



# Mosaic structure of the fungal community in the Kislo-Sladkoe Lake that is detaching from the White Sea

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## Abstract

The major part of the north polar region is intensely rising by postglacial crustal movement. This process gives rise to the separation of different basins from seas and oceans, which affects a combination of freshwater and marine organisms. Gradually losing contact with the seas, many near-shore lakes of the Arctic are mostly desalted and form bogs. Fungi as decomposers play an important role in all ecosystems. However, the diversity and role of fungi in Arctic aquatic ecosystems is largely unknown. It is also not clear how the taxonomic structure of the fungal community is affected by the process of gradual desalinization and waterlogging. We investigated the diversity of filamentous culturable fungi in different parts of the brackish Kislo-Sladkoe Lake (White Sea, Russia). Annually, 42 samples of the bottom and coastal soils have been collected at the lake from which fungi were recovered on standard and selective media. Based on morphological and molecular markers, a total of 127 taxa have been identified. The fungal community appeared to be influenced by its sea origin and comprised both marine (*Paradendryphiella salina*, *Acremonium* spp.) and terrestrial soil species of *Penicillium*, *Talaromyces*, *Mucor*, *Umbelopsis*, *Cladosporium*, *Cadophora*, *Sistotrema*, *Helotiales*, *Pleosporales*, sphagnum moss destructors (*Oidiodendron* spp.) and insect-associated species of *Tolypocladium*. The results indicate that the composition of the fungal community in the rising polar White Sea region reflects the dynamics of global changes in physical–chemical parameters and animal and plant associations because of separation from the sea.

**Keywords** Fungal diversity · Brackish lake · Glacioisostatic movement · Coastal rising

## Introduction

Some Arctic oceanic and marine coastal zones are intensely rising because of glacio-isostatic movement (Krasnova et al. 2013). As a result, water bodies are detaching from seas and

oceans and transforming to meromictic lakes in northern parts of North America and Eurasia (Fig. 1; Dickman 1978; Ludlam 1996; Gibson et al. 2002; Hakala 2004; Lutz and Kaulfuß 2006; Van Hove et al. 2006; Pouliot et al. 2009; Strelkov et al. 2014; Gulati et al. 2017). The coastal line of the White Sea (Russia) is a prominent example of this natural phenomenon. The shoreline of the western part of the White Sea is flat and indented; hence, the consequence of the coastal rising is the separation of the water bodies such as bays, armlets, and small lakes from coastal straits. These lakes gradually lose their connection with the sea and transform into different types of coastal basins varying in depth. Lakes with a depth of more than 6 m transform into meromictic lakes, while those with a depth of 1–6 m transform into boggy fresh lakes. If the depth is less than 1 m, the lakes form marshes (Pantiulin and Krasnonva 2011).

The lakes that are separating from the White Sea combine both marine and continental features and represent a unique environmental niche. These basins are still connected to the sea in a varying degree, while also having inflow of

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**Fig. 1** The studied meromictic lakes in the Arctic region are marked by the circles; WSBS the White Sea biological station

fresh water from inland. Therefore, the main feature of these lakes is a strong vertical stratification governed by dynamics of fresh and salt water influx (Vinogradov et al. 2015). As a result, such detaching lakes harbour unique biota comprising a combination of freshwater and marine organisms (Krasnova et al. 2013). The shore-lines of these lakes are equally interesting biotopes. Horizontal nearshore currents distribute inflowing marine and fresh water along the shores. Therefore, coastal soil and littoral ground are composed of different zones with fluctuating environmental conditions, such as humidity, salinity and pH values. The mosaic coastal vegetation of these lakes highlights the soil patchiness (Sidneva 2008). Consequently, these transitional ecosystems represent extreme habitats and good systems for studying the general regularities in the formation of anaerobic conditions. These water bodies also represent unique objects for investigations of the biogeochemical processes that take place in water and sediments, and the interaction of freshwater and marine organisms. Scientists from various disciplines have explored the hydrology, flora and fauna of the water bodies at different stages of isolation from the White Sea (Krasnova

et al. 2013). However, the fungal diversity and the role of fungi in meromictic and brackish lakes are poorly studied, although some data are available on fungi from meromictic lakes of Canada, Japan, France and Germany (Takishitaa et al. 2007; McAndrews and Turton 2010; Oikonomou et al. 2015; Lepère et al. 2016). So far, there have been no studies on the fungal biodiversity in separating lakes in Russia. Yet, littoral zone and marine bottom sediments including polar marine waters are specific ecotopes for fungi, with respect to biodegradation, symbiotic relationships and other functions (Damare et al. 2006; Damare and Raghukumar 2008; Bubnova 2017; Furbino et al. 2017; Rämä et al. 2017).

In this study, we further explore the fungal biodiversity in the Kislo-Sladkoe Lake, which is detaching from Kandalaksha Bay of the White Sea. This lake is a well-investigated site as a model of separating water reservoirs located in this region. It is located near the Pertsov White Sea Biological Station (WSBS; 66°34'N, 33°08'E; Online Resource 1) of the Lomonosov Moscow State University and is at an initial stage of peatland formation because of its shallow waters (4.5 m maximum depth). Over recent

years, the Kislo-Sladkoe Lake has been examined with a wide range of approaches (Sidneva 2008; Krasnova et al. 2013, 2014, 2015; Malyshko et al. 2015; Vinogradov et al. 2015). We began to study the diversity of fungi in this lake from the year 2008. Presumably, soil from the coast is more habitable for fungi than bottom sediments; therefore, we mainly focused on the investigation of the lake's shore-line, whereas fungi from the bottom sediments were described in less detail. The fungi were isolated from the peat, soil and sediment samples taken from the coast, littoral and bottom of the lake in 2008–2010. Our isolation method recovered culturable basidiomycetous, ascomycetous and zygomycetous fungi, which were identified using morphological and cultural features as well as DNA sequence data (ITS rDNA region, along with LSU rDNA). To make the present work more complete, we combined our new data with selected data from our previous studies of microfungi in the lake (Grum-Grzhimaylo et al. 2016). We characterized the composition, abundance and spatial distribution of fungal species from different parts of the lake and related these data to the ecological peculiarities of the lake, such as its freshening and a contact with the Sea.

## Materials and methods

### Description of the Kislo-Sladkoe Lake

The Kislo-Sladkoe Lake ["Sour-Sweet" Lake; its other name is Polupresnoe ("Brackish") Lake] is located near the Karelian coast of Kandalaksha Bay of the White Sea 2 km east of the Pertsov White Sea Biological Station (Online Recourse 1). The climatic and geomorphological characteristics of this area have been given earlier (Grum-Grzhimaylo et al. 2016). The Kislo-Sladkoe Lake is oval-shaped, 100 m long, 60 m wide, occupying 1.6 ha, with an average depth of 1.5 m and a maximum depth of 4.5 m. The drainage area of the lake is 157.0 m<sup>2</sup>. In 2010, its elevation was 0.4 m above sea level (Schaporenko et al. 2005; Krasnova et al. 2014). Its basin is being formed as a result of separation of a water area between a small unnamed island covered with pines and the mainland shore of Kindo Cape (Fig. 2). In the past, the bed of the strait was bound from two sides by two submerged ridges, which rose to the surface because of the general rise of the land and formed isthmuses separating the lake area from the Great Salma Strait. One of the isthmuses is covered with grass and appears to be inundated only during seasonal snow melting when the lake is overfilled. The second isthmus consists of rolled boulders and pebble. A weak water exchange with Kandalaksha Bay takes place in the surface layer through this isthmus during high tides (once a month for 2–3 days). A weak water exchange is likely to exist through the entire northern shore of the lake. The

southern shore is swampy and very weak though permanent creek flows into the sea over a steep slope of the peninsula. Underwater springs, which freshen the lake water, can also exist (Schaporenko et al. 2005). The vertical structure of the lake is characterized by seasonal and interannual variations (Vinogradov et al. 2015). Water characteristics from each of the layers in 2010 are shown in the Online Recourse 2.

