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Diversity and Cellulase Activity of *Chaetomium globosum* (Kunze), A Destructive Cellulolytic Fungus

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Abstract: Fungi invading objects containing cellulose are called cellulolytic fungi. These fungi produce “cellulolytic enzymes or cellulases”, which decompose cellulose (a polysaccharide) into first oligosaccharide and then monosaccharides, and ultimately to glucose molecules. *Chaetomium globosum* (Kunze) is reported to be a destructive cellulolytic fungus causing damage to our cultural heritage of library books, monuments, paintings, archival materials etc. In the present investigation, diversity of *C. globosum* in different library books of District - Gorakhpur (located in North- Eastern Uttar Pradesh, India) has been studied. Cellulase activity of *C. globosum* isolated and cultured has been assayed using Carboxyl Methyl Cellulose (CMC) Agar medium technique. Gram's iodine solution is used as indicator. The Enzyme Index of *C. globosum* has been reported as 3.6, which shows that it is industrially important fungus, which may be exploited for the production of cellulase enzyme at industrial level. Therefore, this research work strongly supports the concept that screening of cellulase producing ability of *C. globosum* and other microorganisms is very important.

Keywords: Cellulolytic Fungi, Biodeterioration, Cellulase Activity, *Chaetomium globosum*, Mesophilic, Enzyme Index (EI).

Introduction

Chaetomium globosum (Kunze) is a mesophilic cellulolytic fungus. It has a vast range of substratum for its growth including plants, soils, debris of plants and animals, papers, textiles and clothes etc. (Sharma *et al.* 2025). In addition to its saprophytic nature, strain of *Chaetomium globosum* has been reported to be “Endophytic”, assisting in cellulose decomposition of plant cells (Chapman *et al.* 1975). Its habitat ranges from forest plants to soils of various biomes (Domsch *et al.* 1980; Liu *et al.* 2007); to indoor environment including books and wood (Provost *et al.* 2013). Rarely

some strains of *Chaetomium globosum* are reported to be human allergens, causing Mycosis and neurological infections (Kim *et al.* 2008; Provost *et al.* 2013).

In humid tropical and sub-tropical countries of the world, biodeterioration of various cultural commodities in textiles, papers, wood and leather is a very common phenomenon (Tiano, 2009; Ortega - Morales *et al.* 2019).

Our cultural heritages are heavily damaged and destroyed by these micro-organisms (Koestler and Vedral, 1991). The two most important microbes responsible for this biodeterioration are fungi and bacteria. Fungi of various groups play most active role and cause maximum damage to these important objects (Arroyo, 2007).

District Gorakhpur is situated in North - Eastern Uttar Pradesh, India, in foot hills of Himalayas at the bank of Rapti river (coordinates; 26.7637152° N; 83.4039116° E), with average annual temperature of 26°C (79°F); elevation- 75m (246ft). Its high relative humidity and moderate temperature from July to March are very suitable for growth of Mesophilic microbes including fungi (Srivastava, 2007).

Fungi having ability to hydrolyse cellulose (a polymer of glucose) into its simpler units are called cellulolytic fungi (Arnthong *et al.* 2020). Twenty-six species of these cellulolytic fungi belonging to 15 genera have been reported causing biodeterioration of Webster's Dictionary in Gorakhpur (Srivastava *et al.* 2011). These fungi by producing cellulase enzymes, cause biodegradation of our cultural heritage containing cellulose, including important library books *etc.* (Mesquita *et al.* 2009; Sterflinger *et al.* 2013). Their activities are supported by high moisture content, moderate light intensity and temperature (Garg *et al.* 1995; Sterflinger *et al.* 2012; Coronado-Ruiz *et al.* 2018).

Biodeterioration of papers and archival material caused by fungi is very common in India (Agrawal, 1995). Lignocellulosic biomass is heavily decomposed by these cellulolytic fungi (Anwar *et al.* 2014). These fungi cause biodegradation of cellulose containing objects by producing three types of cellulases - "Endo 1,4-β-D-glucanase (EG), Cellobiohydrolase (CBH) or exo 1,4, β-D-glucanase and β-glucosidase" (Teeri, 1997; Bayer *et al.* 1998; Zhang and Lyund 2004; AL-Kharousi *et al.* 2015). Degradation of cellulose is achieved by the combined effects and actions of these three cellulase enzymes (Qin *et al.* 2010).

Chaetomium globosum with other cellulolytic fungi has been reported to degrade cellulose, causing damage of important documents. (Pinzari *et al.* 2006; Michaelsen *et al.* 2009).

