

1 **Mobilization of geochemical elements in the active layer of**
2 **permafrost to surface water in Russian Arctic**

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Abstract

The predicted increasing ground temperatures in the Arctic results in the deepening of the active layer and intensification of geochemical processes, which could affect the geochemical composition of surface waters. Determining the responses of the riparian soil systems to surrounding hydrological flows under changing climate conditions is important for understanding the seasonal changes in hydrological processes. Therefore, in this study, one soil core from the study area polygon rim (close to the Taz River, TA) and two soil cores from the riverain terrace (close to the Syoyakha River, SY and Murtyyakha River, MU) in Russian Western Siberia and their supra-permafrost water, adjacent stream flows and river water were sampled for analysis of geochemical elements. The results showed that most elements above their respective detection limits (Mn, Sr, Fe, Mg, Cr, Co, V, Pb, Al and Ca) started to accumulate in the downwards gleyed layer during September–October in response to the deepest thaw in the active layer. This study focused on the highly mobile elements, i.e., Mn, Ca, Mg, Al and Ti, in the deepest layer; and found that the transport of organic matter in the upper layer carried these elements to both surface water ponds/flows and supra-permafrost water, and further, to the rivers. The best linear correlation for both stream flows and river water were Mn, which may be a proxy for predicting the processes occurring within the active layer during the annual summer-autumn thaw. Finally, landscapes with different ice contents may experience changes in the elements transported to surface waters.

Keywords: Geochemical elements, Arctic, Permafrost-affected soils, active layer, surface water, seasonal thawing

39 **Plain Language Summary**

40 The rapid increasing atmospheric temperature in the Arctic has led the feedback of
41 permafrost thawing, which increases the depth of active layer and accelerate chemical
42 weathering. These processes may cause chemical substances further releasing to adjacent
43 aquatic landscapes. Here we investigated the horizontal and vertical transport of geochemical
44 elements through surface flow and supra-permafrost water to river water from three cryogenic
45 soil cores in the Arctic tundra. The results showed that most elements would accumulate in
46 the gleyed layer that was the deepest in the active layer during thawing season. Mn, Ca, Mg,
47 Al and Ti showed the highest mobility from soil to supra-permafrost water to the surface
48 flows/rivers, while Mn was the only element could transport from topsoil and
49 supra-permafrost water to surface water and river, which may be a proxy predicting the
50 processes of active layer during summer-autumn thawing.

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1. Introduction

The increasing atmospheric temperatures in the Arctic have led to permafrost thawing and vertical downward migration of the active layer into formerly frozen ground, during the thawing season [H H Christiansen *et al.*, 2010; R Connon *et al.*, 2018; M T Jorgenson *et al.*, 2010; V E Romanovsky *et al.*, 2010; S L Smith *et al.*, 2010]. The active layer responds to the changing climatic conditions by increasing its depth, thus accelerating chemical weathering [N Colombo *et al.*, 2019; K Keller *et al.*, 2007; 2010] and potentially releasing geochemical components and organic carbon from the soil to adjacent aquatic landscapes [A J Barker *et al.*, 2014; G Grosse *et al.*, 2016; K D Johnson *et al.*, 2013; C E Johnston *et al.*, 2014; S V Loiko *et al.*, 2017; T V Raudina *et al.*, 2018; E A G Schuur *et al.*, 2015]. This would increase the movement of carbon dioxide (CO₂) from the soil to aquatic reservoirs such as rivers and lakes [C D Elder *et al.*, 2019; S M Natali *et al.*, 2015; Y M Polishchuk *et al.*, 2018; B Wild *et al.*, 2019]; this occurs through the transport of solute and water along surface streams and the permafrost table, which is as known as “supra-permafrost flow”. Supra-permafrost water is found in the shallow subsurface of the active layer, where it is typically situated at the boundary between the frozen and thawed fractions of the soil profile [D J Cederstrom *et al.*, 1953]. Owing to the absence of groundwater discharge in the permafrost-affected region, solutes being transported to lakes or rivers from adjacent soils originate primarily from supra-permafrost water.

Permafrost degradation has been associated with an increased contribution of groundwater to stream via surface flows or supra-permafrost water in Canadian Arctic rivers [M A Walvoord and R G Striegl, 2007]. Ice wedge degradation has also been proven to be related to the water balance of lowland across the Arctic landscapes, and to increase streamflow [A K Liljedahl *et al.*, 2016]. Several studies have observed substantial increments in discharges from the major Eurasian Arctic rivers due to the warming climate [N J Brown *et al.*, 2019; D Feng *et al.*, 2019; R B Lammers *et al.*, 2001; D Q Yang *et al.*, 2002; L Zheng *et al.*, 2019]. T V Raudina *et al.* [2018] observed a northward shift of the permafrost boundary under climate change scenarios, and that the concentrations of dissolved organic carbon (DOC), major and trace elements, and greenhouse gases are expected to decrease in the supra-permafrost waters at the border between the thawed

and frozen parts from peat soils in the western Siberia lowland. *S V Loiko et al.* [2017] hypothesized that the direct mobilization of soil waters to hydrological networks and the transformation of autochthonous processes were controlled by physical factors in different landscapes. Overall, there appears to be considerable variability and uncertainty regarding how discharges of DOC, elements, and greenhouse gases in supra-permafrost waters/surface flows respond to permafrost thaw and are released to surface water ponds and rivers.

Several studies concerning the biogeochemical cycles of carbon and related metals focused on both the aquatic and the terrestrial parts of the continental permafrost-bearing ecosystem [*L Guo et al.*, 2007; *J W McClelland et al.*, 2007; *D Olefeldt and N T Roulet*, 2012; *O S Pokrovsky et al.*, 2011]. More recently, the biogeochemistry of soil porewater and supra-permafrost water in mineral and organic/peat parts of soil profiles in the permafrost areas have been studied [*A J Barker et al.*, 2014; *S Jessen et al.*, 2014; *D Lamhonwah et al.*, 2017; *S V Loiko et al.*, 2017; *T V Raudina et al.*, 2018; *L E Street et al.*, 2016]. The previous studies concluded that the export fluxes of DOC, greenhouse gases, and trace metals from the active layer of permafrost to the surrounding hydrological landscapes are determined by the amount of water passing through the active layer till the border of the frozen permafrost, before being drained to a lake or river [*Oleg S. Pokrovsky et al.*, 2016; *O. S. Pokrovsky et al.*, 2016; *T V Raudina et al.*, 2018]. *A J Barker et al.* [2014] reported that the concentrations of metals in surface water were related to the increasing active layer thickness/degrading permafrost during late fall and early winter. These results revealed that the dynamics of trace metal concentrations in the transitions from Arctic soils to surface water corresponded to top-down freezing processes in the active layer. However, there is a lack of information connecting geochemical tracers in both the underlying supra-permafrost flow and the upper soil layer to explain the transport of geochemical compositions to surrounding hydrological networks and can be used as a proxy for the seasonal active layer dynamics.

Therefore, this study aims to provide this knowledge by analyzing supra-permafrost water, soil pore water, surrounding small water ponds/flows, and eventually, river surface water from three typical soils close to rivers from continuous permafrost regions within the Yamal and Gydan peninsulas, West Siberian Arctic. Tundra lakes and river floodplains are abundant in the Yamal and Gydan peninsulas; lakes cover, on average, 10% of the Yamal peninsula, occupying 20% of the

floodplains of large rivers [Y A Dvornikov *et al.*, 2019]. Approximately 90% of all lakes in the study region are small ($<1 \text{ km}^2$) water bodies. To analyze the effluxes of permafrost soils from relatively homogeneous landscapes, the sampling sites were set up next to the largest rivers flowing to the Kara sea, such as the Syoyakha, Murtyyakha, and Taz Rivers. Our main working hypotheses are as follows: (i) Based on the response of riverine DOC and element concentrations to increasing active layer thickness reaches the highest contents during late autumn, when the active layer reaches its greatest depth in a year. Besides, when the components' concentrations in the base flow increases, with the yearly maximum values being observed in the Siberian rivers [D Q Yang *et al.*, 2002]. Therefore, the dominant contributor of DOC and trace elements to river waters during this time of the year is expected to be supra-permafrost water and base flow to the rivers. (ii) When mineral weathering processes continue in the deeper soil column during late autumn and early winter, shallow subsurface soil can also store major mobilized components, such as DOC, in late autumn during freezing; these components bind with elements and can contribute to spring thaw in the next year. Formerly mobilized components stored in the shallow subsurface during spring snowmelt can be potentially enhanced to supra-permafrost waters and further to surrounding surface waters. Therefore, with an increasing thawed depth of permafrost and the potentially enhanced mineral weathering fluxes into supra-permafrost waters, biogeochemical fluxes from surrounding surface waters may experience a change during spring and autumn. (iii) As the DOC and elements are controlled by the redox environment of riparian soils, they cannot be consumed by the surrounding biota or depleted by abiotic reaction, and thence persist in supra-permafrost water; they may be transported to the surrounding hydrological network.

The specific objectives of this study were (1) to assess the mobility of releasing geochemical elements from upper soil layer water and supra-permafrost water to surrounding water bodies; (2) to examine whether the concentrations of geochemical elements in the surface water (connected streams and received river) can explain the active layer dynamics of permafrost riparian soils in different permafrost-affected landscapes.