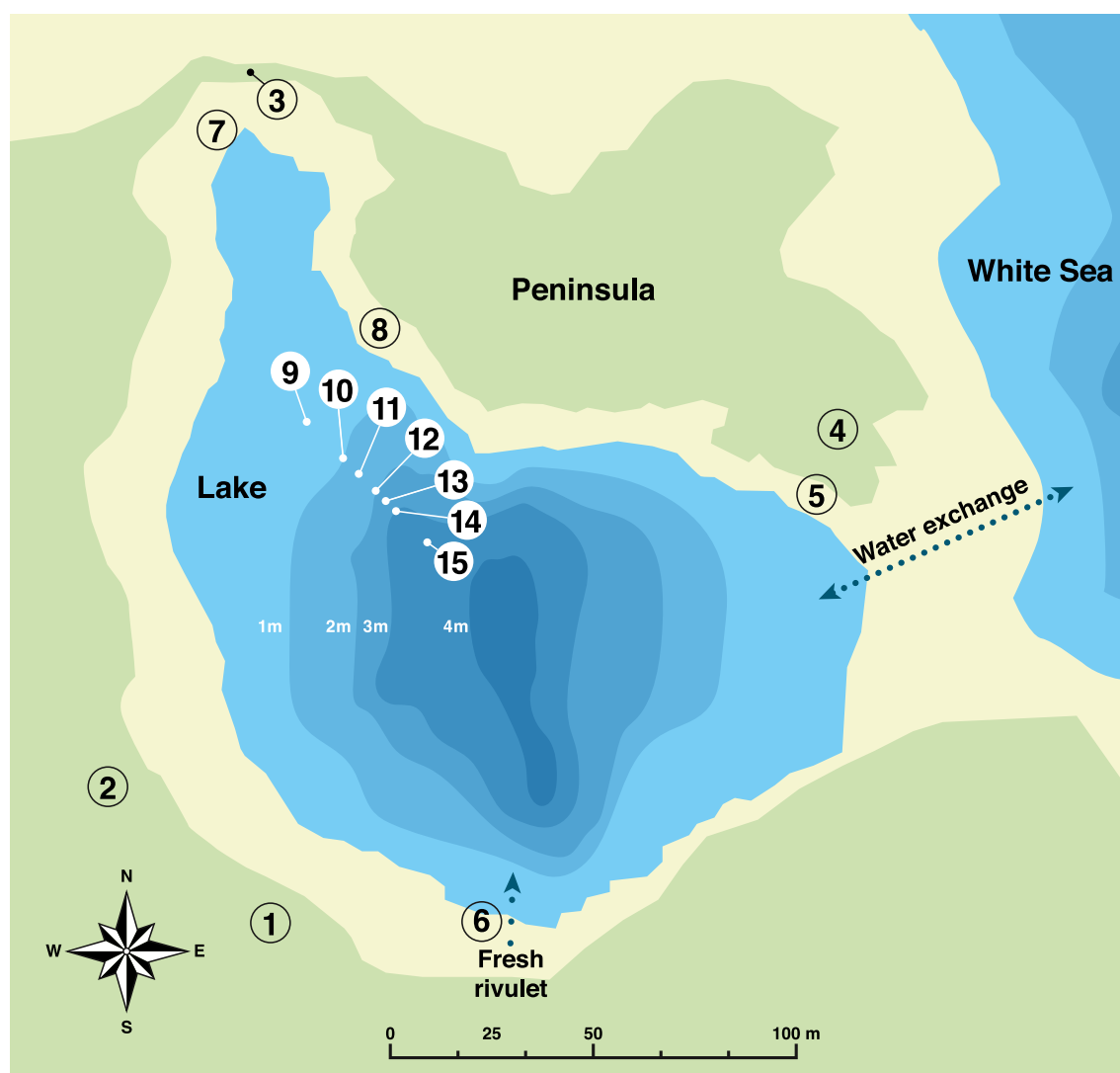
Coastal vegetation of the lake is a mosaic. Different plant communities including littoral, bog and forest vegetation constantly change one after another along the shore. The circular position of the primary sea-coast vegetation is indicative for the coastal rising (Sidneva 2008; Grum-Grzhimaylo 2013; Yatzenko et al. 2017).

### Sampling

The samples of soil, peat and bottom sediments were collected from several locations along the overgrown coast, littoral edge and bottom of the lake (Fig. 2; Tables 1, 2). Soil samples from the coast and littoral at the surface and 0.1 m depth, forest soil at the different horizons (two profile cuts), plants (living and dead parts of the moss *Sphagnum* spp.), and peat samples at 0.15 and 0.3 m depths were taken using a knife and transferred to sterile Petri dishes. The knife was cleaned with water and 96% alcohol following each extraction. The bottom sediment samples were collected every 0.5 m at different depths by SCUBA diving and placed in sterile tubes. All the samples were taken at one replication per season. Some locations were sampled twice (in different years). The collected samples were transported in a cold container to the laboratory and stored at 5 °C for 1–2 days until plating.

### Fungal isolation, cultivation, identification and data analysis

Dilutions were prepared from all the samples using 1 g in 10 mL of sterile distilled water, pounded in a mortar, and further diluted 100-fold. From this final dilution, 0.25 mL was pipetted onto two Petri plates each of six media: malt extract agar, water agar, citric-acid buffer agar, sphagnum extract agar, alkaline buffer agar and brackish water agar with the water from the lake. The media preparation and fungal cultivation were performed as described previously (Grum-Grzhimaylo et al. 2016). The identification of the fungi by morphologic and cultural characters was performed using manuals (Raper and Fennell 1965; Raper et al. 1968; Rifai 1969; von Arx 1981; Ellis 1971; Gams 1971, 2000; Barnett and Hunter 1972; Schipper 1978; Kohlmeyer and Kohlmeyer 1979; Carmichael et al. 1980; Bissett 1982, 1984; de Hoog et al. 2000; Schroers 2001; Zare and Gams 2001; Klich 2002; Samson et al. 2004; Crous et al. 2007; Domsch et al. 2007; Samson



**Fig. 2** Scheme of the Kisko-Sladkoe Lake, the sampling points are: 1 sphagnum peat, 2 boreal podzol, 3 muck land, 4 seashore grassed soil, 5–7 ooze, 8 coarse sand, 9–15 bottom sediments

and Houbraken 2011; Seifert et al. 2011). The morphological analysis was carried out with light microscopes (Mikmed-2, Carl Zeiss Axioskop 40 FL, Leica DM 2500). The information regarding fungal classification generally follows the Index Fungorum (<http://www.indexfungorum.org>) databases. The fungal strains which failed to sporulate following incubation (for 30 days) were considered sterile mycelia. A substantial part of the sterile mycelia and sporulating strains that were hard to identify by morphology were identified by DNA sequence analysis of the ITS and LSU regions of the nuclear-encoded rDNA. The same analysis was conducted for dominant fungal species in order to verify identification by morphological characters. Total genomic DNA (gDNA) was extracted from the mycelium of the 109 fungal strains using the chloroform/isopropanol method (Karakousis et al. 2006; Grum-Grzhimaylo et al.

