Effect of the Wildfires on Sandy Podzol Soils of Nadym Region, Yamalo-Nenets Autonomous District, Russia

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Active land disturbance of soil and environments appears even in remote landscapes of tundra and forest tundra. Wildfires become a frequent factor of soil degradation and intensification of permafrost degradation, also affecting the global balance of carbon, especially content and distribution of polycyclic aromatic compounds. Mature unaffected by fires sandy-textured podzols soils were investigated in comparison with the same soil strongly affected by surface fire in the end of August 2016 in surroundings of the Pangody settlement, Nadym district, Yamal region. Data obtained showed an intensive morphological transformation of the topsoil layers, decreasing total organic matter and apparently increasing the humus enrichment by nitrogen. Wildfires also result in complete transformation of the fractional composition of the polycyclic aromatic compounds. The sum of PAHs increases intensively as well as benzo(a)pyrene content in soils. Therefore, soils of the cryolithozone become more faced to the wildfires during the last decades. Even one-time fire effect results in serious transformation of soil geochemical state.

1. Introduction

Anthropogenic successions caused by climate dynamics and effects of natural (windfalls and pests irruption), natural and anthropogenic (wildfires), and anthropogenic (logging) phenomena are characteristic of tundra and forest ecosystems. The human impact on natural biogeocenoses, which becomes more actual nowadays, leads to a change in ecosystems functioning and makes monitoring work extremely important. An important factor, disturbing natural processes in ecosystems, is wildfires, whose consequences are difficult to predict. Wildfires are given special attention in natural ecosystem surveys. A considerable body of work existing in the science literature is about the forest fires role in the forest cover natural dynamics as they are the most important environmental factor among others that determine the structure and dynamics of forests and, accordingly, the ecological territory state [1–5]. A lot of attention is paid to the problem of postpyrogenic soil change, its properties, soil regime, and functions in a boreal zone. A variety of research studies are devoted to pyrogenic transformation of the water regime, vegetation, and forest landscapes in whole [6–15]. However, data on similar studies of the pyrogenic transformation in polar, tundra, and subboreal landscapes of cryolithozone are extremely rare in the literature. The main directions in this field are research studies on postpyrogenic transformation of plant cover. Therefore, the purpose of this work is to study soil changes after wildfires affect the cryolithozone of tundra and forest-tundra ecosystems.

Besides, scientific interest in PAHs has continued unabated over the last decades since the temperature effects on soil organic matter (SOM) are considered the most common way of the PAH formation.

PAHs’ anthropogenic sources of pyrogenic origin (vehicles, industrial facilities, energy facilities, and others emissions) have been studied in detail so far. At the same
time, there is a trend to reduce the PAH pollution of various landscape components in developed countries due to improved technologies and environmental standards [16, 17]. Therefore, increasingly greater attention is paid to natural pyrogenic sources of PAHs such as volcanism and wildfires. Volcanic sources of PAHs are very local, but forest, steppe and peat fires damage significant areas annually, which is considered as one of the global factors of PAH environmental release. Besides, the world literature contains numerous laboratory experiment data on the possibility of PAH formation from vegetation components; moreover, burning conditions (temperature and oxygen access) and a vegetation type significantly influence on the quantity and structure of the compounds formed.

A more detailed review on this issue was given by Tsibart and Gennadiev [18]. At the same time, the trend of the pyrogenic PAH accumulation in landscapes, especially in soils as a deposit environment, has not been practically studied. Individual publications are devoted to investigation of PAHs in fire-affected soils of different genesis [19–25]. But still there are no enough data in the literature on the composition, quantity, and distribution of PAHs in soils coming from wildfires, particularly in the north region of cryolithozone. North polar ecosystems are characterized by low resistance to anthropogenic impact against the background of high resource potential. There are some works that aimed at peculiarities of PAH accumulation in tundra ecosystems under the different influences, for example, Yakovleva et al. [26], but fire effect was not investigated yet.

The main objective of this work is to investigate PAHs in the initial stage of the demutation postfire sequence of postpyrogenic soils.