2. Studying site and methods

2.1 Geographical setting

The study area was located in the Northwest area of the Yamal-Nets Autonomous Region, across 66° N, with continuous permafrost (**Figure 1A**). Sampling sites were located close to the lowlands of the Syoyakha, Murtyyakha, and Taz rivers. The Syoyakha and Murtyyakha Rivers are situated in the Yamal Peninsula on the Western Siberian plain [M Golovatin *et al.*, 2011]. These geographical features of this region are strongly divided between river valleys, lake hollows, and streams [M Golovatin *et al.*, 2011]. Cryogenic soils in this part of the Yamal Peninsula were formed due to the high ice content of Holocene alluvial sediments [A Sidorchuk and V Grigorev, 1998] and aeolian materials [I Alekseev and E Abakumov, 2018]. This part of the lowland comprises very flat watersheds of the Syoyakha and Murtyyakha Rivers and is covered with frozen bogs and palsa. The Syoyakha is one of the largest rivers of the Yamal Peninsula [Y Bespalaya *et al.*, 2018]; its watercourse is 165 km long and its watershed covers an area of 4,400 km² [Y Bespalaya *et al.*, 2018]. The Murtyyakha River has roughly 580 watercourses and numerous lakes in its basin, which contains 70 rivers more than 10 km long. The Murtyyakha River flows into the Syoyakha River and then enters Kara Sea. Syngenetic permafrost close to these two rivers contains massive Holocene ice deposits [Y Vasil'chuk *et al.*, 2016]. The 1,401 km long Taz River is located in the Gydan Peninsula and drains a basin of 150,000 km²; it flows into the Taz Estuary and ends in the Gulf of Ob. Soils along with the river included alluvial stratified parent deposits from an earlier stage of the Holocene [C Tarnocai, 2009].

We investigated the three poorly drained riparian soils with upper organic layers and a downward mineral profile adjacent to each river (**Figure 1A**). The soil close to the Taz River site (TA) was located at the top of the polygon rim, where the landscape is polygonal tundra, and the center of polygon was wet-depressed and an accumulation of organic matter poorly degraded by anaerobic conditions; the surrounding elevated polygon rim showed the evidence

of cryoturbation in most horizons in the active layer (**Figure 1B**). The soil profile was gleyed with frost heaving of the clay stratum. The surrounding vegetation consists predominantly of wet acidic tussock sedge, grass, and sphagnum moss. Soils sampled nearby at the Syoyakha (SY) and Murtyyakha (MU) Rivers were both situated from the floodplain. The SY site consisted of moist acidic moss, lichen, and sedges, and MU site consisted of lingonberry, willow, moss, and sedge. Both soil profiles had shallow active layers that were strongly gleyed and under reducing conditions, while the deeper layers at MU showed obvious signs of cryoturbation (**Figure 1C and D**). The formation of the SY and MU stratified soils may have resulted from the constant changes in the river water levels. All three soil profiles for TA, SY, and MU have been depicted as relatively light chroma color with oxidation of iron minerals and dominated by mineral substrates with allochthonous organic matter or autochthonous peat.

2.2 Soil core and supra-permafrost water collection and analysis

Three soil cores were extracted up to the permafrost table using a portable Snow-Ice Permafrost-Research-Establishment (SIPRE) auger set consisting of a coring auger and an engine from Jon's Machine Shop, Fairbanks, Alaska, USA. The location of each soil core was directly adjacent to the borehole into which a string thermometer was inserted (**Figure 1A**). The thermometers, which were put in cases filled with grease as an inert material, were placed roughly 10 cm from permafrost table for 24 h in July 2016 in order to reach thermal equilibrium with surrounding soil material. The thermometers were connected to a data logger for recording hourly temperatures in degrees Celsius (Modular GM10, Yokogawa Electric Corporation, Tokyo, Japan), powered by a battery with a three-year life and attached to a stainless-steel stake inserted into the soil cores. The excavated local soils were moved back to the soil cores. The cores were collected in May 2016, sealed with Polyethylene covers, transported at a stable temperature of -4 °C to freeze the soils, and stored in a refrigerator at the same temperature prior to analysis. We believe the cores can partially reflect the deepest active layer during a year, with approximately 65 cm for Mu, 46 cm for Sy, and 90 cm for Ta

shown by the thawing materials. The soil pits were excavated in July 2017 adjacent to the coring location (**Figure 1**), and site restoration procedures followed the standard protocols for permafrost-affected soils [C-L Ping *et al.*, 2013]. Excavating soil pits may provide an insight into the vertical depth between the active layer and permafrost that contain external flows to rivers. Each 35 × 35 cm soil pit was excavated with a shovel and hatchet until reaching frozen materials and the genetic horizon and name of each soil were identified using the World Reference Base for Soil Resources WRB [2015] classification system. The stagnic condition of soils meant that we had to use 2,2'-Dipyridyl to spray soils for 30 s to test for reducing conditions, as per the tests previously conducted on gleyed soils [C L Ping *et al.*, 1998].

The frozen soil core was cut into 5 cm sections with a diamond wire saw (LKH-8, Kanghua Company, Guilin, China), with a cutting section of diamond-impregnated wire mounted on idler pulleys, in a cold room in the laboratory of Arctic Logistics Center, Salekhard, Yamal-Nenets Autonomous Region, Russia. The cut sections of each soil core were then placed in tightly sealed plastic bags (9.5 cm x 18 cm, Bag Whirlpak Clear 7oz, Nasco, FL, US) and kept at room temperature (approximately 23 °C) to thaw for 12 h. Completely thawed soils were used to collect soil porewater with an SPS 200 lysimeter soil solution sampler (SDEC, Reignac-sur-Indre, France). The collected water was filtered through a 0.45 µm polypropylene filter, decanted to glass test tubes, and acidified with 0.5(wt) % ultrapure nitric acid (HNO₃, CAS: 7697-37-2 Fisher Scientific, Hampton, USA) before processing through inductively coupled plasma-tandem mass spectrometry (8800 ICP-MS/MS, Agilent, Santa Clara CA, USA). To remove the spectral interferences of ions through ion/neutral reactions, the collision reaction cells (CRCs) mode in the ICP-MS was operated using He or NH₃ as the inert collision gas, following the protocol developed by E McCurdy and G Woods [2004]. The elements V, Cr, Mn, Fe, Co, Ni, Cu, Zn, and As were analyzed in CRCs mode with He gas to remove unidentified polyatomic particles arising from the variables Cl⁻, S⁻, and C⁻. Ti was analyzed in CRCs mode with NH₃ to remove interference from S-based polyatomic ions. The elements Sr, K, Ca, Mg, Al, Ba, Be, B, Cd, Pb, Au, Mo, P, Sc, Ag, Tl, Sn, and rare earth

elements (REEs) were analyzed using normal (non-CRCs) mode.

Four calibration standards (1×10^{-2} , 1×10^{-1} , 1×10^1 , $1 \times 10^2 \mu\text{g L}^{-1}$) were prepared by diluting $1 \times 10^3 \mu\text{g L}^{-1}$ stock standard solution for each element and mixed REEs (TraceCERT®) purchased from Sigma-Aldrich, St. Louis MO USA. The 2% (w/w) ultrapure HNO_3 was taken as the blank. Calibration was performed before every analytical test with a good calibration curve ($R^2 \approx 0.999$), and three blanks and check standards were run before every ten samples. The analytical uncertainty for each sample was determined by the analysis of triplicate within an error of $\pm 3\%$. The detection limits for all elements were set at the level of $1 \mu\text{g L}^{-1}$. To correct the magnitude of the signal suppression or enhancement, a standard mixture solution of Bi, Ge, In, Li, Sc, Tb, and Y ($10 \mu\text{g /mL}$) prepared in 2% ultrapure HNO_3 was used as the internal standard added in all standards, blanks, and samples.

Then, each soil core section was transported to the Applied Ecological Laboratory, Saint Petersburg State University. They were dried in a vacuum drying oven for 16 h, homogenized by grinding using a roller mill, processed through a 2-mm sieve, and finally, soils were pressed into a powder pellet with a vertical hydraulic jack. Each pellet was analyzed through scanning electron microscopy with energy dispersive X-ray spectroscopy (SEM-EDX) (JSM-6390 LA, EX2300, JEOL, Tokyo, Japan). EDX revealed the characteristic peak for each metal in each sample with atomic-level precision, which can directly transfer the concentration in terms of mg kg^{-1} . The soil certified reference material CRM027 and SQC001-30G (Sigma-Aldrich, St. Louis, USA) for trace metals were used for calibration. Samples and the reference material were analyzed in triplicate, and the detection limits were determined based on the minimum value of the reference material.

The location of stagnant supra-permafrost waters was collected adjacent to the previous soil cores in September 2016. To prevent the contamination of water samples due to the previous methods of collecting supra-permafrost waters with soil pit excavation, an alloy probe rod was first inserted into the soil to confirm approximately the same depth of the active layer–permafrost boundary as the previous soil core. A direct push device (SP16

Groundwater Sampler, Geoprobe Systems®, Salina KS USA) comprising a polyvinyl chloride screen was driven to the permafrost table within a sealed steel sheath. Then, the water samples were pumped through the tube to the surface and stored in 25-mL PVC serum bottles by oscillating the tubing up and down.