2016). We amplified and sequenced the ITS rDNA for all strains and LSU rDNA for the most of them using common primer sets. The primer sets, thermocycling programs and sequencing procedures were performed as described previously (Grum-Grzhimaylo et al. 2016). Sequences were compared with the data from GenBank using BLAST similarity searches. Newly generated sequences were deposited in the GenBank. The GenBank Accession Numbers of 36 species from the lake are in our previous work (Grum-Grzhimaylo et al. 2016), and the others given in the Online Resource 3.

Data on the total number of fungi were expressed as colony-forming units (CFU)  $\text{g}^{-1}$  of dry peat, as described previously (Grum-Grzhimaylo et al. 2016). The Sørensen index was used to compare fungal communities between the different parts of the lake (Zak and Willig 2004).

**Table 1** Characteristics of the samples of the different parts of the Kislo-Sladkoe Lake shore

Characteristics	Soil from the coast		Soil from the littoral zone						
No. of sampling place (see Fig. 2)	1	2	3	4	5	6	7	8	
Date of sampling	July 2008, July 2010	July 2008	July 2009, July 2010	July 2009, July 2010	July 2009, July 2010	July 2010	July 2010	July 2010	July 2010
Location (compass point/depth, m)	SSW	W	N	NE	ENE	S	N	NNE	
Main vegetation	<i>Sphagnum</i> spp.	<i>Pinus sylvestris</i> , <i>Betula pubescens</i>	<i>Juncus atrofuscus</i> , <i>Triglochin maritima</i> , <i>Blysmus rufus</i> , <i>Eleocharis uniglumis</i>	–	–	<i>Carex</i> spp.	–	–	
Type of ground	Peat	Boreal podzol	Muck land	Seashore soil	Ooze				Coarse sand
Sample (horizon/ type/depth, m)	LS DS 0.2	L F+H B 0.3	S 0.1	S 0.1	S 0.1	S 0.1	S 0.1	S 0.1	S 0.1
pH of samples	6.0	5.0	6.5	5.0	8.5	8.0	8.5	8.0	8.5

LS living part of *Sphagnum* spp., DS dead part of *Sphagnum* spp., L dead soil cover, ground litter, F + H fermentation and humification horizons, B illuvial horizon with surface gleying, C parent rock material (sand), S surface

**Table 2** Characteristics of the samples of the different parts of the Kislo-Sladkoe Lake bottom sediments

No. of sampling place (see Fig. 2)	9	10	11	12	13	14	15
Date of sampling	July 2010						
Depth, m	0.5	1.0	1.5	2.0	2.5	3.0	3.5
Type of ground	Oozy bottom						
Sample (horizon)	Surface						
pH of samples	6.0	6.0	5.5	5.0	6.0	6.0	5.5

## Results

In total, 313 fungal isolates were obtained from the 42 samples of different types of soils around the lake and its bottom sediments. These isolates belonged to 127 taxa, representing 40 genera and 28 sterile unidentified morphotypes (Table 3). Of these taxa, 111 (71.2%) were Ascomycota, 8 (5.1%) Basidiomycota, and 8 (5.1%) Zygomycota. The number of viable fungal propagules ranged from  $10^2$  to  $10^5$  CFU g<sup>-1</sup> of the sample dry weight. Taxonomic identification of all taxa was based on morphological characters, while 40 of them (109 isolates) were identified using molecular markers (ITS and LSU rDNA), including the majority of the sterile mycelia (86 of 114 isolates), together with 13 species which were impossible to identify by morphology and a few frequent species (10 taxa).

The dominance of the fungi from the Ascomycota division was a common characteristic of the mycobiota from all parts of the lake due to abundance of the anamorphic species. The predominant genera were *Penicillium* (mostly *Monoverticillata*) and *Talaromyces* (Table 3).

The following genera were widely represented in all parts of the lake: *Acremonium*, *Cladosporium*, *Cadophora*, *Trichoderma*, *Lecanicillium*, *Mucor*, *Umbelopsis*, *Aspergillus*, *Tolypocladium*, *Cordyceps* (was presented by anamorphic stages from the genera *Beauveria* и *Parengyodontium*), *Talaromyces* and *Penicillium*.

In the sphagnum peat (Figs. 2 point 1, 4), the following species dominated: *Acremonium* spp., *Cadophora luteo-olivacea*, *Hypocreales* sp., *Penicillium montanense*, *Metapochonia bulbillosa*, *Tolypocladium cylindrosporum*, *Cladosporium antarcticum*, *C. allicinum*, *C. cladosporioides*, and *C. herbarum*. *Cadophora luteo-olivacea*, *Talaromyces funiculosus* and *Tolypocladium inflatum* were frequently found in grassed soil (Figs. 2, points 3, 4, 4); *Penicillium glabrum*, *P. montanense*, *Trichoderma polysporum* and *Umbelopsis ramanniana* in the forest soil (Figs. 2, point 2, 4). *Acremonium* spp., *Antrodia* sp., *Paradendryphiella salina*, *Sistotrema brinkmannii*, *Penicillium aurantiogriseum*, *P. funiculosum*, *P. glabrum*, *P. thomii*, *Penicillium* sp., *Tolypocladium cylindrosporum*, *T. inflatum*, *Trichoderma harzianum* and *T. viride* predominated in the littoral soils (Figs. 2, points 5–8, 3), and *Cladosporium cladosporioides* in the benthic ooze

(Figs. 2, points 9–15, 3). The sterile mycelium morphotypes were often isolated from all components of the lake, except for the forest and silted soils (Fig. 2, points 2, 3, 6, 7).

The greatest level of similarity in the fungal species composition was identified for the mycobiota of littoral zone and the lake benthic ooze, while the lowest level was found between the ooze and the sphagnum peat. The similarity of the fungal communities between the different parts of the lake ranged from 0.09 (between the sphagnum peat and the bottom sediments) to 0.23 (between the littoral zone and the bottom sediments) (Table 4).

In the 24 samples taken from different points along the lakeshore that were not flooded during high tides (Fig. 2, points 1–4; Table 1), 88 fungal morphotypes were distinguished. The largest number of fungal taxa was found in samples of the living part of the sphagnum moss and sphagnum peat (Fig. 2, point 1) and grass-overgrown soil (Fig. 2, point 4). *Penicillium montanense* was identified in all layers of the sphagnum peat deposit, and all horizons of the forest and grass-overgrown soil. *Penicillium glabrum* and *Trichoderma polysporum* were also isolated from all the horizons of the forest soil (Fig. 2, point 2). *Cladosporium antarcticum*, *C. allicinum*, *C. herbarum*, *Tolypocladium cylindrosporum* and *Acremonium* spp. were recovered from the peat-bog layers (Fig. 2, point 1). *Talaromyces funiculosus* and *Tolypocladium inflatum* were found to be common in the grass-overgrown soil (Fig. 2, point 4), and *Acremonium* sp. 2 and *Plectosphaerella* sp. in the silted soil (Fig. 2, point 3). *Cladosporium*, *Cordyceps* (*Beauveria* spp., *Parengyodontium album*), *Tolypocladium*, *Penicillium*, *Umbelopsis*, *Cadophora* and sterile mycelium were the most widely represented fungi in the sphagnum peat (Fig. 2, point 1). In the grass overgrown soil (Fig. 2, point 4), *Acremonium*, *Cadophora*, *Cordyceps* and *Penicillium* were found with high frequency, while *Penicillium* was maximally represented in the forest soil (Fig. 2, point 2), and *Acremonium*, *Fusarium* and *Trichoderma* in the silted soil (Fig. 2, point 3). Two species of *Oidiodendron* were found only in the sphagnum peat (Fig. 2, point 1; Table 3).

