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AIR BORNE MYCOFLORA OF EXTRA AND INTRAMURAL ENVIRONMENT OF COLLEGE LIBRARY

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ABSTRACT

The present investigation was undertaken to find out the concentration of the various air borne fungal spores in the outdoor and indoor environment of the College library. The mycoflora of indoor and outdoor environment of library in Ozar (MIG) College, Dist. Nashik (M.S) was investigated for the period of one year i.e. from October 2013 to September 2014 by Himedia Petri plate air sampler No. LA-030 with Himedia readymade Rose Bengal Chloramphenicol Agar culture plates techniques. There were total 21 air borne fungal colonies were isolated and identified in the ambient air of college library. *Aspergillus* (19.82% and 28.2%), *Cladosporium* (8.99% and 7.20%) *Penicillium* 8.23% and 5.30%), *Fusarium* (5.34 % and 4.50%) and *Alternaria* (6.10 % and 9.50%) were found to be all time occurrence and shown predominance in the outside and inside the library environment respectively. Lignocellulosic degrading fungus like *Chaetomium* (5.18% and 4.19 %) was also recorded in high humid and low temperature conditions, *Alternaria*, *Fusarium Helminthosporium*, *Trichoderma*, *Rhizopus*, *Curvularia*, *Tetraploa*, *Periconia*, *Stemphyllum* and some unidentified colonies were recorded in high concentration during the investigation with little variation in their concentration. The Total Colony Forming Units per M3 of air was more inside the library (1782/M3 of air) than outside (1680/M3 of air). The air borne fungal spores recorded in the library environment would be the main cause of some health hazards like sneezing, watery eyes, coughing and uneasy feelings in some sensitive visitors of the library. The air borne fungal spore concentration also shows close relation with the meteorological parameters like temperature, humidity and rainfall.

Keywords: *Library, Allergy, Airborne Fungi, Weather Conditions*

INTRODUCTION:

Due to suspended air borne particles in the library environment, both the readers and books get infected at more or less percentage. The enclosed environment of the library offer unique substrate such as binding glue, paper and other organic material which support the active fungal growth. Because of light weight and microscopic nature, the spores of fungi remain suspended in the air for long period. The concentration of these bioparticles shows great variation from time, season, altitude and weather conditions (Lacey 1991). Since the study of allergic aerospora of library environment is of great importance and continuous exposure to the library causes the respiratory disorders. A very meagre studies were reported on similar matter (Bürge 1978, Vittal 1985, Tripathi 1987, Singh 1990, Agashe 1996, Ghani 1993, Verma 1996, Tilak 1997, Saoji 1997, Atluri 2002, Md. Shabbir 2003, Upadhyay 2005, Giri 2012,) hence the present work is undertaken to

study the allergic air borne fungal spores in the ambient air of library. Respiratory allergy is most prevalent among all populations with increasing trend all over the world. The bioparticulates implicated to cause allergic symptoms are pollen grains, fungal spores, insect debris, house dust mites, animal dander, chemicals and food etc. Today, more than 300 million of the world population is known to suffer from one or other allergic ailments affecting the socioeconomic quality of the life. Most people spend maximum of their time in the indoor environments of house, offices, factories and other indoor work places. This continuous exposure to the close indoor environment to many air borne pollutants comes through inhalation of air. The respiratory allergy is the most severe disorder including asthma, rhinitis, dermatitis, hay fever, skin rashes, sneezing, coughing, running nose, watering eyes etc. The major causative agents for the respiratory allergy are pollen grains, fungal spores, dust, mites, insect debris etc (Pande 2012).

MATERIAL AND METHOD:

The town Ozar (MIG), Maharashtra, is well known for its grape and onion production and similarly for air craft production at Hindustan Aeronautics Limited. The arts, Science and Commerce College, Ozar (MIG) was established along with central library in the year 1985. The total carpet area of the library is 1901 sq. fts consisting total 24,130 valuable books including text, reference books, periodicals and journals worth of Rs.27,38018/- up to year March 2014. The books were arranged subject wise and kept on open and closed steel racks keeping 3" distance between the racks.

The air sampling in the College library was done by using Himedia air sampler No. 030 with readymade Himedia Rose Bengal Chloramphenicol nutrient media plates. The air sampler was loaded with nutrient media plates and placed on the reading table for 10 minutes every forth night between 11.00 am to 11.30 am, i. e during peak hours of book transactions. The sampling of the air was done outside and inside the library environment from the months of October 2013 to March 2014 in the first season and April to September 2014 in the second season. After the exposure, the petriplates are brought to the botany laboratory for the incubation at the temperature 25-27 °C for 4-5 days. The Colony Forming Units (CFU) were counted and Total Fungal Counts (TFC) were expressed as Colony Forming Units per cubic meter of air (CFUs/M³).