For the lower floodplain (SY and MU), water samples were collected after pumping the tube for 15–20 min to flush water with permafrost materials. The time required for pumping water in TA was relatively longer than that for the other sites, by approximately 30 min, with less volume of inflowing water because of the higher depth of polygon rim. For all sites, the first portion of water containing permafrost materials was discarded, and the clearer water with fewer impurities was collected. Unfiltered water Samples in the serum bottles were treated with 0.2 mL of mercuric chloride (HgCl_2 , Sigma-Aldrich) and capped without air bubbles or headspace using butyl rubber stoppers pierced by a needle attached to a 3-mL syringe that allows air and water escape from the bottle when the stopper is inserted. With the bottle inverted, 10 ml of methane-free helium (Sigma-Aldrich) was added, meanwhile 10 ml of water was removed through needle displacement and the syringe. Then, another 10 ml of methane-free helium was added to the headspace without removing water, and water and headspace were equilibrated in the bottle by shaking for 2 min. The triplicates of subsample headspace gas (into the syringe) were analyzed for CH_4 and CO_2 by gas chromatography (GC-456, Bruker, Billerica MA, USA), with flame-ionization detection (FID) and electron capture detection (ECD), respectively. After each set of ten samples, the detectors were calibrated based on Air Liquide ($\text{CO}_2 = 246.6 \mu\text{mole/mole}$, $\text{CH}_4 = 302.3 \mu\text{mole/mole}$). The results of triplicate injection showed a repetition rate within $\pm 2\%$. The gas solubility of CH_4 and CO_2 [P Kastanidis *et al.*, 2018] were taken by calculating the total concentration of CH_4 and CO_2 in the vials and then converting to $\mu\text{mol/L}$ of the initial samples. All analysis procedures for analyzing CH_4 and CO_2 were carried out in the Center for Chemical Analysis and Materials Research of St. Petersburg State University. The water temperature, dissolved oxygen, pH, and specific conductivity were field measured with a portable multiparameter

meter (Orion Star™ A329, ThermoFischer Scientific, Waltham MA, USA). Stagnant supra-permafrost waters in this study were oxygenated with an average O₂ saturation in samples ranging from 10.2 to 82.3% with an uncertainty of 3-4%, which is very close to the previous reports in Western Siberian lowlands [S V Loiko *et al.*, 2017; T V Raudina *et al.*, 2018]. No significant difference was observed in the O₂ concentrations from water samples in each soil site. The average temperature of supra-permafrost waters did not vary significantly as 12.3±1.8 °C for SY, 13.4±1.6 °C for MU, and 15.6±2.0 °C for TA. The dissolved organic carbon (DOC) concentrations were analyzed by a TOC-L TOC Analyzer (Shimadzu Kyoto, Japan) with an uncertainty of 5% and a detection limit of 4 µg/L.

2.3 Surface water collection and analysis

All surface water samples were collected in September 2016. All sampling sites were adjacent to the location of the soil cores. We collected water samples (streams connected to each river) from shallow (< 10-30 cm) permafrost subsidence and hollows with the size of < 1 m²; deep depressions (< 1 m) of palsa bogs with sizes ranging from 5 to 15 m²; small thaw ponds with sizes ranging 10-150 m²; and each river closest to the soil core within 5–10 m (depth < 5 m). Neoprene gloves were used during water sampling, and water from the shoreline was collected by a standard PVC MP² two-stop peristaltic pump tubing (1.09 mm I.D., White/Red, Pkg. 12, PerkinElmer, Waltham MA, USA) outfitted with a pre-sterilized Durapore® 0.45 µm capsule filter (MilliporeSigma, Burlington MA, USA). The water passing through the tubing and filter capsules for the first 30 s was not collected, and polypropylene (PPCO) bottles with white PP closure precleaned by 2% nitric acid were used for water collection. The analysis of trace metals in water samples was the same as in subchapter 2.2. The samples used for chloride determination were not acidified. Chloride determination was performed by titration with silver nitrate (AgNO₃, 0.02 mol L⁻¹) as titrant according to ISO 9297: 2000 “Water quality — Determination of chloride — Silver nitrate titration with chromate indicator (Mohr's method).” Titration calibration was used by

standardizing AgNO_3 solution versus NaCl solution with the same molar concentration (0.02 mol L^{-1}). The relative standard deviation (RSD) of titrant was $<1 \%$.

3. Results

3.1 Characteristics of soil profile and chemical composition in the soil column

The detailed horizon and vegetation information in the soil pit profiles in the photographs for SY, MU, and TA is shown in **Figure 1**. The soil group belongs to Histic Gleysols (Stagnic), Histic Gleysols (Turbic), and Turbic Cryosols for SY, MU, and TA, respectively. The top fractions of soil horizons from three soil pits all show a high chroma color, with TA being relatively lighter (**Figure 1B**), and all the lower fractions show strong gleyed conditions, which may indicate a redox boundary. The lower fractions of soils were further tested by a few drops of alpha-alpha-dipyridyl for 30–40 s, showing that no pink color was observed for SY at the depth of 6–9 cm, with a strip-shaped iron oxidized boundary to the next horizon. A positive pink color was observed during the reaction in the next gray color horizon (9–18 cm), suggesting the occurrence of reduced ferrous oxides under the reducing condition. Compared with the previous horizon, this horizon had few plant roots and little accumulation of organic matter. The lowest section (18–35 cm depth) had a sporadic positive pink color. This difference may be due to the different redox spots visually observed in the excavated soil pit (**Figure 1C**). The lower 18–35 cm showed heterogeneous rusty-gray color as the ferric irons, as well as aluminum, and inclusions of organic materials due to the cryogenic mass exchange through bottom-up transportation of water by thawing/freezing processes that bring parent materials to the upper layers, where cracks can be observed in 18–35 cm. In the upper 6–9 cm horizon, the oxidation of ferrous iron to ferric iron is mostly due to the leaching process of upper thawing water containing oxygen. Therefore, the low density of strictly reduced gleyed horizon (6–9 cm) separated the two different geochemical transport pathways between the upper and lower horizons.

For soil pit MU, the pink reaction color was observed at the depth of 7–25 cm (the shape

shown as B_{g@} horizon in **Figure 1D**), and the middle part of depth 25–60 cm and 60–65 cm depth. In the B_{g@} horizon, there were more disbursed rust-orange spots and stains that followed the path of roots; these ferrous irons were oxidized by the atmosphere through the growth upper vegetation's roots growing. In the 25–60 cm depth, the apparent presence of cryoturbation with a gray-brown color was observed alongside the dark-gray inclusions from the lower permafrost. The oxidized iron in this horizon results from the multi-year freezing/thawing and lower activities of congeliturbation that transport the materials from both the upper and lower layers. The 60–65 cm depth was completely under the reducing condition with loamy soils in gray color.

For soil pit TA, the reducing conditions indicated by a positive pink color were observed below 55 cm. TA shows a mixed O and A horizon within 30 cm that has a brownish-gray color and very few roots, which is common under hummocks in Western Siberia. Within the 0–55 cm depth, the lateral ring-shaped layers and mud spots can be observed, which were caused by frost heave when a rounded knoll of ice rises during the freezing time. The evidence of cryoturbation can be seen below 55 cm with the dark gray color, and some moss residues are observed in the fragments of cryoturbation clusters.

The vertical elemental concentrations in three soil cores measured by EDX are listed in **Table S1 (Supporting Information)** and plotted as logarithmic concentrations in soil core depth in **Figure 2**. The other elements not shown in **Figure 2** did not exhibit a regular pattern under reducing conditions within the soil cores. Mn and Co were enriched in the top mineral layer with organic matter accumulation, which likely because of their complexation with natural organic matter. In contrast, Cr, V, Sr, Ca, Mg, Al, Ti, Cd, Mo, Zr, and Sc increased in lower mineral soil horizons in the three soil cores. In particular, for the MU site, Fe, Cr, and V were only higher in the first gleyed horizon with cryoturbation (7–25 cm) and decreased gradually with increased depth. Al, Ti, Cd, and Sc kept increasing until 40 cm (second gleyed horizon) and then decreased gradually or maintained relatively comparable levels in the deeper layers. As, Mo, and Zr decreased until 40 cm and then increased in deeper mineral

horizons mixed with frozen material and at the frozen horizon. Zn, Sr, K, Ca, Na, Mg, and Y were higher in all gleyed horizons.

For the SY site, three trends were observed: *i*) relatively slight decrease from top to down within the gleyed horizons for As, Cr, V, Mo, Zr, and Se; *ii*) continuous increase in gleyed horizon depth for Sr, K, Ca, Va, Mg, Al, Ti, and Pb; *iii*) only the Cd content increased sharply in the upper gleyed horizon (9–18 cm) and decreased significantly below 18 cm. For the TA site, the maximum concentration was observed in the gleyed horizon for Cr, V, Sr, Ca, Mg, Al, Ti, Cd, Pb, Mo, Zr, Y, and Sc. These elements were apparently enhanced in reducing fractions, concentrations of which, with respect to the upper layers of soil column, can reflect the soils with increased reducing soil conditions. This obvious demarcation between elements with concentrations either increasing in the upper layers of soil column or accumulating in the lower part of soil column correlates to the variation of redox conditions in the soil pit profile as well. The fluctuation of elemental concentrations in the lower soil cores may also be caused by the effect of cryoturbation.

The elemental concentrations of soil porewater are listed in **Table S2** and plotted as the function of depth along the cores (not containing supra-permafrost water) in **Figure 3**. The results showed Ca, Mg, and Ti as soluble species that may be available for transporting supra-permafrost water for all three sites because of the increased concentrations in the deeper fraction of the soil core. Additionally, Al in TA site, Mg in MU site, and Fe in SY site may also have the same potential. To some extent, Mn, Ca, Mg, Al, Ti, and Fe all can be soluble species in both upper and deeper fractions of the soil core, indicating both the transport capability from surface flows and supra-permafrost waters to the river.

