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Aquatic hyphomycetes communities occur as potential bioindicators of water quality

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Abstract

With a profound knowledge of how physico-chemical parameters affect these communities, microbial communities could be used as indicators for environmental changes and for risk assessment studies. We studied aquatic hyphomycete communities in ponds from sites shaped by intense human activities (namely the “Mansfeld region”) and Urmil river, there are no sources of man made effluents in Chhatarpur district. Environmental stress factors such as high concentrations of chloride and calcium as well as low concentrations of oxygen significantly reduced the diversity and biomass of hyphomycetes in the investigated samples. Fungi were negatively correlated with nutrient concentrations. We propose that aquatic hyphomycete communities can be used as sensitive and integrative indicators for freshwater quality.

Key words: Aquatic hyphomycetes; freshwater; community structure; bioindicator

1. Introduction

Aquatic hyphomycetes, an important part of the fresh-water microbial communities, are excellent candidates for bioindicators. They play a vital role in the functioning of aquatic ecosystem and fulfill crucial activities as part of food webs and in nutrient cycling. They initiate the degradation of organic material by e.g. increasing the palatability of litter and wood. In addition, aquatic hyphomycetes can degrade organic compound in harsh habitats such as rivers highly enriched in nutrients (Pascoal *et al.*, 2003) [10].

There is a general trend that, independent of its type, pollution usually leads to a decline in the diversity of aquatic hyphomycetes. A drastic decline in diversity and sporulation of aquatic hyphomycetes, accompanied by changes in species dominance (e.g. top conidia producers) has been reported in habitats highly polluted by heavy metals (Krauss *et al.*, 2001) [6]. However, monitoring the relative abundance of individual microbial species within their polluted habitats is of prime importance to detect relevant species that can act as bioindicators in risk assessment monitoring programs.

The objectives of this study were to (1) extract the main environmental factors responsible for constraining the community structure of aquatic hyphomycetes; and (2) investigate whether species assemblages of aquatic hyphomycetes might be used as potential indicators for the detection of minute environmental changes. We focused on fresh-water ponds and river in Chhatarpur district.

2. Material and methods

2.1 Site location

Three water bodies in the Chhatarpur district were selected for the field investigation. Two were the Kishore sagar and Pratap sagar ponds, which was chosen because it represents a pond with a high nutrient load. Indeed, numerous houses and shops are located along its shores. The other water body was Urmil river, which passes through the teak (*Tectona grandis* L.), Salai (*Boswellia serrata* L.), Khair (*Acacia catechu*) and Bamboo (*Dendrocalamus strictus*) forest area. Except for occasional dwellings and small farms along its banks, there are no sources of man-made effluents which might contribute nutrients to the stream.

For mycological analysis, samples of foam, submerged leaves and twigs were collected between 15th and 30th of february and processed as described by Nilsson (1964) [9]. Water samples were also collected for physico-chemical analysis.

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2.2 Water chemistry

The physico-chemical parameters of water viz., Temperature, water pH, dissolve oxygen (mg/l), nitrate (mg/l), phosphate (mg/l), chloride (mg/l) and calcium hardness (mg/l) were analyzed following the methods of A.P.H.A. (2005) [1].

Temperature was measured by using a centigrade thermometer by dipping it at a depth of 5-8 cm in water for 5 minutes, at the time of sample collection. pH was recorded on spot with the help of a digital portable pH meter (Hanna) at the time of sample collection. Dissolve oxygen content (mg/l) was determined on the spot by making a composite sampling of water at each month following the Winkler's azide modification method. Nitrate (mg/l), phosphate(mg/l), chloride (mg/l) and calcium hardness (mg/l) were analyzed following the methods of A.P.H.A. (2005) [1].