The littoral zone of the lake (Fig. 2, points 5–8) is characterized by different soil properties and plant communities, which vary depending on the distance to the sea or freshwater basins. Eleven soil samples from the littoral zone



**Table 3** Isolated species with total number (CFU  $\times 10^2$  g<sup>-1</sup> dry soil) and media of isolation

pH of the samples	No. of sampling places (see Fig. 2)	Soil from the coast					Soil from the littoral zone					Bottom sediments				Media
		5.0–6.0	6.5	6.5–8.5	5.0	8.0–8.5	7.0–8.0	8.5	8.0	5.5–6.0	5.0–6.0	9	10	11		
		1	2	3	4	5	6	7	8	8.0	9, 10	11–13	14, 15			
Taxa																
<i>Acremonium</i> cf. <i>charticola</i>		–	–	5	–	–	–	–	–	–	–	–	–	WA		
<i>Acremonium</i> cf. <i>fuci</i>		–	–	–	–	–	–	–	5	–	–	–	–	SWA		
<i>Acremonium</i> sp. 1		10	5	–	10	10	20	–	–	–	800	–	–	WA, AA, SWA, MA		
<i>Acremonium</i> sp. 2		–	–	20	–	20	–	50	5	–	–	–	–	AA, MA, SWA		
<i>Acremonium</i> sp. 3		30	–	–	5	15	–	–	–	–	–	–	–	MA, AA		
<i>Acremonium</i> sp. 4		–	–	–	–	5	–	–	–	–	–	–	–	MA		
<i>Acremonium</i> sp. 5		–	–	10	–	50	10	–	–	–	–	–	–	SWA, MA		
<i>Acremonium</i> sp. 6		–	–	–	–	28	10	–	5	–	–	–	–	AA		
<i>Alternaria alternata</i> (Fr.) Keissl.		15	–	–	–	–	–	–	–	–	–	–	–	CA		
<i>Alternaria</i> sp. 1		–	–	–	–	–	–	–	5	–	–	–	–	MA		
<i>Alternaria</i> sp. 2		5	–	–	10	–	–	–	–	–	–	–	–	CA, MA		
<i>Antrodia</i> sp.		–	–	–	–	15	–	–	–	–	–	–	–	CA		
<i>Ascomycota</i> sp. 1		35	–	–	–	–	–	–	–	–	–	–	–	CA		
<i>Ascomycota</i> sp. 2		–	–	–	–	–	–	–	5	–	–	–	–	MA		
<i>Ascomycota</i> sp. 3		–	–	–	5	–	–	–	–	–	–	–	–	MA		
<i>Ascomycota</i> sp. 4		–	–	–	–	–	–	–	–	–	–	–	20	WA		
<i>Ascomycota</i> sp. 5		–	–	–	–	–	50	–	–	–	–	–	–	SWA		
<i>Aspergillus proliferans</i> G. Sm.		–	–	–	–	–	45	–	–	–	–	–	–	SEA		
<b><i>Aspergillus tubingensis</i> Mosseray</b>		–	–	–	–	–	–	15	–	–	–	–	65	MA		
<i>Aspergillus ustus</i> (Bainier) Thom and Church		–	–	–	5	–	10	–	–	–	–	–	–	MA		
<i>Aspergillus versicolor</i> (Vuill.) Tirab.	10	–	–	–	–	–	10	–	–	–	–	–	–	CA		
<b><i>Aspergillus</i> sp.</b>	–	–	–	5	15	–	–	–	–	–	–	–	–	MA, SEA		
<i>Aureobasidium pullulans</i> (de Bary and Löwenthal) G. Arnaud	–	–	5	–	–	–	–	–	–	–	–	–	–	MA		
<i>Basidiomycota</i> sp.	–	–	–	–	–	–	–	–	–	–	–	–	–	MA		
<i>Beauveria bassiana</i> (Bals.-Criv.) Vuill.	15	–	–	–	–	–	10	–	–	–	–	–	–	AA, CA, MA, SEA, WA		
<i>Beauveria brongniartii</i> (Sacc.) Petch	21	–	–	–	–	–	–	–	–	–	–	–	–	MA		
<i>Beauveria</i> sp.	–	–	–	–	14	–	–	–	–	–	–	–	–	MA		
<b><i>Botryotinia</i> sp.<sup>a</sup></b>	550	–	–	–	–	–	–	–	–	–	–	–	–	WA		
<i>Botrytis cinerea</i> Pers.	50	–	–	–	–	–	–	–	–	–	–	–	–	CA		
<b><i>Cadophora fastigiata</i> Lagerb. and VKM F-4772<sup>a,b</sup></b>	5	–	–	–	5	–	–	–	–	–	–	–	–	AA, CA, SWA		
<b><i>Cadophora luteo-olivacea</i> (J.F.H. Beyma) T.C. Harr. and McNew VKM F-4773<sup>b</sup></b>	250	–	10	15	–	–	–	–	–	–	10	–	–	CA, SEA, WA		

Table 3 (continued)