3.2 Thermal dynamics within soil depth

The temperatures in multiple depths from the installed thermometers are provided in **Figure 4**. In general, the surface (0 cm) soil temperature corresponds to the variation in ambient temperature in this region. The soils at four different depths in TA, MU, and SY were completely frozen by the end of April as our measurements were initiated. In TA, the thawing

of frozen soils (temperature rose above 0 °C) occurred on May 30, 2016 at 0 cm, on June 7, 2016 at 20 cm, on June 18, 2016 at 40 cm, and on June 30, 2016 at 75 cm. In MU, the thawing occurred on June 2, 2016 at 0 cm, on July 14, 2017 at 25 cm, on June 25, 2016 at 40 cm, and on July 4, 2016 on 59 cm. In Sy, the thawing occurred on June 1, 2016 at 0 cm, on June 11, 2016 at 18 cm, on June 22, 2017 at 26 cm, and on July 2, 2016 at 38 cm. The frozen soils in the Arctic notably thaw from top down during the spring-summer season. The lower latitude of TA results in a relatively earlier thawing time than that of MU and SY.

3.3 Chemical composition in the supra-permafrost water

All measured elemental concentrations are plotted as the function of sampling time in **Figure 5** and the pH values for supra-permafrost water in TA, SY, and MU for different sampling months are presented in **Figure S1** with a detailed description in **Text S1**. In TA, the contents of Zn, Mn, Ti, and Sr increased generally from the beginning of summer (June) while Sr and Ti increased slightly with an obvious increasing trend at the end of summer (September 15, 2016). Zn reached its highest concentrations (15.93-17.50 $\mu\text{g L}^{-1}$) at the beginning of autumn, and this level was maintained until October 2016. Unlike Zn, the peak value of Mn was observed in October. Fe concentrations started to rise significantly at the end of September and peaked at the end of October 2016 (2049.71 $\mu\text{g L}^{-1}$). Mg, Cr, V, Pb, Al, and Ca concentrations exhibited less fluctuation over summer with low concentrations until an increasing trend was observed at the beginning of autumn. Na was the only element that showed a gradual increase in concentration over time till October 15, 2016. For Cd and Mo, there was no particular trend, and their levels were low. SY and MU essentially followed the pattern of TA, while the highest Fe concentration in SY was found at the end of October 2016.

3.4 Chemical composition in the surrounding hydrological streams and rivers

We measured the elemental concentrations of the surface water surrounding the soil cores, including the connected hydrological flows (i.e., shallow, depression, and thaw ponds) and

rivers (the sites receive the flow), which are shown in **Figure 6**. The pH values for TA, MU, and SY for different months are shown in **Figure S2** with detailed description in **Text S2**. The concentrations of CO₂, CH₄, and DOC in the surrounding streams are plotted as the function of sampling time in **Figure S3** and described in **Text S3**.

The remaining detected elements were not shown here because they were below the detection limits. The three sites showed strong similarity of elements' distribution in the connected surface flows and the adjacent rivers with the passage of time. Most elements (i.e., Fe, Mn, Zn, Na, Pb, Al, and Ca) showed a dramatically increasing trend from the beginning of September to the beginning of October, while Co, V, and Sr significantly increased beginning in August, and only Cr increased starting in July. These elements in the surrounding hydrological streams or rivers reached their highest concentrations at the beginning of October, except for Mo, Cd, and Ti, which had very low concentrations at all time-scales with a slightly increased concentration in October. The concentrations of Fe, Zn, Al, Co, and Sr were consistent with the observation of enhanced soluble concentrations in supra-permafrost water for each of the corresponding months. This may be because of the solubility of these metals, except Co and Mn, which remained at low levels in the top layers of soil cores, and because of the different solubilities of the metals during snow melting, which are related to the bonding mechanism in the soil, especially for top soil layers with increased organic matter. The highest to lowest concentrations for metals were approximately $\text{Fe} > \text{Mn} > \text{Zn} > \text{Sr} > \text{Ti} > \text{V} \approx \text{Na} \approx \text{Al} \approx \text{Pb} > \text{Cd} \approx \text{Mo}$.

4. Discussion

4.1 Variability of elemental concentrations in solid soil and soil porewater

In the TA soil column, higher elemental concentrations in soil porewater, such as Mn, Mg, and Ti, were found in the oxidizing zone (~10-15 cm, the layer with organic matter accumulations). Compared with the deeper depth of soil column (~30-83 cm, reducing zone),

enhanced elemental accumulations were also observed, especially for Mg and Ti. The lesser extent of Ca and Al concentrations in the top organic layer (**Figure 2**), which may be due to the excess of Ca in the soil solutions, occupy more than 90% of total cation concentration [J O Leckie, 1986]. Ca cation can control the soluble stage of trace elements in soils, due to Ca's presentation as an organic complex in soil solutions, preventing the precipitation of soil deposits [J O Leckie, 1986]. For instance, fulvic acid in this organic layer can significantly interfere with the crystallization of aluminum hydroxide polymorphs [H Kodama and M Schnitzer, 1980]. Besides, the aqueous Ti concentrations in both the oxidizing and reducing zones are similar, while Mn is relatively higher in oxidizing zones, and Mg is more enhanced in the reducing zone. Compared with the distribution of these elements in the solid soil core, only Ca and Al showed a similar trend. This difference is due to the soluble and chemical fraction of species.

In MU soil column, the concentrations of Mn, Ca, Mg, Al, Fe, and Ti in soil porewater were increased below 7 cm in the reduced zones. However, the solid soil phase shows high levels of Mn in the oxidizing zones (~0-7 cm), with the other elements' distribution corresponding to the soluble forms. In SY soil column, Mn, Mg, and Fe contents in soil porewater were enhanced in the upper oxidizing zones, and Ca content was only obviously enhanced in reduced zones, while Al and Ti both accumulated in the oxidizing and reduced zones. The Mn concentrations were found to be the highest in the oxidizing zones in solid phase, whereas other elements (Ca, Mg, Al, Ti, and Fe) increased in the reduced zones.

In both MU and SY, Mn concentrations are found higher in top layer of soil, which is consistent with the low solubility of Mn in lower pH soils (pH = 4.7 for Mu and 4.8 for Sy) [F Scholz and H Kahlert, 2015]. We speculate this is because of a more integral gleyed layer (~10-40 cm) caused by cryoturbation, which makes the soils share the same origin. SY revealed more heterogenous materials that divided several soil layers through cryoturbation, which results in elemental fluctuations in terms of soluble elements.

The three soil cores' depth represents the typical maximum vertical extent during the

thawing period in autumn. Among the three soil cores, differences between elemental concentrations in soil porewater and solid soil are still substantial. Taking as an example the concentrations of Mn in the Ta soil column, the concentration of Mn in the oxidizing layer is 1039 mg/kg⁻¹ with approximately 0.11 mg/L⁻¹ in soluble form. There is approximately 0.045 mg/L⁻¹ mobilized within thawing water to the reduced zones with 388 mg kg⁻¹ presented in solid soil of this layer.

To better understand the solubility of elements, the partitioning coefficients (K_d) were calculated as shown in **Figure 7**. Despite the discrepancy of K_d below the top layer, the decreasing solubility can be observed within the gleyed layers, which is consistent with the theory that metals are hindered in this layer [I Antcibor *et al.*, 2014]. However, Al and Mn were observed with clearly increasing solubility in all soil cores above the permafrost table. Besides, the high partitioning from the solid to the aqueous fraction was observed in Ti for Sy, and Fe and Mn for Mu. Unlike the top organic-rich layers mainly reflecting biogeochemical cycle, minerogenic layers reflect mineralogical weathering. Therefore, the difference may be due to different soil textures and frost processes.

The enrichment of Mn and Al for soil materials in Western Siberia was previously investigated [E Abakumov *et al.*, 2017; I Antcibor *et al.*, 2014; L Evgeny *et al.*, 2017; X Ji *et al.*, 2019a; X Ji *et al.*, 2019b]. The further soil water flows into supra-permafrost water, the more it creates consistent input of elements such as Mn and Al from soil to solution. The previous study showed that the solubility of Mn increases with increasing soil acidity [E Andrade *et al.*, 2002], which is consistent with our observation. A J Barker *et al.* [2014] first reported that trace metals concentrations (Al, Ba, Fe, and Mn) correlate with the seasonal thawed active layer in the Alaskan arctic. This also proves that elemental transport occurred through thawing soil porewater with the potential for those elements to enter surrounding water bodies.

4.2 Elemental fluctuation in supra-permafrost water and surface water by seasonal controls

To some extent, except for Mn and Al, some other elements also showed high solubility in the bottom of the three soil cores (**Figure 7**). We expected to see high mobilization of Mn, Al, Ca, and Ti to the supra-permafrost water during spring snow thawing. However, except for Mn, no or very slightly increasing concentrations in the late May / beginning of June can be found in the supra-permafrost water (**Figure 5**), which may be due to the low volume of thawing water or the equilibrium between elemental pool size in solid soils and snowmelt water in the reducing zones at this time. We found the peak concentration time for most elements were shown in late autumn / the beginning of winter. Compared with elements in soil porewater, more elements were above the detection limits and at distinguishable higher levels during the August to October time period. This may indicate that the origin of elements in supra-permafrost water may not only originate from the dilution of snowmelt water within the soil core, while other water flows through the slope of the permafrost table in the landscape. The relatively higher levels of elements such as Mn, Zn, and Fe in TA, than that of SY and MU, could result from the frost heaving of the polygon rim, which released more frozen materials. During summer, thawing starts in the permafrost-affected soils leading to a deeper active layer, which can be shown by the slight increase in Mn, Fe and other elemental concentrations in the supra-permafrost water over the course of June, July, and August. This increase is correlated with the low coefficients in the bottom of the soil core, and with the soil temperature as well. The soil thermal stratification shows that soil temperatures were above 0 °C at 0, 20, and 40 cm for TA from late June; in 0, 25, and 40 cm for MU from beginning June; and in 0, 18, and 25 cm for SY from late June whereas the deeper depth (75 cm for TA, 59 for MU, and 38 for SY) was still frozen at this time.