2. Materials and Methods

2.1. Study Site and Soil Sampling. The research object is located near Pangody settlement, Nadym district, Yamal region, Western Siberia, Russia (see Figures 1 and 2) (N 65-50-356, E 074-24-210). The study plots are represented by forest-tundra ecosystem, sublayered by stratified parent material, namely, sand sublayered by silty textured permafrost grounds (on the depth of 1.7 m, morphologically measured). The objects of investigation for comparison fire effect are two soils pits (2 soil profiles) presented by mature soil, podzol (according to WRB podzols), and fire-affected soil, which was initially the same with mature podzol soils. The control plot contained plant cover and soil unaffected by wildfire and was situated close to the fire-affected plot. The surface fire event occurred in 2016, at the end of the summer. Soil samples were taken in 2016 too, immediately after the fire. Soil samples were selected from each horizon in three replications with an aim to analyze basic chemical properties: (1) podzol, mature soil, unaffected by wildfire; stratification of layers is folic-albic-spodic-podzol (according to WRB) and O-E-BF-BC-C (according to Russian soil classification); (2) podzol, affected by fire, stratification of layers is folic$_{pyr}$-albic$_{pyr}$-spodic-podzol (according to WRB) and Opyr-Epyr-BF-BC-C (according to Russian soil classification). So, the fire effect on soil macromorphology is
pronounced only in the topsoil or superficial soil horizons. Two soil pits were investigated (at unburned site as a control and at surface fire plot); totally, more than 30 individual soil samples were sampled for further chemical analyses in laboratory and were analyzed.

2.2. Chemical Analyses of the Fine Earth. Air-dried soil samples were grounded and passed through a 2 mm sieve for further analysis of fine earth. Analyses were conducted in the certified laboratory of St. Petersburg State University at the Department of Applied Ecology, Russia. Soils were analyzed according to the following methods: determination of actual acidity (pHw) and potential soil acidity (CaCl₂). Soil microbial respiration was determined using incubation chambers. An element analyzer Euro EA3028-HT Analyzer was used for determination of C and N content in SOM, a pH meter pH-150M for assessing pH in water and in salt suspensions.

Concentrations of following 14 polynuclear aromatic hydrocarbons (PAHs) and its total sum were studied:

- Acenaphthene (ANA)
- Anthracene (ANT)
- Benzo(a)anthracene (BaA)
- Benzo(a)pyrene (BaP)
- Benzo(b)fluoranthene (BbF)
- Benzo(g,h,i)perylene (BPE)
- Benzo(k)fluoranthene (BkF)
- Chrysene (CHR)
- Dibenz(ah)anthracene (DBA)
- Fluoranthene (FLT)
- Fluorene (FLU)
- Naphthalene (NAP)
- Phenanthrene (PHE)
- Pyrene (PYR).

National standard method PND F 16.1:2:2.2:3.62–09 [27], US EPA method 8310 [28, 29], and publications of D.N. Gabov [30, 31] were used for PAH content analysis. The US EPA method 3550b [29] was used for PAH extraction. It consists of ultrasonic treatment via Branson 5510 ultrasonic bath (USA, power 469 W, working frequency 42 kHz) at room temperature with methylene chloride (high purity grade).