pH of the samples	Soil from the coast										Soil from the littoral zone					Bottom sediments			
	5.0–6.0	6.5	6.5–8.5	5.0	8.0–8.5	7.0–8.0	8.5	8.0	5.5–6.0	5.0–6.0	9	10	11						
No. of sampling places (see Fig. 2)	1	2	3	4	5	6	7	8	9, 10	11–13	14, 15								
<i>Cadophora malorum</i> (Kidd and Beaumont) W. Gams	–	–	–	–	–	–	–	5	–	–	–	–	–	CA					
<i>Cadophora melinii</i> Nannf.	30	100	–	5	–	–	–	–	–	–	–	–	–	MA, SEA, WA					
<i>Cadophora</i> sp. 1	25	–	–	–	–	–	–	–	–	–	–	–	–	SEA					
<i>Cadophora</i> sp. 2	–	–	–	–	–	–	–	5	–	–	–	–	–	CA					
<i>Chaetosphaeria</i> sp.	–	–	–	–	–	10	–	–	–	–	–	–	–	CA					
<i>Cladosporium allicinum</i> (Fr.) Bensch, U. Braun and Crous	400	–	–	5	–	–	–	–	–	–	–	–	–	AA, CA, MA, SWA					
<i>Cladosporium antarcticum</i> K. Schub., Crous and U. Braun 2007	200	–	–	5	–	–	–	–	–	–	–	–	–	AA					
<i>Cladosporium</i> <i>Cladosporium cladosporioides</i> (Fresen.) G.A. de Vries <sup>a</sup>	650	–	–	–	–	10	–	–	15	–	–	–	20	MA, SEA, SWA					
<i>Cladosporium herbarum</i> (Pers.) Link <sup>a</sup>	60	–	–	–	–	20	–	–	–	–	–	–	–	MA, SWA					
<i>Cladosporium oxysporum</i> Berk. and M.A. Curtis	200	–	–	–	–	–	–	–	–	–	–	–	–	MA, SEA					
<i>Cladosporium</i> sp. 1	–	–	–	–	10	–	–	–	–	–	–	–	–	AA					
<i>Cladosporium</i> sp. 2	10	–	–	–	–	–	–	–	–	–	–	–	–	SEA					
<i>Cladosporium</i> sp. 3	–	–	5	–	–	–	–	–	–	–	–	–	–	CA					
<i>Cladosporium sphaerospermum</i> Penz.	10	30	–	–	–	–	–	–	–	–	–	–	–	SEA					
<i>Coniochaeta</i> sp.	–	5	–	–	–	–	–	–	–	–	–	–	–	CA					
<i>Cordyceps militaris</i> (L.) Fr.	–	–	–	–	–	10	–	–	–	–	–	–	–	AA					
<i>Cylindrobasidium laeve</i> (Pers.) Chamuris	–	–	–	5	–	–	–	–	–	–	–	–	–	WA					
<i>Dothidea</i> sp. <sup>a</sup>	–	–	–	–	–	–	–	–	–	–	–	5	–	MA					
<i>Dothideomycetes</i> sp. 1 <sup>a</sup>	–	–	–	–	–	–	–	–	–	–	–	–	65	SWA					
<i>Dothideomycetes</i> sp. 2 <sup>a</sup>	–	–	–	–	–	–	–	–	–	–	–	–	–	MA					
<i>Fusarium avenaceum</i> (Fr.) Sacc.	10	–	10	–	–	–	–	–	–	–	–	–	–	MA					
<i>Fusarium graminearum</i> Schwabe	–	–	10	–	–	–	–	–	–	–	–	–	–	CA					
<i>Fusicladium pini</i> Crous and de Hoog	10	–	–	–	–	–	–	–	–	–	–	–	–	SEA					
<i>Helotiaceae</i> sp. <sup>a</sup>	10	–	–	–	–	–	–	–	–	–	–	–	–	WA					
<i>Helotiales</i> sp. 1 <sup>a</sup>	35	–	–	–	–	–	–	–	–	–	–	–	–	CA, WA					
<i>Helotiales</i> sp. 2 <sup>a</sup>	25	–	–	–	–	–	–	–	–	–	–	–	–	CA					
<i>Helotiales</i> sp. 3	–	–	–	15	–	–	–	–	–	–	–	–	–	MA					
<i>Helotiales</i> sp. 4 <sup>a</sup>	–	–	5	–	–	–	–	–	–	–	–	–	–	SWA					
<i>Helotiales</i> sp. 5 <sup>a</sup>	–	–	–	–	–	–	–	–	–	–	–	10	–	SEA					
<i>Helotiales</i> sp. 6	–	–	–	–	–	–	–	–	–	–	–	5	–	CA					
<i>Lecanicillium evansii</i> Zare and W. Gams	–	–	–	–	15	–	–	–	–	–	–	–	–	CA					
<i>Lecanicillium</i> sp. 1	10	–	–	–	–	–	–	–	–	–	–	–	–	AA					
<i>Lecanicillium</i> sp. 2	35	–	–	–	–	–	–	–	–	–	–	–	–	SWA					



Table 3 (continued)

	Soil from the coast					Soil from the littoral zone							Bottom sediments			
	5.0–6.0	6.5	6.5–8.5	5.0	15	8.0–8.5	7.0–8.0	8.5	8.0	5.5–6.0	5.0–6.0	5.5–6.0	9	10	11	
pH of the samples	1	2	3	4	5	6	7	8	8.0	9, 10	11–13	14, 15				
No. of sampling places (see Fig. 2)																
<i>Lecanicillium</i> sp. 3	–	–	–	–	15	–	–	–	–	–	–	–	–	–	–	SEA
<b>Massarinaceae</b> sp. <sup>a</sup>	25	–	–	–	–	–	–	–	–	–	–	–	–	–	–	CA
<i>Memnoniella dichroa</i> (Grove) L. Lombard and Crous	30	–	–	–	–	–	–	–	–	–	–	–	–	–	–	SEA
<i>Metapochonia bulbilosa</i> (W. Gams and Malla) Kepler, Rehner and Humber VKM F-4674 <sup>a,b</sup>	30	–	–	–	–	–	–	–	–	–	–	–	–	–	–	CA
<i>Mortierella elongata</i> Linnem. <sup>a</sup>	–	–	–	–	–	–	–	–	–	–	200	–	–	–	–	MA
<i>Mucor circinelloides</i> Tiegh.	–	5	–	–	–	–	–	–	–	–	–	–	–	–	–	CA, MA
<i>Mucor hiemalis</i> Wehmer VKM F-4774 <sup>b</sup>	–	25	–	15	15	–	–	–	–	–	–	–	–	–	–	MA, SEA, WA
<i>Mucor racemosus</i> Fresen.	–	–	5	–	–	–	–	–	–	–	–	–	–	–	–	WA
<b>Nectriaceae</b> sp.	15	–	–	–	–	10	–	–	–	–	–	–	–	–	–	MA
<i>Oidiodendron ambiguum</i> Peyronel and Malan	25	–	–	–	–	–	–	–	–	–	–	–	–	–	–	MA
<i>Oidiodendron griseum</i> Robak	10	–	–	–	–	–	–	–	–	–	–	–	–	–	–	MA
<b>Paradendryphiella salina</b> (G.K. Sutherland) Woudenberg and Crous	–	–	5	–	–	–	–	5	–	–	–	–	–	–	–	MA
<i>Paragynodontium album</i> (Limber) C.C. Tsang, J.F.W. Chan, W.M. Pong, J.H.K. Chen, A.H.Y. Ngan, Cheung, C.K.C. Lai, D.N.C. Tsang, S.K.P. Lau, P.C.Y. Woo	30	–	–	–	15	–	–	–	–	–	–	–	–	–	–	CA, MA, SEA
<i>Penicillium aurantiogriseum</i> Dierckx	–	–	5	–	–	–	–	–	–	–	–	–	–	–	–	SEA
<i>Penicillium decumbens</i> Thom	–	–	–	–	–	35	–	–	–	–	–	–	–	–	–	CA
<i>Penicillium dierckxii</i> Biourge	10	–	–	–	–	–	–	–	–	–	–	–	–	–	–	WA
<i>Penicillium glabrum</i> (Wehmer) Westling	10	750	–	–	–	20	–	5	–	–	–	–	–	–	–	CA, MA, SEA, WA
<i>Penicillium griseofuuum</i> Dierckx	–	–	–	–	–	–	–	–	–	–	10	–	–	–	–	CA
<b>Penicillium montanense</b> M. Chr. and Backus	100	250	–	15	–	10	–	–	–	–	–	–	–	–	–	CA, MA, SEA, SWA, WA
<i>Penicillium multicolor</i> Grig.-Man. and Porad.	–	–	–	–	–	–	–	–	15	–	–	–	–	–	–	WA
<i>Penicillium restrictum</i> J.C. Gilman and E.V. Abbott	–	50	–	–	–	–	–	–	–	–	–	–	–	–	–	WA
<i>Penicillium roseopurpureum</i> Dierckx	–	–	–	–	–	–	–	–	–	5	–	–	–	–	–	WA
<b>Penicillium thomii</b> Maire VKM F-4664 <sup>b</sup>	–	20	–	5	15	–	–	–	–	–	200	–	–	–	–	CA, MA
<i>Penicillium waksmanii</i> K.M. Zaleski	15	10	–	–	–	–	–	–	–	–	–	–	–	–	–	MA, SEA
<i>Penicillium</i> sp. 1	–	–	–	–	–	10	–	–	–	–	–	–	–	–	–	SWA
<i>Penicillium</i> sp. 2	–	–	–	–	–	20	–	–	–	–	–	–	–	–	–	CA, MA, SWA
<i>Penicillium</i> sp. 3	–	–	5	–	–	–	–	–	–	–	–	–	–	–	–	SWA
<i>Penicillium</i> sp. 4	–	–	–	–	–	–	–	10	–	–	–	–	–	–	–	CA, MA, SWA
<i>Peniophora</i> sp.	–	–	–	–	–	–	–	–	–	–	5	–	–	–	–	WA