Most elements revealed relatively high solubility with small K_d values at the deepest depth. In Ta, Mn follows this trend at the depth 18-35 cm, whereas at the deeper depth of 35-40 cm the solubility decreases. The high mobility of these elements in the deeper soil column, along

with the latter thawing for these layers, could account for high concentrations in supra-permafrost water and surface water in late September and beginning of October. This pattern is relatively comparable to other elements in TA, MU, and SY, except for Ca and Ti in MU. Generally, elemental fluctuations in the surface flow and river were the same as the supra-permafrost water. The concentrations of Zn, Mn, and Ti in the rivers from July to October were higher than the surface flow, indicating the mobility of these elements to the rivers from the surface flow and include the possibility of contribution to this input from supra-permafrost water. The previous study also showed some metals like Fe, Al, Mn, and Ba decreased in the active layer with subsequent thawing and increasing surface water concentrations during summer to autumn and could indicate elemental transport signals corresponding to the depth dynamics of the active layer [A J Barker *et al.*, 2014]. Our results are consistent with the theory that major and trace elements export to the lakes and rivers through the boundary between the thawed and frozen layers [T V Raudina *et al.*, 2018]. However, we cannot exclude the possibility of abrupt permafrost collapse in the surrounding water body and rivers, which has been shown to enhance metals release into surface water [S V Loiko *et al.*, 2017]. This may also be a reason that not all elements in the surface water were above the detection limits in the soil porewater.

4.3 Potential elemental signatures as a function of organic matter in the surface flow

The discharge and source of water are important for the transport of geochemical components such as elements and organic carbon to surface waters. However, it is difficult to determine water sources in cryogenic landscapes with large amounts of ice and multiple flows. In the present study, DOC is used as the “participating media” to estimate elements transported to the surface water; these are considered to be the major carriers of trace elements in boreal and permafrost-affected organic-rich surface waters [B Lyvén *et al.*, 2003; Q Ma *et al.*, 2019; E Neubauer *et al.*, 2013; R D Rember and J H Trefry, 2004; E V

Vasyukova et al., 2010]. Due to no pollution from domestic and industrial activities in our sampling sites, the riverine organic carbon is usually subdivided into two origins as the allochthonous pool derived from terrestrial organic matter (topographical erosion and soil leaching) and the autochthonous pool derived from in-situ phytoplankton production [*D Hope et al.*, 1994]. Additionally, an empirical model for organic carbon exports showed DOC exports to surface runoff is mostly driven by the unfrozen zones during permafrost thawing [*C Fabre et al.*, 2019]. Therefore, the surface streams with smaller water volume connected between the soils and the rivers were investigated for the relationships between DOC and elements, and the elemental relationship among the streams, rivers, and supra-permafrost waters.

The soil in the three study fields all have a top organic layer with sporadic organic distributions in the mineral layers. Previous studies reported that precipitations would form in the rapid surface runoff through supper soil layers with a depth of approximately 15–20 cm [*K H Bishop et al.*, 1993; *D Hope et al.*, 1994]. DOC would be exported to the surface water from the top layer without infiltration in the riparian zone. In TA, the highest concentrations of CO₂ and CH₄ were observed in the summer snowmelt season with an increasing thawing degree while low concentrations were observed (**Figure 8**), which may have been due to an abrupt thawing process from the top organic layer. This was consistent with the maximal microbial metabolism occurring at the boundary between thawed and frozen permafrost soils in the Western Siberia lowland [*Y N Morgalev et al.*, 2017]. However, *P J Dillon and L A Molot* [1997] reported that the release of CO₂ into the boreal waters was increased when the DOC concentration was elevated in the thermokarst lakes. We found a slight decrease in CO₂ and CH₄ when the DOC concentrations reached its peak in Autumn in TA. However, a different pattern was observed in SY and MU in that CO₂ and CH₄ did not increase significantly during snow-ice melting in the spring and summer seasons, while an increasing trend was observed for DOC contents in the surface water. *R M Manasypov et al.* [2015] reported no significant enrichment of CO₂ and CH₄ in the thermokarst water bodies during the

spring flood through the base flow and further observed that CO₂ levels only increased in small depressions (<10 m²), which is consistent with the ponds (approximately 1 m²) that we sampled. Although some obvious buildup of CO₂ was found in the summer for SY, the results are still comparable to the phenomena in high latitude lakes [*J Karlsson et al.*, 2013]. The reason for the contrast between TA and the other sites (SY and MU) could be caused by the low volume of the water pond in SY, and in MU due to the low ice contents and a short period suitable for this type of accumulation in low elevations. However, there is also a possibility that more ice existing in polygon rim and buried talik would lead more the more unfrozen water flowing to the surrounding streams. Combination of the results of three sites shows that the response mechanism of enhancing CO₂ in the streams during snow melting is heterotrophic respiration of allochthonous DOM, as approved by the constantly elevated DOC concentrations in this period. However, we suggest the bio-utilization of these carbon types in small ponds may vary according to the contents of materials by different water volumes.

The elements with high mobility in the bottom gleyed layer of the soil column and obvious higher concentrations during the snowmelt period were plotted against the DOC (**Figure 8**). In Ta, Mn increased sharply in the summer while Al, Ca, and Mg did not significantly increase. Peak values of Mn, Al, Ca, and Mg with DOC were observed in autumn. The same pattern was observed for SY and MU, especially for Mn. This may indicate the release of micronutrients as Mn during vegetation activity and upper moss litter leaching in the warmer water, which allows higher mobility within the soil column. Because we did not observe a high concentration of Mn in the uppermost soil layer (**Figure 2**), we speculate this contribution of Mn may be from the supra-permafrost water.

Another difference found in the TA site was that elemental concentrations (Al, Ca, and Mg) in autumn were significantly higher in comparison to MU and SY, which may be a result of upwelling via the icy cracks beginning as early as October, producing organic- and Fe-accumulated allochthonous ice crystallized at the pond surface. The maximum values of these elements peak during autumn in the surface water are about 2 to 3 times greater than in

spring and summer, representing the highest water flow including DOC and dissolved elements.

Additionally, the strong linear correlation of Mn between surface streams and rivers was found for all three sites, and that of Al between the surface streams and river was found for Ta (Figure 9). However, when comparing the surface streams and supra-permafrost water, Mn was correlated for all three sites; Al only for TA and SY; and Ca only for TA (Figure 10). For Mn, the snowmelt flows had higher Mn contents attributed to the increasing mobility of Mn species to other elements in superficial layers of soil as previously reported [A J Barker *et al.*, 2014]. Supra-permafrost water also initially contributes to the surface flows and then further flows to the river. The differences for Al and Ca may be due to the different water–soil interactions. Therefore, Mn could be a seasonal signature of autumn representing the dominance of overland surface flow and maximum interactions with the active layer when the active layer is the deepest.

5. Conclusion

This study found that the levels of elements did not begin to increase when the surface temperature is above 0 °C, and the deeper soil column temperature is above 0 °C in late summer/early autumn. The patterns for surrounding water ponds, adjacent river surface waters, and supra-permafrost waters were generally comparable for most elements above detection limits (Zn, Mn, Sr, Fe, Mg, Cr, Co, V, Pb, Al, and Ca) and were highest during the period from September to October, corresponding to the deepest depth of the active layer. Although a geochemical barrier in the reducing gleyed layers was observed in the soil column, some elements (Mn, Ca, Mg, Al, and Ti) still have high mobility above the permafrost table. Additionally, Mn may flow into surrounding water flows (to the river) by both upper soil and supra-permafrost waters transporting organic matter including CH₄ and CO₂. However, heterogenous landscapes with more ice and cryoturbation may cause more elements to move to the surface waters in the present study. Mn in surface flow may be a proxy for the active

layer process during the period of summer and autumn. Highly dynamic Arctic hydrological systems that receive geochemical elements from surrounding permafrost soils and delivering more DOC and elements to adjacent hydrological systems. Future climate conditions will intensify the soil–stream–river process due to permafrost degradation and the dynamics of the active layer.

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Figure captions

Figure 1. Location of the sampling area and installed thermometer (A). Soil pit profiles and their upper vegetation species near rivers: (B) Taz river (67°30'13.3"N, 78°40'56.2"E), (C) Syoyakha river (69°57'03.6"N, 71°23'15.2"E), and (D) Murtyyakha river (70°06'04.7"N, 68°40'24.1"E) in Yamal-Nets Autonomous Region. All soil profiles are gleyed and have oxidation/reduction horizons in the upper active layer, substantial top organic layer, and cryoturbation in the lower part of the active layer.

Figure 2. Vertical metal/element distribution in soil cores collected from TA (Taz river), SY (Syoyakha river) and MU (Murtyyakha river) sites in the Yamal-Nets Autonomous Region in beginning May 2016. The logarithmic transformed distribution shows the elements' enrichment relevant to the top layer with humus accumulation in an oxidizing condition (Mn and Co) and the lower gleyed layer in a reducing condition.

