from over the open dug wells changed from 1.1 to 288.5 ppm,
- values of methane emission from the closed wells varied from -7.8 to 18660 mg m\(^{-2}\) d\(^{-1}\) and that from the open wells were from -62.2 to 42.4 mg m\(^{-2}\) d\(^{-1}\),
- concentrations of carbon dioxide varied from 0.07 to 0.22 vol.% over the closed wells and from 0.07 to 0.15 vol.% over the open wells,
- values of carbon dioxide emission from the closed wells changed from about -10 to about 47 g m\(^{-2}\) d\(^{-1}\) and then from the open wells varied from -9.9 to 225.8 g m\(^{-2}\) d\(^{-1}\),
- average methane emission was 1.9 g m\(^{-2}\) d\(^{-1}\) and average carbon dioxide emission was almost 27 g m\(^{-2}\) d\(^{-1}\),
- higher emission values of methane and carbon dioxide from the closed oil dug wells may result from microbial generation of these gases within the rubble which fills the wells; this component contributes to the flux of gases from deep sources,
Figure 5. Dynamics of changes of concentrations measured in sample stations Nos. 1–10 (A-J)
the measurements were taken within a short period of time (one week), so the seasonal changes could not have had any influence on the established emission value.

In the area of the Pusty Las "oil mine", significant methane emission occurs from abandoned oil dug wells, although average emission values were only 1 order of magnitude higher than those measured in the petroleum-prone areas in the Carpathians. Moreover, the maximum methane emission measured over one of sampled dug wells in the Pusty Las area was several orders of magnitude lower than the emissions from mud volcanoes from various localities in the world.

The results demonstrate that old oil dug wells still present in the areas of historical hydrocarbon exploitation in the Carpathians may be a significant source of greenhouse gases released to the Earth atmosphere.

**REFERENCES**


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<td></td>
<td>$n_{CH_4}$</td>
<td>$E_{CH_4}$</td>
</tr>
<tr>
<td></td>
<td>(ppm min$^{-1}$)</td>
<td>(mg m$^{-2}$ d$^{-1}$)</td>
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<tr>
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</tr>
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