Table 3 (continued)

	Soil from the coast					Soil from the littoral zone										Bottom sediments			
	5.0–6.0	6.5	6.5–8.5	5.0	8.0–8.5	7.0–8.0	8.5	8.0	5.5–6.0	5.0–6.0	5.5–6.0	5.0–6.0	5.5–6.0	5.0–6.0	5.5–6.0	9	10	11	
pH of the samples	1	2	3	4	5	6	7	8	9	10	11–13	14,15							
No. of sampling places (see Fig. 2)																			
<i>Phaeoacremonium</i> sp.	375	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	SEA
<i>Phialophora</i> sp.	–	–	–	5	–	–	–	–	–	–	–	–	–	–	–	–	–	–	SWA
<i>Phialophora verrucosa</i> Medlar	15	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	CA
<i>Plectosphaerella</i> sp.	–	–	20	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	AA
<i>Pleosporeles</i> sp. 1	–	–	–	–	–	–	–	5	–	–	–	–	–	–	–	–	–	–	MA
<i>Pleosporeles</i> sp. 2	–	–	10	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	MA
<i>Pleosporeles</i> sp. 3	–	–	–	–	–	–	–	5	–	–	–	–	–	–	–	–	–	–	MA
<i>Pseudogymnoascus pannorum</i> (Link) Minnis and D.L. Lindner	35	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	SEA
<i>Sarocladium implicatum</i> (J.C. Gilman and E.V. Abbott) Giraldo, Gené and Guarro	–	–	–	–	–	10	–	–	–	–	–	–	–	–	–	–	–	–	WA
<i>Scopulariopsis brumptii</i> Salv.-Duval	–	–	–	–	–	–	5	–	–	–	–	–	–	–	–	–	–	–	MA
<i>Sistotrema brinkmannii</i> (Bres.) J. Erikss. VKM F-4640 <sup>b</sup>	–	–	10	15	30	–	–	5	–	–	20	20	–	–	–	–	–	–	AA, CA, MA, SWA, WA
<i>Sporothrix</i> sp.	–	–	–	–	–	–	10	–	–	–	–	–	–	–	–	–	–	–	CA
<i>Stachybotrys echinatus</i> (Rivolta) G. Sm.	–	500	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	SEA, WA
<i>Sydowia polyspora</i> (Bref. and Tavel) E. Müll. <sup>a</sup>	275	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	CA, SEA
<i>Talaromyces aculeatus</i> (Raper and Fennell) Samson, N. Yilmaz, Frisvad and Seifert	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	MA
<i>Talaromyces funiculosus</i> (Thom) Samson, N. Yilmaz, Frisvad and Seifert	–	–	–	250	30	20	10	5	–	–	–	–	–	–	–	–	–	–	CA, MA, SEA, WA
<i>Talaromyces variabilis</i> (Sopp) Samson, N. Yilmaz, Frisvad and Seifert	–	20	–	30	–	–	–	–	–	–	200	–	–	–	–	–	–	–	CA, SEA
<i>Talaromyces verruculosus</i> (Peyronel) Samson, N. Yilmaz, Frisvad and Seifert	–	10	–	–	15	–	–	–	–	–	5	–	–	–	–	–	–	–	CA, SWA
<i>Tolypocladium cylindrosporum</i> W. Gams VKM F-4627 <sup>b</sup>	25	–	–	–	–	10	10	5	–	–	–	–	–	–	–	–	–	–	AA, MA, SEA
<i>Tolypocladium inflatum</i> W. Gams VKM F-4675 <sup>b</sup>	5	60	5	15	5	–	5	5	–	–	200	–	–	–	–	–	–	–	AA, MA, WA
<i>Tolypocladium nubicola</i> Bissett	5	–	–	5	–	–	10	–	–	–	–	–	–	–	–	–	–	–	MA, SWA, WA
<i>Trichoderma asperellum</i> Samuels, Lieckf. and Nirenberg	–	–	–	–	–	10	–	–	–	–	–	–	–	–	–	–	–	–	WA
<i>Trichoderma harzianum</i> Rifai	50	–	–	–	10	–	10	–	–	–	–	–	–	–	–	–	–	–	CA, SEA, SWA, WA
<i>Trichoderma koningii</i> Oudem.	5	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	CA
<i>Trichoderma parapiluliferum</i> (B.S. Lu, Druzhin. and Samuels) Jaklitsch and Voglmayr	–	5	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	MA
<i>Trichoderma polysporum</i> (Link) Rifai VKM F-4630 <sup>b</sup>	–	–	5	–	–	–	–	–	–	–	–	–	–	–	–	–	–	–	CA, WA
<i>Trichoderma viride</i> Pers.	–	–	5	–	–	10	–	–	–	–	–	–	–	–	–	–	–	–	SWA
<i>Trichoderma</i> sp.	–	–	–	–	–	10	–	–	–	–	–	–	–	–	–	–	–	–	CA

**Table 3** (continued)

	Soil from the coast					Soil from the littoral zone					Bottom sediments		
											9	10	11
	5.0–6.0	6.5	6.5–8.5	5.0	8.0–8.5	7.0–8.0	8.5	8.0	5.5–6.0	5.0–6.0	5.5–6.0	5.0–6.0	5.5–6.0
pH of the samples	1	2	3	4	5	6	7	8	9, 10	11–13	14, 15		
No. of sampling places (see Fig. 2)													
<i>Umbelopsis isabellina</i> (Oudem.) W. Gams	875	15	–	–	–	–	–	–	–	–	–	–	CA
<i>Umbelopsis ramanniana</i> (Möller) W. Gams	25	40	–	–	–	–	–	–	–	–	–	–	CA, WA
<i>Umbelopsis vinacea</i> (Dixon-Stew.) Arx	10	–	–	–	–	–	–	–	–	–	–	–	CA
<i>Umbelopsis</i> sp.	65	–	–	–	–	–	–	–	–	–	–	–	SWA
<i>Xylabobolus</i> sp.	–	–	–	–	–	–	–	–	–	–	–	–	AA, SEA
No of sterile isolates (28 morphotypes)	3	–	–	5	4	2	1	3	5	2			AA, CA, MA, SEA, SWA, WA

Species identified by morphological and molecular approach are in bold

MA malt extract agar, CA citric agar, AA alkaline agar, WA water agar, SEA sphagnum extract agar, SWA semi-saltwater agar

<sup>a</sup>GenBank Accession Numbers of these species are in our previous work (Grum-Grzhimaylo et al. 2016)