Table 1: Hydrochemical characteristics of the sampling sites

Chemical characteristics	Urmil river	Kishore sagar pond	Pratap sagar pond
Temperature	19.2	20.3	20.2
pH	7.6	9.4	9.1
Dissolved oxygen (mg/l)	7.3	5.9	4.5
Phosphate (mg/l)	0.00	0.36	0.49
Nitrate (mg/l)	0.09	4.00	1.89
Chloride (mg/l)	29.5	245.9	125.3
Calcium hardness (mg/l)	28.4	62.8	48.2

3. Results

3.1 Fungal diversity

A total of 20 species belonging to 14 genera of water borne conidial fungi viz., *Anguillospora*, *Beltrania*, *Campylospora*, *Clavariopsis*, *Dactylella*, *Flabellospora*, *Flagellospora*, *Isthmotricladia*, *Lemonniera*, *Lunulospora*, *Setosynnema Trichocladium*, *Tricladium* and *Triscelophorus* were isolated from Urmil river (Table 2). These species were isolated from different decomposed leaf litter of known and unknown species and foam.

Water of Kishore sagar and Pratap sagar pond get polluted by washing of the clothes and bathing by the people. Under these circumstances, aquatic hyphomycetes may be reduced. However, only one fungal species *Lunulospora curvula* was recorded from the Kishore sagar and Pratap sagar pond. According to Manoharchary (1989) [7], aquatic hyphomycetes disappear from such habitats as a result of pollution and anaerobic conditions.

3.2 Abiotic environment

Water temperature at all sites ranges between 19.2 °C to 20.3 °C. The pH value at all sites ranged from slightly basic (7.6) to alkaline (9.4). Concentration of dissolved oxygen at all sites fluctuate between ranged between 5.9-7.3 mg/l. The lower values (5.9 and 4.5 mg/l) were measured in the ponds water. In river water dissolved oxygen concentration was 7.3 mg/l. These fungi require a fresh oxygenated environment for their occurrence (Webster & Towfic, 1972). During study period, two ponds showed the alkaline condition (9.1 and 9.4) but the river water showed the slightly basic condition (7.6). pH greatly affects the decomposition activities of aquatic hyphomycetes in running fresh water bodies. The occurrence and degradative ability of these fungi colonizing on submerged leaf litter is influenced by the hydrogen ion concentration (pH) of water (McKinley & Vestal, 1982) [8]. Species richness of aquatic hyphomycetes declines noticeably at pH ≤ 4.5 and ≥ 8.0 (Barlocher 1987; Chamier 1992) [4, 5].

The nitrate value tended to be higher at Pratap sagar pond (1.89 mg/l) and Kishore sagar pond (4.00 mg/l) whereas lower at river Urmil (0.09 mg/l). Similarly, phosphate value be slightly higher at Pratap sagar pond (0.49 mg/l) and Kishore sagar pond (0.36 mg/l) whereas not found at river Urmil (0.00 mg/l). The concentration of chloride (245.9 mg/l) and calcium (62.8 mg/l) was much higher at Kishore sagar pond.

Table 2: Distribution of aquatic hyphomycetes found in the sampling sites

	Urmil river	Kishore sagar pond	Pratap sagar pond
<i>Anguillospora crassa</i> Ingold	+	-	-
<i>A. longissima</i> (Sacc. and Sydow) Ingold	+	-	-
<i>Beltrania rhombica</i> Penzig,	+	-	-
<i>Campylospora chaetocladia</i> Ranzoni	+	-	-
<i>Clavariopsis aquatica</i> de Wildeman,	+	-	-
<i>Dactylella submersa</i> (Ingold) Nilsson,	+	-	-
<i>Dactylella rhombospora</i> Grove	+	-	-
<i>Flabellospora crassa</i> Alasoadura	+	-	-
<i>Flabellospora verticillata</i> Alasoadura	+	-	-
<i>Flagellospora penicilloides</i> Ingold	+	-	-
<i>Flagellospora penicilloides</i> Ingold	+	-	-
<i>Isthmotricladia gombakiensis</i> Nawawi,	+	-	-
<i>Isthmotricladia laeensis</i> Matsushima	+	-	-
<i>Lemonniera cornuta</i> Ranzoni	+	-	-
<i>Lunulospora curvula</i> Ingold	+	+	+
<i>Lunulospora cymbiformis</i> Miura	+	-	-
<i>Setosynnema indica</i> sp. nov.	+	-	-
<i>Trichocladium angelicum</i> Roldan and Honrubia	+	-	-
<i>Tricladium indicum</i> Sati and Tiwari	+	-	-
<i>Triscelophorus monosporus</i> Ingold	+	-	-
Species richness	20	1	1

The table shows the presence (+) and absence (-) of the individual aquatic hyphomycete species.