The fungal colonies detected per unit volume of air are calculated as under;

$$\text{CFUs/M}^3 = \frac{\text{Number of Colonies on nutrient medium plates (N C)} \times 25}{\text{Sampling time in minus (T)}}$$

After the incubation period of 5-6 days the colonies were appeared on the nutrient medium petriplates. These colonies were counted and identified up to genus level with the help of available literature (Barnett 1971, Tilak 1985).

RESULTS AND DISCUSSION:

Studies reveals the total 21 types of fungal colonies recorded in the ambient air of the library. High concentrations of fungal colonies were recorded in the library environment (326) than outside the library. Highest concentration of colonies was recorded to be of *Aspergillus* (inside 28.57 % and outside 17.485 %), *Cladosporium* (13.19% and 12.14 %), *Penicillium* (8.9 % and 4.29%), *Helminthosporium* (6.13 % and 5.0%) and *Alternaria* (4.29% and 7.86%) to the total colonies recorded inside and the outside the library environment respectively. (Table 01 and 02) In addition to the cellulose, the paper also contains the lignin, hemicellulose, pectin, waxes, tannins and minerals useful for the fungal growth. Beside the impurities formed during the paper production are helpful for the nutrition of fungi (Tilak S T 1985). The cellulytic activity of some paper deteriorating fungi produces endogluconase enzymes which damages the paper quality of the library book (Verma K S 1987) indicate the role of fungi in book deterioration.

Present studies indicates the high concentration of the fungal colonies outside the library in the months of October to March 2013 while there was gradual increase the concentration inside the library from the Months of April to September 2014 might be due to high humid and low temperature conditions. (Fig.01 & 02). Some fungal types like *Drechslera*, *Epicoccum*, and *Geotrichum* were only reported in the rainy months of July to September 2014. High concentration of fungal colonies (713) inside the library environment indicates humid and damp environmental conditions which would be favourable for the growth of moulds on the books. Chauhan et al (2006) reported the similar observation by studying 7 different libraries at Agra. The adverse effects of fungal components not only deteriorate the paper material but also affect the library staff. Agashe (1991) also reported 23 types of fungal colonies in the indoor environment of library at Bengaluru. Giri S K (2012) reported 36 fungal spore types from the ambient air of library at Nagpur adding highest concentration of *Aspergillus*, *Cladosporium*, *Alternaria* and *Rhizopus* spores. Predominance of *Aspergillus* and *Cladosporium* in the present studies is in relation with the reports of Vittal (1985) and Singh (1990). The cellulose degrading fungi like *Chaetomium* was also recorded (5.18% and 4.19 %) indicating the threat to the stored books in the library environment. Dhawan (2005) critically reviewed the role and aerobiodeteriogens on paper and museum material indicating high concentration of versatile fungi like *Aspergillus*, *Penicillium*, *Cladosporium* and *Fusarium*. As indicated by Nandimuthan (1995) and Tripathi (1987) the present studies also reveals high concentration of fungal colonies during the high humid, low temperature conditions in the month of December 2013 and January 2014 than in high temperature and low humid conditions in the month of February and March 2014. Since the study of allergic airospora of library environment is of great importance and continuous exposure to the library causes the respiratory disorders, A very meagre studies were reported on similar matter (Tilak S T (1982), Verma K S (1987), Tripathi S N (1987), Nandimuthan (1995), Shaney (2001), Sharma (2004), Tiwari K L (2004). And Mujumdar (2005).The studies inside and outside the library environment is of great importance since the library environment is loaded with the biopollutants that can be harmful to the health of the library users, beside reducing the life of books (Agashe S N 1996)

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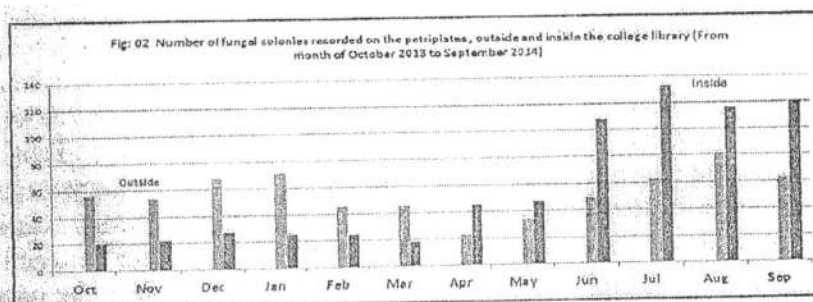
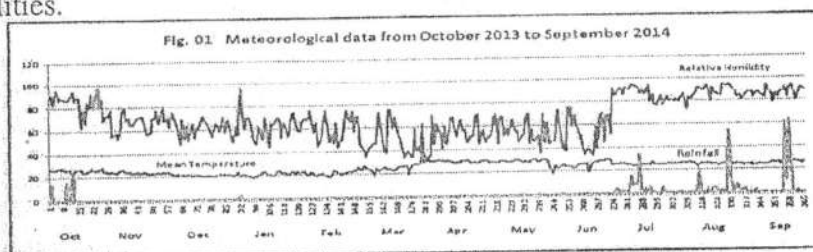