Figure 3. Distributions of soluble metals along the vertical soil water extracted from the soil cores collected adjacent to the Ta, Sy, and Mu rivers in the Yamal-Nets Autonomous Region.

Figure 4. Soil temperature (°C) at different soil depths within the active layer as function of time in 2016. The soil in the Russian Arctic (TA, SY, and MU sites) experiences a top to down freezing process, making soil porewater flow downwards to deeper soils above frozen materials (permafrost table).

Figure 5. Soluble element concentrations (µg/L) in the supra-permafrost water collected from the Ta, Sy and Mu sites as a function of date in 2016, respectively.

Figure 6. Soluble element concentrations (µg/L) in the surface water from connected hydrological streams to rivers and rivers from the TA, SY, and MU sites as a function of date in 2016, respectively.

Figure 7. Plot of partition coefficients (K_d) for the soil cores from the TA, SY, and MU sites, sampled in the late September 2016. K_d values were calculated by the ratio of elemental concentrations (mg/kg) in the solid soil phase to the elemental concentrations (mg/L) in soil porewater.

Figure 8. Elements/CO₂/CH₄-dissolved organic carbon (DOC) relationships for the surrounding hydrological streams in TA, SY, and MU sites over the course of spring to autumn 2016.

Figure 9. Linear correlations of the elements with high mobility in soil columns between the connected surface streams and receiving rivers.

Figure 10. Linear correlations of the elements with high mobility in soil column between the connected surface streams and supra-permafrost water.

Figure 1.

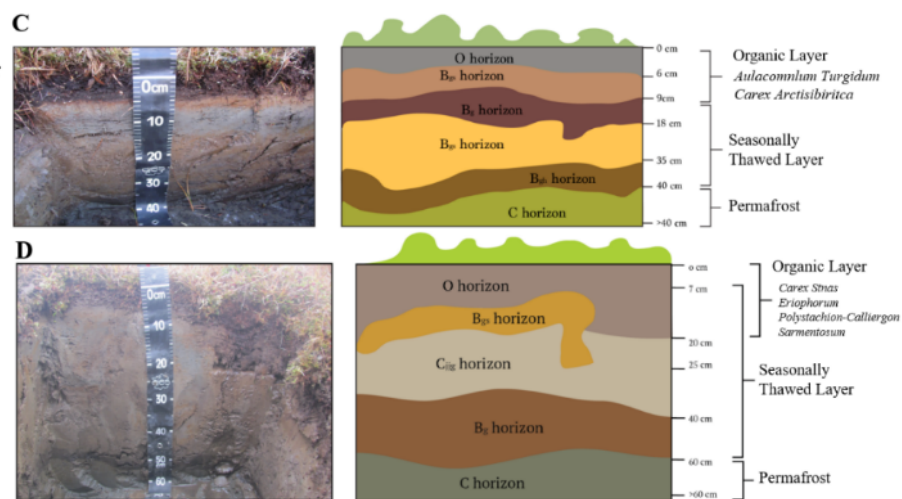
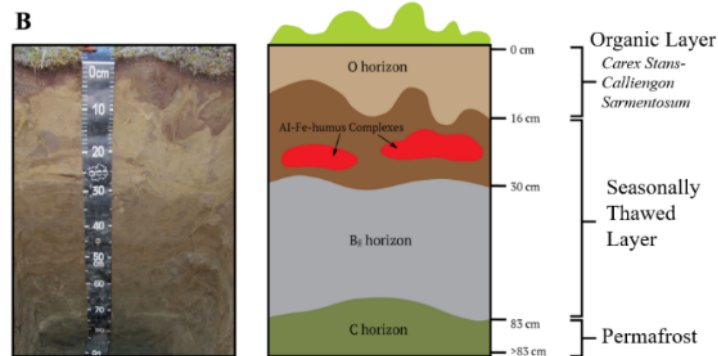
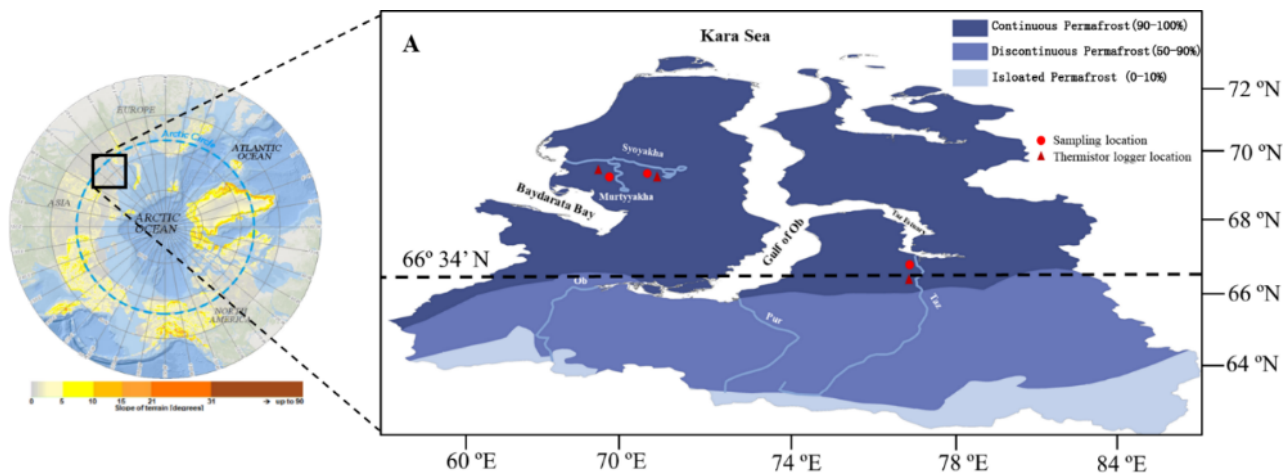


Figure 2.

Figure 3.

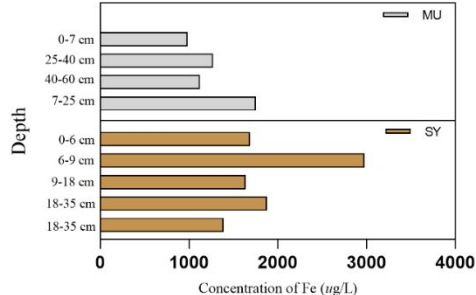
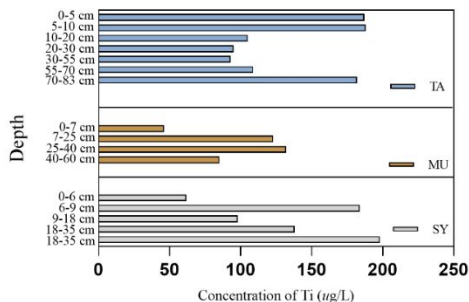
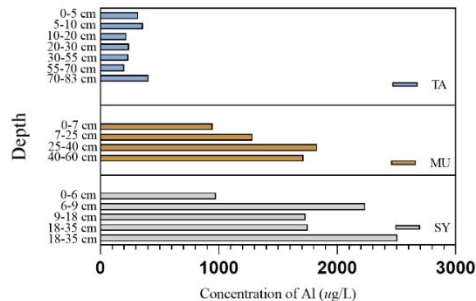
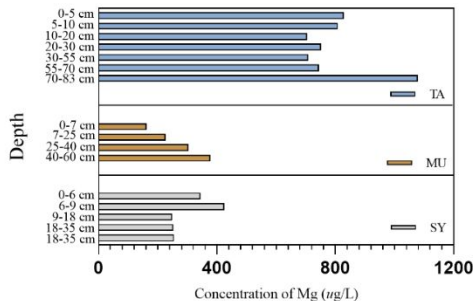
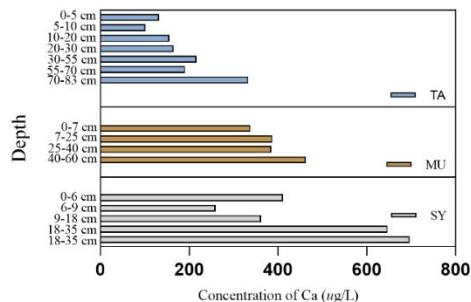
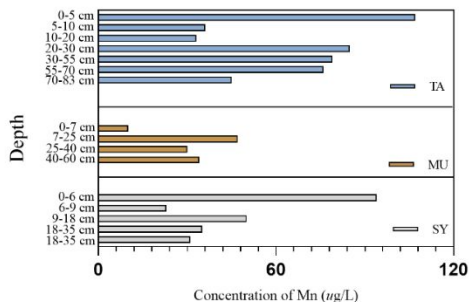