The cellulase producing and decomposing ability of various cellulolytic fungi including *Chaetomium* is reported by several workers (Ames, 1961; Aranyanak, 1995; Al-Kharousi *et al.* 2015).

Exploitation of these cellulolytic fungi and their enzymes at industrial scale is the need of time (Arif *et al.* 2024). Control of these cellulolytic fungi and protection of our cultural heritage in papers and cloths/textiles is a challenging job (Sequeira *et al.* 2017). However, isolation of these cellulolytic fungi from deteriorated objects and their identification is the initial step. Pure cultures of these fungi are then used to evaluate their cellulolytic activities by calculating their enzyme index (EI).

In the present research work, isolation of cellulolytic fungi from deteriorated samples of papers and cloths/textiles has been done to study the presence of *C. globosum*. Cellulase activity of isolated *C. globosum* has been determined by calculating its enzyme index (EI).

Materials and Methods

1. Isolation of Test Fungus - *Chaetomium globosum* Kunze

Various sites of District Gorakhpur (such as Government District Library; Central Library of D.D.U. Gorakhpur University and Central library of St. Andrew's Post-Graduate College; Buddha Museum and whole sale market of clothes/textiles) were visited and infested samples of papers and cloths/textiles were collected from these sites.

For isolation, 3 techniques were used – Direct Observation, Standard Blotter Method and Czapek Dox Agar Method.

Six species of *Chaetomium* were isolated and reported from deteriorated samples of papers and cloths/textiles, which were given Isolate/Culture Numbers (**Table – 1**).

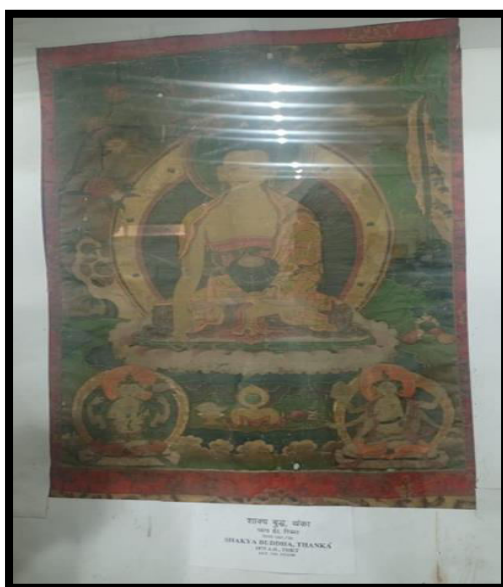
Table – 1
Frequencies of Species of *Chaetomium* Kunze Isolated from Deteriorated Samples of Papers and Cloths/Textiles

S. No.	Species of <i>Chaetomium</i> Isolated	Isolate/Culture No.	DO	SBM	CDA
1.	<i>Chaetomium dolichotrichum</i> Ames	SP/AML/0122	+	+	++
2.	<i>C. funiculosum</i> Corda	SP/AML/0123	+	+	
3.	<i>C. globosum</i> Kunze	SP/AML/0124	+++	++	+++
4.	<i>C. strumarium</i> Minter	SP/AML/0125	+	+	++
5.	<i>Chaetomium</i> sp. (US) - 1	SP/AML/0126	+	-	+
6.	<i>Chaetomium</i> sp. (US) - 2	SP/AML/0127	+	-	-

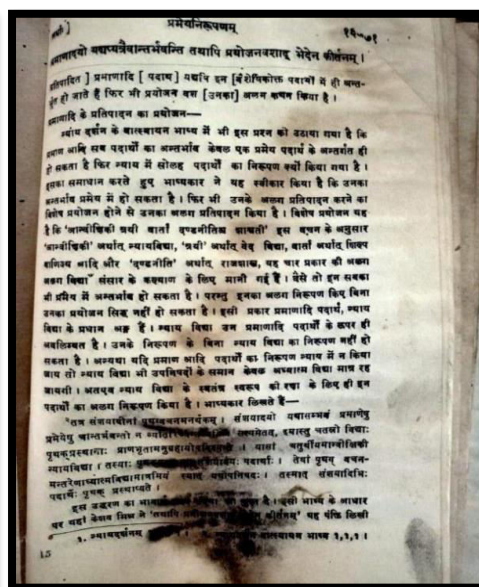
+++	:	Most Frequent	DO	:	Direct Observation
++	:	More Frequent	SBM	:	Standard Blotter Method
+	:	Less Frequent	CDA	:	Czapek Dox Agar
-	:	Absent	US	:	Unidentified Species

As it is evident from **Table – 1**, maximum frequency observed was of *C. globosum*, which was selected as the test fungus. It was cultured on Potato Dextrose Agar (PDA) Medium.