Kuderna Danish concentrator (Supelco) was used for solvent removal (evaporation). The US EPA purification method 3660c (1996) for refining PAHs fractions is focused on consecutive chromatography in columns filled with aluminum oxide (Brockman activity grade 2–3, Neva Reaktiv) and silica gel (Fluka). The absence of peaks in the blank chromatogram meant the purity of PAH fractions. Standard PAH solutions were prepared using a standard mixture of 14 PAHs (Supelco) (concentrations of each component in the range of 100–2000 μg/cm³). A reverse-phase HPLC (high-performance liquid chromatography) was used for analysis of components and quantities of PAHs in soils. This analysis was made in gradient mode with spectrofluorimetric detection via chromatograph “Lymahrom" (“Lumex", Russia) at 30°C on a column Supelcosil™ LC-PAH n5 μm (25 cm × 2.1 mm). Mobile phase was provided with acetonitrile-water gradient. A special valve was used for injection of 10 μl volume samples. Individual PAHs were identified by the time of retention. And then components’ fluorescence spectra from the column were compared with standard PAH spectra. PAH quantity was determined using the external standard method. Moreover, the same procedure was performed for standard reference materials (1944 New York/New Jersey Waterway Sediment (National Institute of Standards and Technologies NIST, USA)) that contain a mixture of 14 PAHs in order to assure the quality. An error of PAH determination (in particular benzo(a)pyrene) in the soils was 35% in the range...
of 5–40 ng/g and 25% in the range of 40–2000 ng/g with a confidence probability of $P = 0.95$. There are standards for the benzo(a)pyrene content in the soils (20 ng/g) in Russian legislation (GN 2.1.7.2041–06). The results obtained were compared with these standards. In order to compare the PAH content with standards for benzo(a)pyrene in soils, benzo(a)pyrene total potency equivalents were calculated by multiplying the concentration of each analyzed individual PAH in the sample by its benzo(a)pyrene toxic equivalency factor according to the World Health Organization (WHO/IPCS 1998) scheme. PAH sources apportionment was performed using special molecular markers and ratios, proposed by Yunker et al. [32], and a total combustion PAH index, reported by Hwang and Cutright [33]. Applied markers, such as ANT/(ANT + PHE), FLT/(FLT + PYR), BaA/(BaA + CHR), IPY/(IPY + BPE), CombPAH/14PAH, and BaP/BPE, are applied in order to distinguish between pyrogenic and petrogenic sources of PAHs. The sum of combustion PAHs (combPAH/14PAH) was used as the tracer of pyrogenic sources. The combPAH/14PAH marker indicates the proportion of the sum of combustion-specific compounds in total PAH content, which are fluorene, pyrene, chrysene, benzo(a)anthracene, benzo(k)fluoranthene, benzo(b)fluoranthene, benzo(a)pyrene, benzo(g,h,i)perylene, and indeno(1,2,3-cd)pyrene [34].

All the measurements were performed in triplicate; average values were given with standard error. In order to determine the differences between studied objects, experimental data were statistically treated using Sigma-Plot 8.0 software (calculation of average values, standard deviation, one-way ANOVA, and post hoc test).

3. Results and Discussion

In the mature soil, the uppermost horizon was presented by natural forest floor with the thickness of about 5–7 cm, while the thickness of the postfire remnants of the fire-affected soil was only 1-2 cm and spatially was expended sporadically, namely, in discontinuous form. A new shallow pyrogenic horizon, whose properties are determined by the presence of abundant coals, is formed [35]. Folic$_{pyr}$ and Albic$_{pyr}$ (Opyr and Epyr, respectively) horizons are black layers due to the deposition of an ash and charcoal at the soil surface. Organic matter and plant residues are lost from the surface horizons of the soil. It is related to the destruction of the organic horizons, the mineralisation of root residues, and the almost complete absence of fresh plant waste, which could be a material for humification. Moreover, morphological characteristics are changed in case of forest fires [36]. A decrease in the porous media was recognised as the main soil development process after the fires as a result of soil porous infilling by ash and charred organic material of different natures. Fire-affected soil has a higher pH of 5.2 in comparison with increased acidity of 4.7 pH units in a mature soil unaffected by fire.

The total organic carbon content was quite different in soils affected by fire, compared with benchmark ones (see Figure 3(a)). This fact could be interpreted as a result of brief combustion and following losses of soil organic matter under the spontaneous high-temperature fire effect.

The content of nitrogen (see Figure 3(b)) was decreased in soil of postfire plot, but not in proportional ratio with the TOC content. This resulted in the values of C/N ratios (see Figure 4) used for estimation of the humus enrichment degree by nitrogen. This index showed that the soil organic matter of fire-affected soil is more enriched by nitrogen organic substances of the mature soil. This was not the result of real changes in organic matter quality due to humification or stabilization. It could be result of apparent enrichment of the humic substance systems by additional accumulation of mineral form of the nitrogen compounds under effect of fire-induced mineralisation and transformation of the SOM. Data obtained are in well correspondence with other pyrogenic soils of Siberia [7, 8, 11, 15, 37]. A clear maximum of PAH concentrations in the surface horizons (due to enrichment with organic matter, which is explained by the high absorbing capacity of organic matter in comparison with
mineral matter and the main amount of polyarenes from the atmosphere) was not revealed in studied soils as, for example, in podzols near Baikal lake [38].