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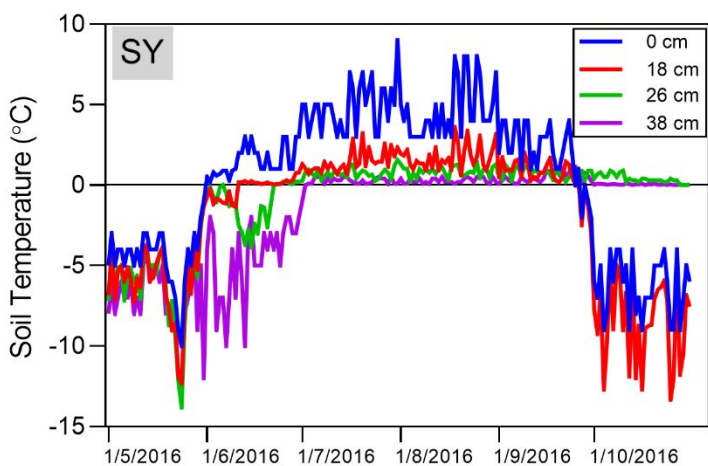
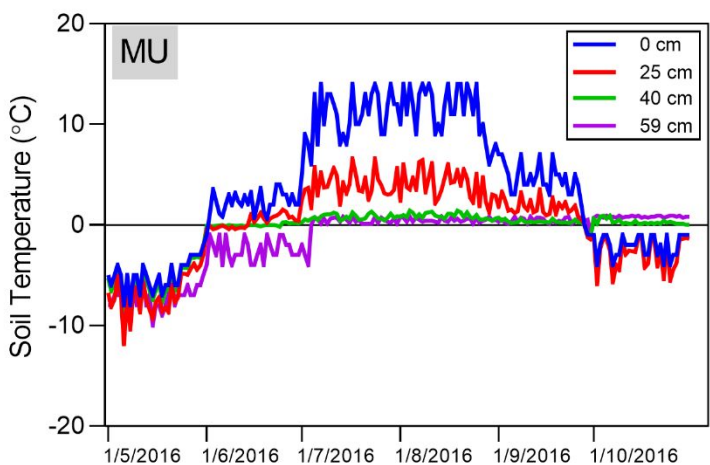
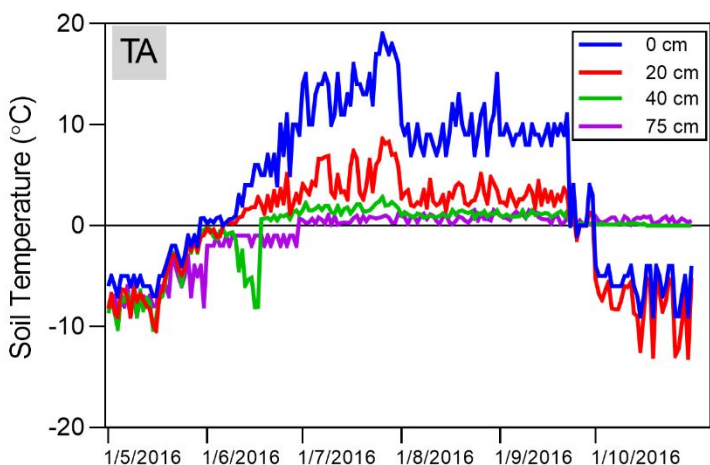
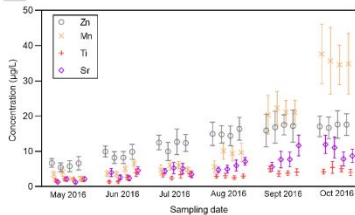
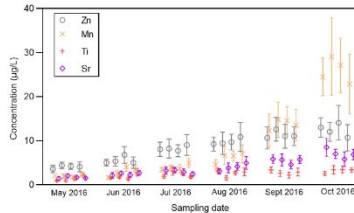


Figure 5.

TA



SY



MU

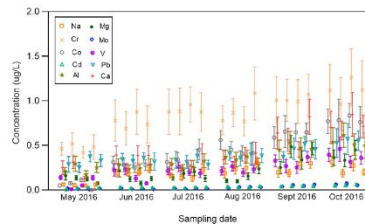
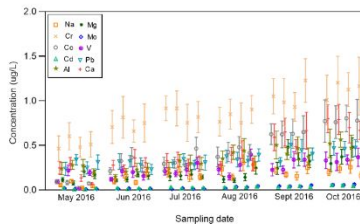
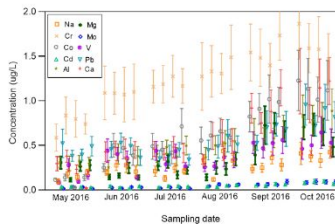
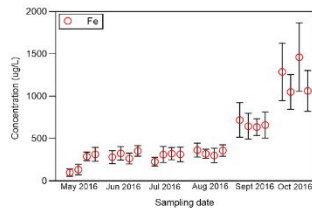
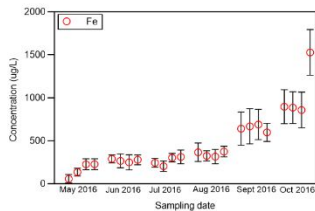
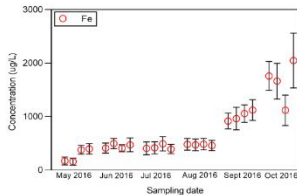
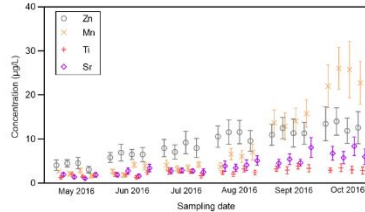
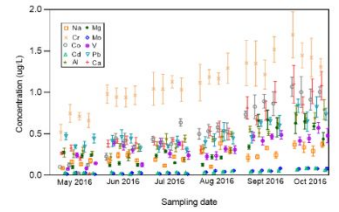
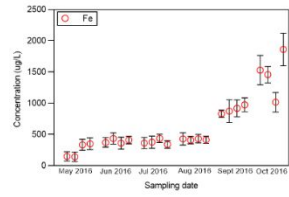
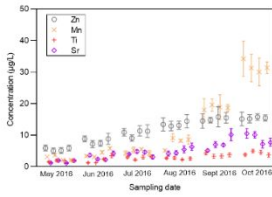


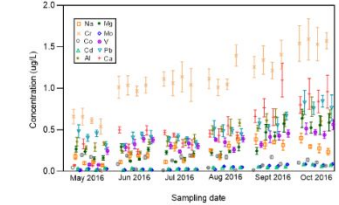
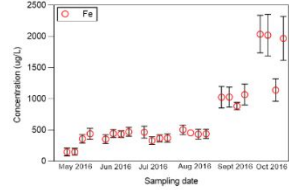
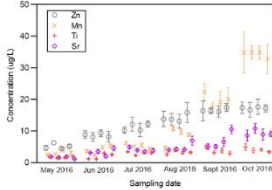
Figure 6.

TA

Surface flow

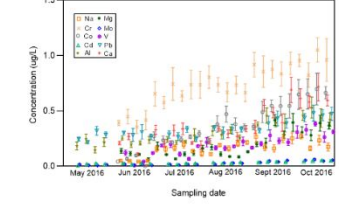
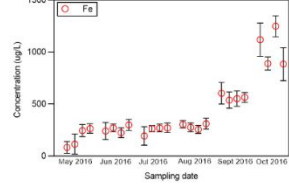
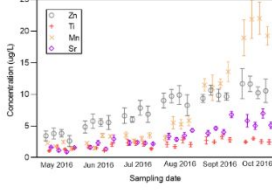


River

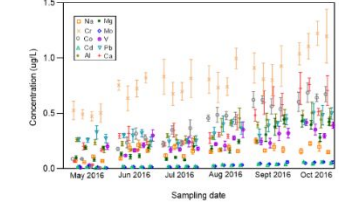
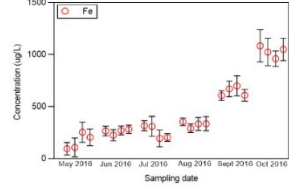
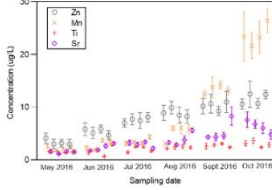


MU

Surface flow

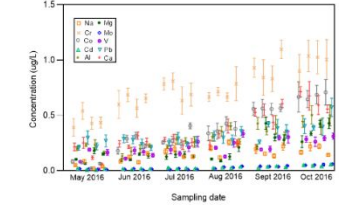
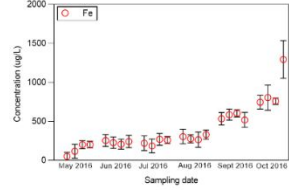
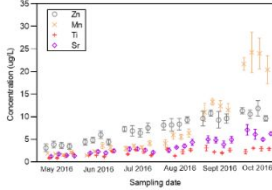


River



SY

Surface flow



River

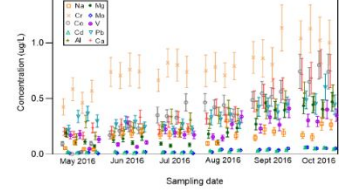
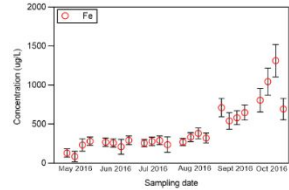
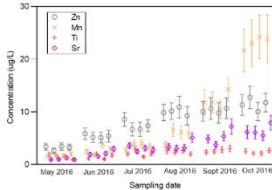