4. Discussion

At the investigated sites, the pollution was mainly caused by increased concentration of nitrate and chloride (Table-1). In our study, the maximum concentrations of nutrients measured at Pratap sagar pond and Kishore sagar pond. Consequently fungal biomass and sporulation rate were clearly inhibited in these sites compared to the Urmil river (Table-2) suggesting that the positive role of nutrients on fungal growth might be true up to a certain threshold, beyond which biological activity might be inhibited.

Aquatic hyphomycetes live preferentially in clean and well aerated water habitats (Barlocher, 1992, 2005) [3, 2]. Results of present study showed that growth of aquatic hyphomycetes were generally inhibited with the amount of nutrients in the water. Sridhar *et al.*, 2000 [11]; Pascoal *et al.*, 2003 [10] also observed same results.

5. Conclusion

Aquatic hyphomycete communities presented in this paper can be used as sensitive and integrative indicators for freshwater quality. Microbial communities are suitable for detection of stress in natural systems because of their ubiquity and rapid response to environmental changes. Present study indicate that variations in water chemistry cause a significant proportion of the change in fungal community. This paper provide a future application of aquatic hyphomycetes as indicators for pollution.

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7. References

1. APHA. American Public Health Association Standard Methods for the Examination of Water and Waste Water, 17th edition, American Water Works Association. Water Pollution Control Federation Publication Washington, DC. 2005.
2. Barlocher F Freshwater fungal communities. In: Deighton J, White J F Jr, Oudemans P Editors. The fungal community: its organization and role in the ecosystem, Taylor & Francis, CRC Press, 2005, 39-59.
3. Barlocher F Community organization, In: F. Barlocher (ed.), The ecology of aquatic hyphomycetes. Springer. Verlag, Berlin Germany. 1992, 38-69.
4. Barlocher F Aquatic hyphomycete spora in 10 streams of New Brunswick and Nova Scotia. Canadian Journal of Botany. 1987; 65:76-79.
5. Chamier CA Water chemistry. In: The ecology of aquatic hyphomycetes. Edited by F. Barlocher. Springer. Heidelberg and New York. 1992, 152-172.
6. Krauss G, Barlocher F, Schreck P, Wennrich R, Glasser W, Krauss GJ. Aquatic hyphomycetes occurrence in hyper polluted water in Central Germany. Nova Hedwigia. 2001; 72:419-428.
7. Manoharachary C Glimpses on water borne conidial fungi from India. In: Perspective in Aquatic biology (ed. R.D. Khulbe), Papyrus Publishing house, New Delhi. 1989, 71-77.
8. McKinley VL, Vestal JR. Effect of acid on plant litter decomposition in an arctic lake. Applied and Environmental Microbiology. 1982; 43:1188-1195.
9. Nilsson S Fresh water hyphomycetes. Taxonomy morphology and Ecology. Symbolae Botanicae Upsalienses. 1964; 18:1-130.
10. Pascol C, Pinho M, Cassio F, Gomes P. Assessing structural and functional ecosystem condition using leaf breakdown: studies on a polluted river. Freshw Biol. 2003; 19:109-128.
11. Sridhar KR, Barlocher F. Initial colonization, nutrient supply and fungal activity on leaves decaying in streams. Applied and Environmental Microbiology. 2000; 66:1114-1119.
12. Webster J, Towfic FH. Sporulation of aquatic hyphomycetes in relation to aeration. Transactions British Mycological Society. 1972; 59:353-364.