The PCM (potential mineralizable carbon) values of the mature soil were essentially higher in superficial layers of mature soil than in lower ones. This was also an indicator of the fact that the deepness of fire effect in case of one-time surface fire is not high, and this effect is not pronounced in the deepest soil layers (see Figure 5). Data obtained are comparable with those obtained previously [16, 17, 39, 40].

The total content of PAHs (see Figure 6) in fire-affected soil was higher than in control, and this is caused by pyrogenic factor. The total concentration of the PAH in control soil was comparable with those founded for mature soils of Belyi islands [41] and for soils of benchmark merged areas of the Russian Arctic [42]. However, data obtained for podzol are less significant than corresponded data shown by other authors [43, 44] and much more than for soils in forest steppe [45]. Pyrogenic processes lead not only to formation of increased total content of the PAHs but also accumulation of benzo(a)pyrene in soils (see Figure 7) and formation of completely different component compositions of the PAHs (see Figures 8 and 9), where the pyrogenic components became dominant.

The calculation of the indexes of the PAHs origin showed that soils under the fires showed accumulation of the petrogenic compounds \( \text{FLT}/(\text{FLT} + \text{PYR}) = 0.42, \ \text{IPY}/(\text{IPY} + \text{BPE}) = 0.46, \ \text{CombPAH}/\sum \text{PAH} = 0.79, \ \text{and BaP}/\text{BP} = 1.19 \). These indexes were essentially lower in natural soils, which indicate the petrogenic sources of the main PAHs in this soil. This well corresponds with data on PAH content in soils of the Belyi island with different functional loads, collected previously [40]. However, Dymov et al. [43] showed that content of PAHs such as CHR, FLU,
Table 1: Statistical data analysis.

| PAH | PHE | ANT | FLT | PYR | BaA | CHR | BbF | BkF | BaP | DBA | BPE | IPY | \( \Sigma \text{PAHs} \) |
|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----------------|
| P value | ANOVA (a) | <0.05 | <0.05 | <0.05 | 0.22 | 0.73 | <0.05 | 0.5 | <0.05 | <0.05 | 0.52 | <0.05 | <0.05 | <0.05 |
| Significance of differences | Significant | Sign | Sign | Insignificant | Insignificant | Sign | Insignificant | Sign | Sign | Insignificant | Sign | Sign | Sign |

NAP, PYR, and ANT significantly increased in OPIR (L) horizon after wildfires. Only some evidence of the transport pollution has been fixed in natural soils (BaA/ (BaA + CHR) = 0.24 and IPY/(IPY + BPE) = 0.43). So, the fire affects not only total concentration of the PAH but also their component composition.

One-way ANOVA analysis revealed significant differences in the content of most individual PAHs (phenanthrene, anthracene, fluoranthene, chrysene, benzo(k) fluoranthene, benzo(a)pyrene, benzoperylene, and idenopyrene), as well as the total PAH content between the control soil and postfire soil (P < 0.05) (Table 1).

4. Conclusions

Soils of the cryolithic zone have faced more wildfires during the last decades. Wildfires become more and more frequent, especially in tundra and forest-tundra in the Western Siberia [46, 47] (http://wiki-fire.org). The podzols soils have been investigated in two plots: not affected by fire and under the influence of the surface fire in August 2016 in surroundings of the Pangody settlement, Nadym district, Yamal region. Data obtained showed an intensive morphological transformation of the topsoil layers decreasing total organic matter and apparently increasing the humus enrichment by nitrogen. Wildfires also result in complete transformation of the fractional composition of the polycyclic aromatic compounds. The sum of PAHs increases intensively as well as benzo(a)pyrene content in soils. The structural composition of the PAHs showed changes under the fire effect: the components of the pyrogenic origin were prevailing in fire-affected soils. Thus, even one-time fire effect results in serious transformation of soil geochemical state.

Data Availability

The data used to support the findings of this study are available from the corresponding author upon request.

Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this paper.

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