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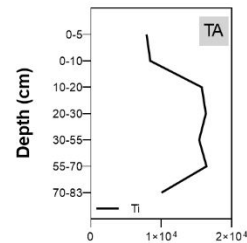
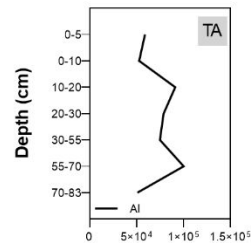
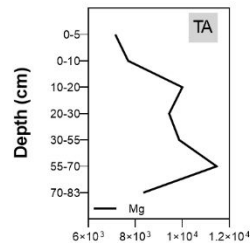
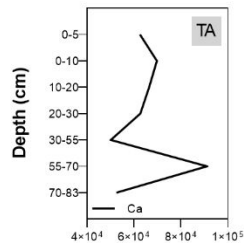
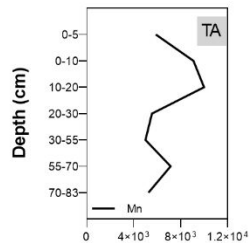
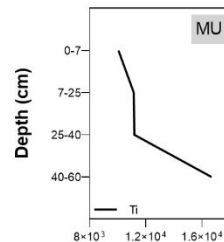
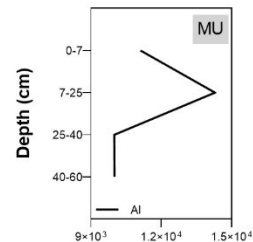
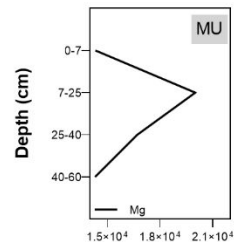
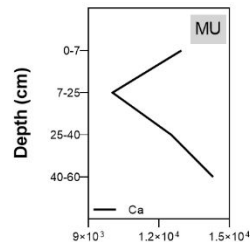
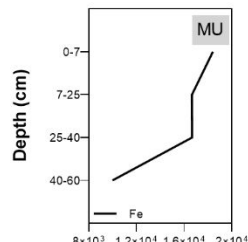
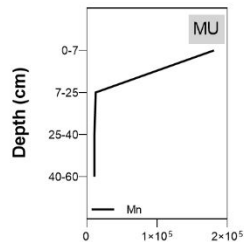
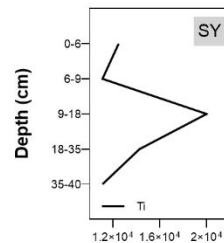
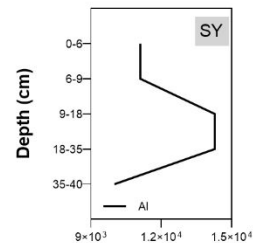
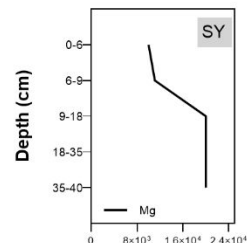
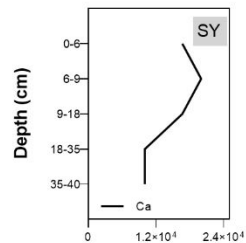
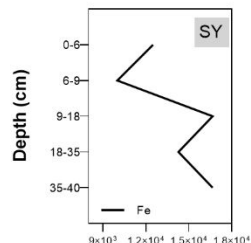
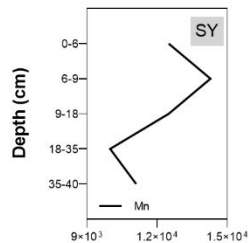


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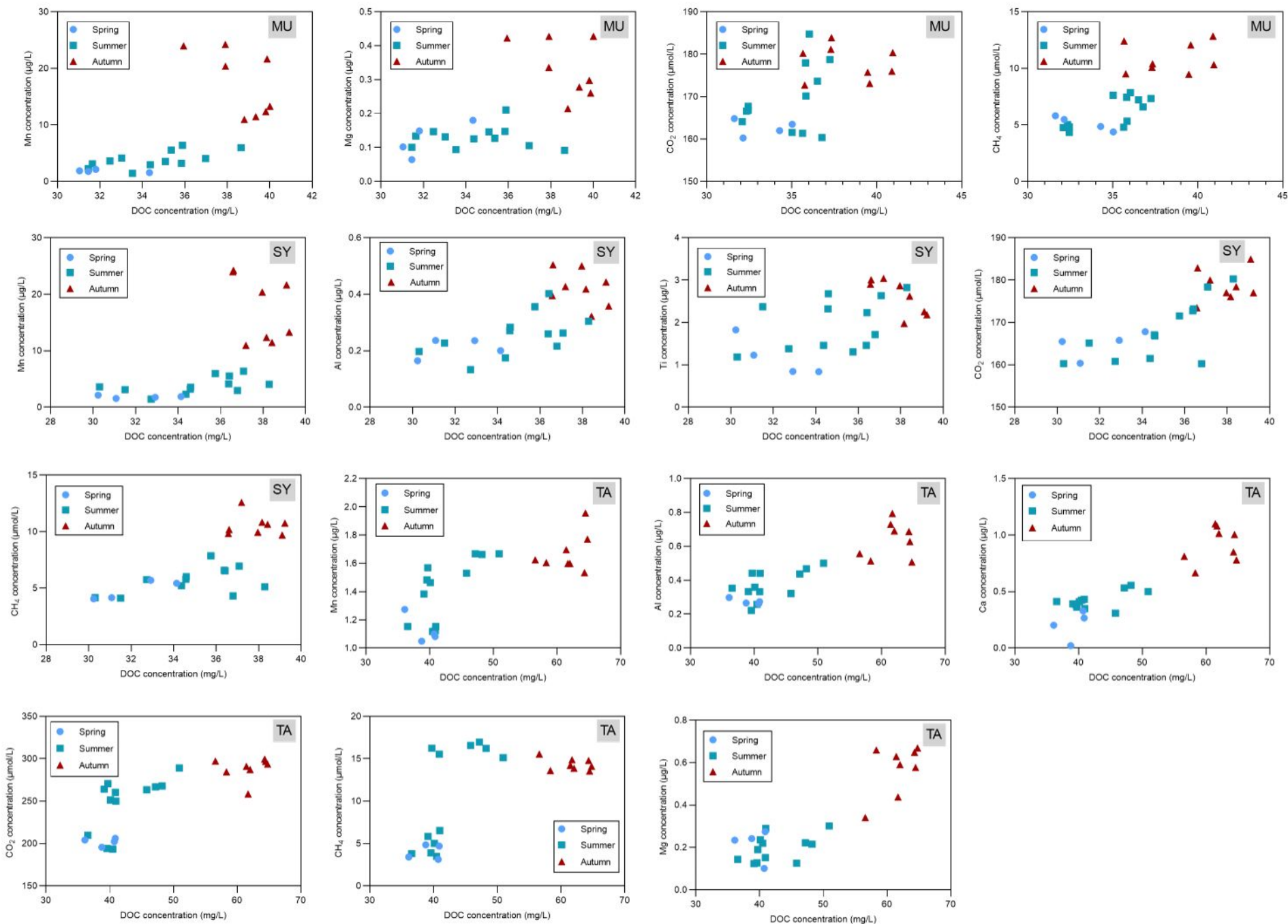


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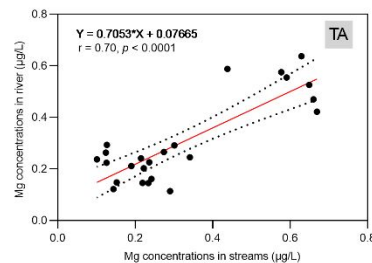
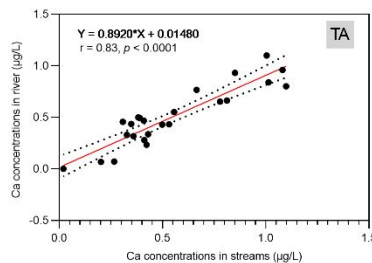
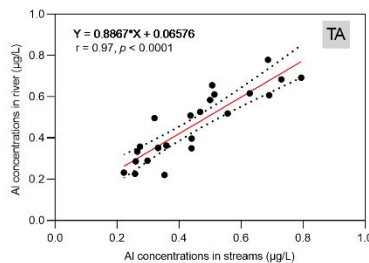
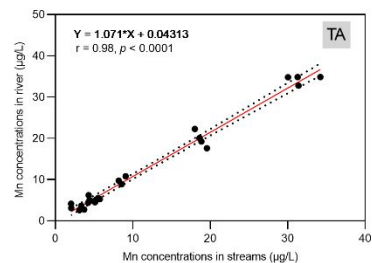
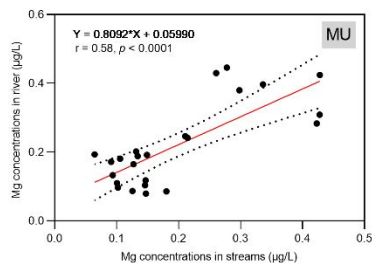
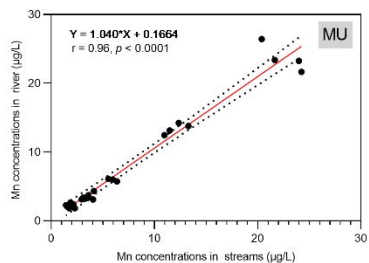
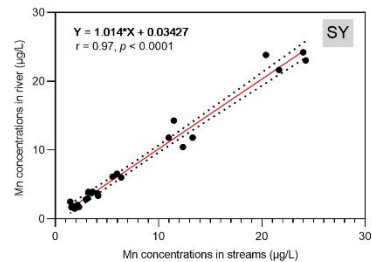
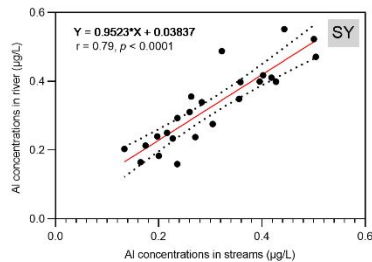
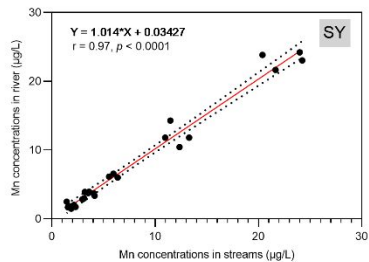


Figure 10.