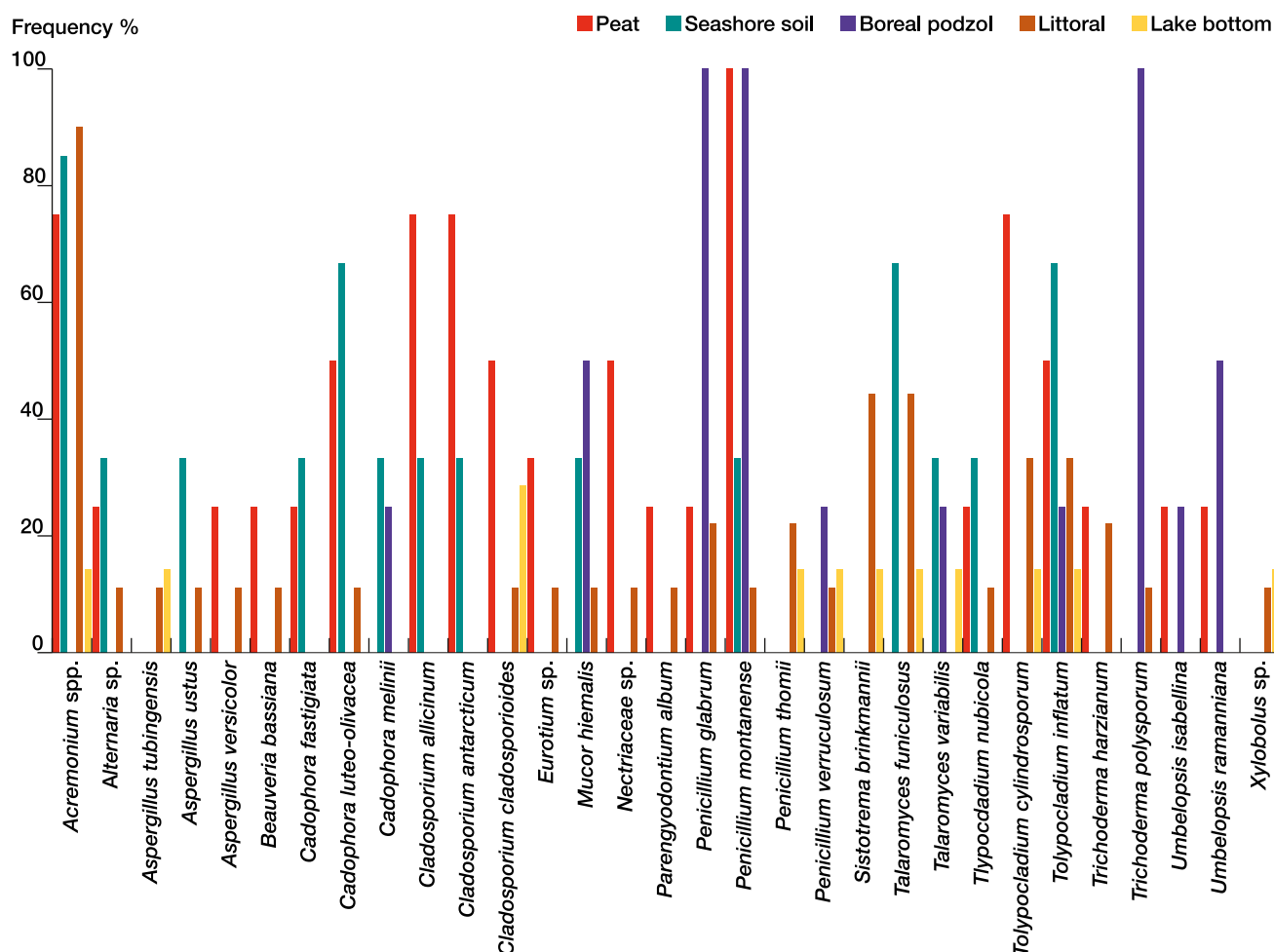
<sup>b</sup>Species deposited to All-Russian Collection of Microorganisms—VKM (the numbers are near the species names)

of the lake yielded 63 morphotypes of fungi. Samples that were collected from the surface yielded about 3–4 times more fungal taxa than in samples collected at a depth. The maximum number of fungal taxa (21) was observed in a sample of surface silt taken close to the river confluence area (Fig. 2, point 6). In general, the soils at the lake always contained *Penicillium* spp. and sterile mycelia. Genera such as *Acremonium*, *Tolypocladium*, *Trichoderma*, *Aspergillus* and *Cladosporium* were also widely represented (Table 3). The analysis of the spatial frequency of occurrence showed that the predominant fungi in the soil of the Kislo-Sladkoe Lake littoral were *Acremonium* spp., *Talaromyces funiculosus*, *Sistotrema brinkmannii*, *Tolypocladium cylindrosporum*, *T. inflatum*, *Paradendryphiella salina*, *Penicillium glabrum*, *P. thomii*, and *Trichoderma viride* (Fig. 4).

In total, 33 fungal taxa were identified in the samples collected from the lake's bottom sediment (Fig. 2, points 9–15). The number of fungi from the bottom sediment in various samples varied from 4 to 7. The lake benthic ooze harbored *Cladosporium* spp., *Talaromyces* spp., *Penicillium* spp., *Tolypocladium* spp., and *Sistotrema* spp., while the remaining genera were represented by a single species, and the sterile mycelia by 10 morphotypes (Table 3).

## Discussion

The Kislo-Sladkoe Lake is an interesting system to study, as it shares features of both fresh- and seawater environments. We hypothesized that such a transitional ecosystem would contain a unique composition of fungal biodiversity. Earlier studies on typical terrestrial peatlands in the same area showed a drastically different set of fungi (Grum-Grzhimaylo and Bilanenko 2012; Bilanenko and Grum-Grzhimaylo 2016; Grum-Grzhimaylo et al. 2016). Thus, our current study highlights the unusual hydrology of the lake, which in turn has resulted in the establishment of a specific fungal community. This could be explained by the distinctive physical, chemical and hydrological conditions, characteristic to the lake, which possesses a direct connection to the sea and, as a consequence, contains coastal vegetation, algoflora, microorganisms and animals different from the continental water basins that serve as a source of food for the fungi. At the level of higher rank taxa, such as divisions and classes, the fungal communities in the peatlands of the Kindo peninsula are characterized by a similar taxonomy. As the distance from the sea increases, the composition of the dominant species of fungi in the peat and swamp ooze of the peatlands changes. We found similar species of fungi in a specific component (e.g., peat or ooze) between the peatlands, rather than in the peat and ooze of a single peatland. The sphagnum peat of the lake coast (Fig. 2, point 1) is thin in comparison with other studied peat bogs in the area (Grum-Grzhimaylo

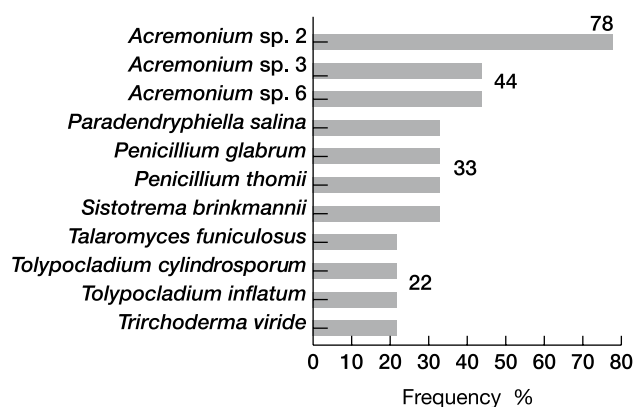


**Fig. 3** The spatial frequency of the most abundant fungal species in the Kisko-Sladkoe Lake

**Table 4** Similarity of fungal communities of the different parts of the Kisko-Sladkoe Lake (Sørensen index)

	Grassland soil	Forest soil	Littoral zone	Lake bottom
Sphagnum peat	0.20	0.18	0.21	0.09
Grassland soil	–	0.21	0.20	0.15
Forest soil	–	–	0.11	0.12
Littoral zone	–	–	–	0.23

et al. 2016). From the lake, despite the unfavorable conditions of the sphagnum peat (recalcitrant cell walls of sphagnum moss, abundant phenolic compounds, anoxia), we isolated a large diversity of fungi, including *Acremonium* spp., *Alternaria* spp., *Aspergillus versicolor*, *Beauveria* spp., *Botryotinia* sp., *Botrytis cinerea*, *Cadophora* spp., *Cladosporium* spp., *Oidiodendron* spp., *Lecanicillium* spp., *Penicillium* spp., *Talaromyces funiculosus*, *Tolypocladium*