The pure culture of isolated *C.globosum* (**Isolate/Culture No. SP/AML/0124**) was deposited in NFCCI (National Fungal Culture Collection of India), Pune, Maharashtra, India to get Accession No. (**Accession. No. – NFCCI5963**). Also, the pure culture of *C. globosum* was procured from ITCC (Indian Type Culture Collection), Indian Agricultural Research Institute (IARI), New Delhi (**ITCC. No. - 3680**).



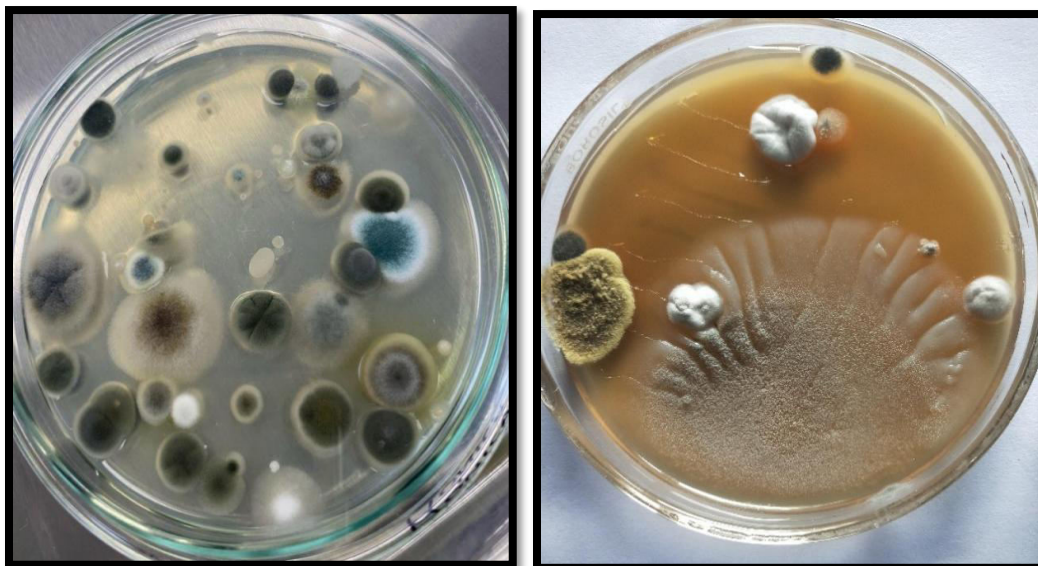
**Shakya Buddha, Thangka
(1875 AD, TIBBET) infested by
Cellulolytic fungi**



**Infested page of
“Hindu Dharmkosh”**



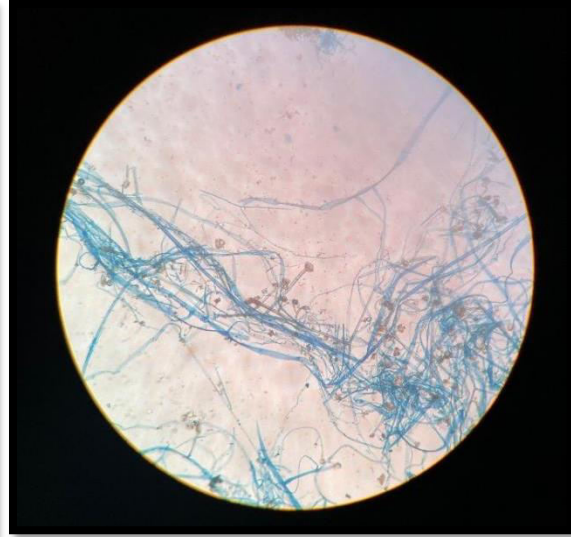
Damaged Book entitled “SUKH SAGAR” Isolation of Cellulolytic Fungi (1856)



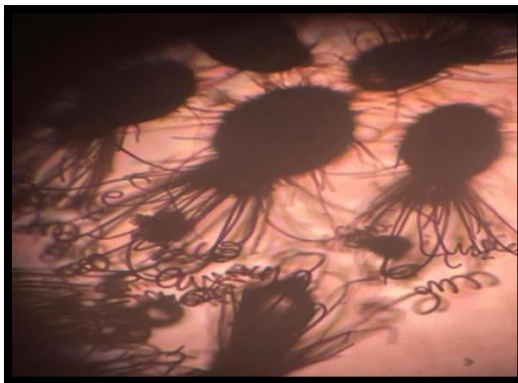
Mixed Culture of Cellulolytic Fungi Isolated