**Fig. 4** The spatial frequency of fungal species (with the spatial frequency  $\geq 22\%$ ) in the littoral of the Kisko-Sladkoe Lake

spp., *Trichoderma* spp., *Umbelopsis* spp., sterile mycelia, some of which (*Cadophora* sp. 1, *Cadophora luteo-olivacea*, *Cladosporium* sp. 2, *Cladosporium sphaerospermum*,

*Fusicladium pini*, *Helotiales* sp. 5, *Lecanicillium* sp. 3, *Penicillium aurantiogriseum*, *Phaeoacremonium* sp., *Pseudogymnoascus pannorum*, and *Stachybotrys dichroa*) appeared only on the media which were based on the sphagnum extract (Table 3). Contrary to typical terrestrial peat, the sphagnum peat of the lake harbored a substantially larger fraction of fungi which are indicative of marine habitats, such as *Acremonium*, *Cadophora*, *Cladosporium*, and *Tolypocladium* (Online Resource 4). The fungal composition in the peat bogs of the terrestrial lakes in the area (Verkhneye, Krugloye and Yershovskoye Lakes), which have already separated from the sea, is similar, despite the different age and altitude above sea level (Grum-Grzhimaylo et al. 2016).

The benthic ooze (Fig. 2, points 9–15) is a more dynamic system due to differences in the organic matter inflow, exposure to external physical, chemical and hydrological conditions, and to the presence of marine sediments therein. Therefore, the fungal composition of the ooze was unique and differed between different lakes, except for sterile mycelia, the proportion of which was much higher compared to the peat. The ooze of both the terrestrial lakes and brackish lakes contained typical marine fungi of *Acremonium* which, by BLAST similarity search, are close to *A. fuci*, *A. potronii* and *A. strictum* (Jones et al. 2009). Thus, the presence of typical marine-borne fungi in the ooze of all the studied lakes reflects the process of their separation from the sea.

To make a direct comparison with the marine habitats, we compared the fungal composition of the lake with the fungal community in the bottom sediments of the Velikaya Salma Strait (part of the White Sea near the lake). Bubnova et al. (2014) detected three obligate marine species in the sediments (*Acremonium fuci*, *Paradendryphiella arenaria* and *P. salina*), all the rest were classified as land-origin species. The fungal composition was represented mainly by the cosmopolitan species (*Penicillium* spp.) and typical soil species for this region (*Pseudogymnoascus pannorum*, *Penicillium* spp., *Tolypocladium* spp., and *Trichoderma* spp.). The distinctive feature of the fungal composition in the bottom sediments was a high abundance and rich diversity of *Acremonium* spp., which are known to often occupy marine-borne habitats (Bubnova et al. 2014). These fungi were also found in our samples taken from the littoral zone of the lake, especially at sites where sea water flows in. BLAST search results showed that all of them were close to the obligate and facultative marine species of *Acremonium*.

Many of the dominating and frequent species distinguished from different components of the lake appeared to be typical representatives of nearby habitats (*Acremonium* spp., *Penicillium* spp., *Tolypocladium* spp., and *Trichoderma* spp.).

Of note, the littoral samples, which are the closest to the sea (Fig. 2, points 5, 7), were enriched with the fungi typically found in marine habitats (*Paradendryphiella salina*,

*Acremonium* cf. *fuci*, and *Cadophora* spp.). Conversely, the peat samples from the shore, which are desalinated and covered with vegetation (Fig. 2, points 1, 2, 4), contained fungal species commonly associated with peat bogs (*Oidiodendron* spp., *Penicillium thomii*, *P. montanense*). *Penicillium montanense* was recovered from all the samples of the coastal area (forest soil, sphagnum peat, meadow soil and the place of the freshwater brook confluence), but was not found in samples taken at places connected to the sea or in benthic ooze sediments. It can be assumed that pH values inhibit the spread of this species. *Penicillium montanense* is often noted in acidic habitats, such as decomposing bogs and fens, needle litter, and the rhizosphere of coniferous (Thormann et al. 2004; Osono et al. 2006), while sea water has an alkaline reaction.

Selective media makes it possible to recover slowly growing oligotrophic and acid-tolerant species that are able to thrive under unfavorable conditions, such as low pH and the presence of recalcitrant compounds of sphagnum moss. The highest number of species was isolated on malt extract agar and citric acid buffer agar media. However, the use of alkaline and neutral media allowed us to identify the alkali-tolerant and neutrophilic species (*Acremonium* spp., *Plectosphaerella* sp.) characteristic of many parts of the lake. The results of using the different media are shown in Online Resource 5. DNA-based identification was most useful with basidiomycetous fungi, though also with ascomycetous species of the Helotiales and Pleosporales. BLAST search results showed that many of our pleosporalean isolates from the lake were close to known marine species, such as *Bissothecium circinans* and *Loratospora aestuarii* (Schoch et al. 2009). Similarly, fungi from the Helotiales have often been found in marine habitats (Wang et al. 2006). In our studies of the fungi in the Kindo peninsula peatland lakes, the major part of the Helotiales were found in the lake (Table 5). Presumably, this is related to the fact that conditions in this lake are more similar to the sea conditions in comparison with the other lakes of the Kindo peninsula. The most common basidiomycete in the lake was *Sistotrema brinkmannii*, which is widely distributed in deciduous and coniferous forests, and known as a late-stage wood destructor. In the current study, this species was found in the swamp peat along the peninsula and from most of the samples taken from different parts of the lake. The presence of *Sistotrema brinkmannii* both in the littoral soil and benthic sediments of the water basins is intriguing, and its exact role in these wood-free habitats warrants further research.

Our investigations revealed that the fungal communities in the Kindo peninsula peat and lakes depend on their origin. Since the lake is similar in its properties to the other lakes that we studied in the WSBS area (depth less than 5 m and presence of a freshwater flow), it could be assumed that the lake would develop in the same way—from eutrophic

**Table 5** Helotiales taxa distribution in the Kislo-Sladkoe Lake, and the Yershovskoye, Krugloye and Verkhneye lakes

Lake	Number of taxa
Kislo-Sladkoe	15
Ershovskoye	8
Krugloye	2
Verkhneye	9

to oligotrophic, through the mesotrophic and upper aapa-type swamp stages (Olunina 2008). Hence, in the process of their waterlogging, as well as in other swamps, there will be a respective change in plant communities, the organic composition of sediments, and, consequently, in the fungal content. The presence of both typical marine species and species characteristic for freshened water basins, soil and peat, indicates the lake's transitional stage from the sea to a groundwater basin, resulting in a combination of abiotic factors. In future research, it would be of interest to study the functional role and adaptation trajectories of fungi in such environments.

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## Compliance with ethical standards

**Conflict of interest** The authors declare that they have no conflict of interest.

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