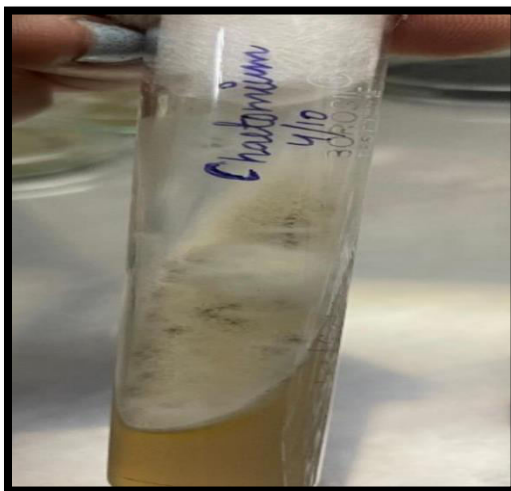
Pure Culture of *Chaetomium globosum*



Mycelium of *Chaetomium globosum*



Spores of *Chaetomium globosum*



Slant of *Chaetomium globosum* for



Pure Culture of *Chaetomium globosum*

Deposition in NFCCI, Pune procured from ITCC, IARI, New Delhi

2. Cellulase Assay of *Chaetomium globosum*

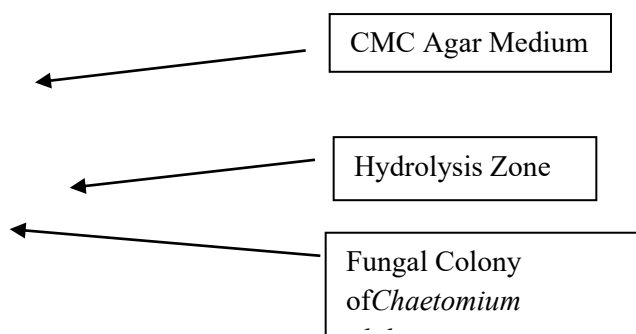
Cellulase assay of the test fungus *C. globosum* was done by using 1% Carboxyl Methyl Cellulase Agar medium (CMC AM).

Carboxyl Methyl Cellulose serves as source of carbon and Gram's iodine solution as indicator to observe cellulose hydrolysis zone (Coronado - Ruiz *et al.* 2018). In this process of qualitative determination of cellulase activity, the Gram's iodine, react with cellulose as well as its decomposed components. The unhydrolyzed and intact cellulose holds the colour of Gram's iodine solution. However, that cellulose which is hydrolysed by fungal cellulase enzymes reflects 'clear zone' (seen as pale-yellow zone). The enzymatic index of *C. globosum* was calculated as-

$$\text{EI of } C. \text{ globosum} = \frac{\text{Diameter of Hydrolysis Zone}}{\text{Diameter of Colony Growth of } C. \text{ globosum}}$$

To evaluate E.I., 6mm. diameter disc of pure culture of *C. globosum* was cultured on 1% CMC Agar Medium and was incubated for 7 days at 28 ± 2 °C temperature. The antibiotic used was Streptomycin. Gram's Iodine solution was used as indicator. Afterwards, this culture was washed with water. Colony diameter of *C. globosum* and clear zone around the fungal disc were measured to calculate enzyme index (Coronado- Ruiz; 2018).

Observations



Cellulase Assay of *Chaetomium globosum*

1. Diameter of hydrolysis zone (Diameter of colony + clear zone) = 36mm.
2. Diameter of colony growth of *C. globosum* = 10mm

Therefore;

$$\text{EI of } C. \text{ globosum} = \frac{36 \text{ mm.}}{10 \text{ mm.}} = 3.60$$

Result and Discussion

Six species of *Chaetomium* have been isolated and cultured from deteriorated samples of papers and clothes/ textiles. Of these 6 species, 4 have been identified as *Chaetomium dolichotrichum* Ames, *C. funiculosum* Corda, *C. globosum* Kunze and *C. strumarium* Minter. However, two unidentified species (US) were also isolated and were given culture numbers for future studies. Of all these 6 species, *C. globosum* was reported to be the most frequent species collected from all the samples.

Enzymatic index (EI) of *C. globosum* has been calculated as **3.60**

The results of present work clearly shows that *C. globosum* is efficient in cellulase enzyme production and cellulase producing ability of this species can be exploited at industrial level. Also, the objects containing cellulose, such as books, archives, cloth and textiles etc. could be a good source to explore and isolate cellulase producing fungi like *C. globosum* etc.

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Authors Contributions

All authors contributed significantly to the final manuscript.

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