TABLE: 01

Fungal Types	Number of colonies recorded on the petriplates (Outside the library)													Total	% Conc	CFU/M ³
	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep				
<i>Alternaria</i>	3	4	1	2	1	3	-	2	4	7	8	5	40	6.10	100	
<i>Aspergillus</i>	12	7	11	12	7	8	6	8	5	15	22	17	130	19.82	325	
<i>Cercospora</i>	3	2	2	-	-	2	-	-	3	-	5	-	17	2.59	42.5	
<i>Chaetomium</i>	3	5	4	6	3	2	3	-	3	-	2	3	34	5.18	85	
<i>Cladosporium</i>	12	3	10	11	4	3	2	4	5	-	3	2	59	8.99	147.5	
<i>Corynospora</i>	-	-	-	-	-	-	2	7	-	5	2	1	17	2.59	42.5	
<i>Curvularia</i>	-	-	3	4	1	-	-	2	5	-	5	-	20	3.05	50	
<i>Drechslera</i>	2	1	-	2	1	1	-	-	-	-	-	-	-	-	-	
<i>Epicoccum</i>	3	2	4	3	5	4	2	1	2	4	3	7	17	2.59	42.5	
<i>Fusarium</i>	2	-	3	-	4	6	-	-	-	-	-	-	35	5.34	87.5	
<i>Geotrichum</i>	2	4	6	4	3	1	-	3	1	-	-	-	15	2.29	37.5	
<i>Helminthosporium</i>	-	-	1	1	2	-	-	-	3	-	3	3	30	4.57	75	
<i>Nigrospora</i>	3	5	7	8	4	2	2	3	3	7	7	3	54	8.23	135	
<i>Penicillium</i>	-	1	-	-	-	-	-	-	-	-	-	-	1	0.15	2.5	
<i>Phoma</i>	-	-	-	1	1	-	-	-	6	3	2	5	21	3.20	52.5	
<i>Periconia</i>	3	5	2	3	1	-	-	-	-	1	-	2	5	0.76	12.5	
<i>Pringsheimia</i>	2	5	3	7	2	1	-	-	2	3	7	3	29	4.42	72.5	
<i>Rhizopus</i>	3	4	2	2	4	5	2	-	2	2	1	2	29	4.42	72.5	
<i>Stemphyllum</i>	3	4	2	2	4	5	2	-	2	2	1	2	29	4.42	72.5	
<i>Trichoderma</i>	4	5	7	4	2	7	4	3	3	7	3	9	58	8.84	145	
<i>Unclassified</i>	4	5	7	4	2	7	4	3	3	7	3	9	58	8.84	145	
Total	57	54	68	71	46	45	23	34	50	63	82	63	656	100	1640	

TABLE: 02

Fungal Types	Number of colonies recorded on the petriplates (Inside the library)													Total	% Conc	CFU/M ³
	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep				
<i>Alternaria</i>	2	3	2	2	-	2	3	-	23	12	9	10	68	9.54	170	
<i>Aspergillus</i>	3	3	7	10	12	5	2	17	33	42	23	44	201	28.19	502.5	
<i>Cercospora</i>	-	-	2	2	1	1	3	-	5	4	2	7	27	3.79	67.5	
<i>Chaetomium</i>	-	-	-	2	1	1	4	-	7	5	7	3	30	4.21	75	
<i>Cladosporium</i>	2	4	5	3	2	1	9	7	2	5	7	4	51	7.15	127.5	
<i>Corynospora</i>	-	-	-	-	-	3	2	4	-	7	-	7	16	2.24	40	
<i>Curvularia</i>	-	-	-	-	2	1	3	-	4	-	7	7	24	3.37	60	
<i>Drechslera</i>	-	-	-	2	-	-	2	-	1	4	-	2	12	1.68	30	
<i>Epicoccum</i>	2	1	-	-	-	-	4	2	4	8	7	4	32	4.49	80	
<i>Fusarium</i>	-	-	-	-	-	-	2	-	3	-	4	6	15	2.10	37.5	
<i>Geotrichum</i>	-	-	2	1	1	-	-	3	5	3	8	28	3.93	70		
<i>Helminthosporium</i>	3	2	1	-	-	-	2	7	3	7	7	6	38	5.33	95	
<i>Nigrospora</i>	-	-	-	1	1	-	-	1	-	7	3	-	11	1.54	27.5	
<i>Penicillium</i>	-	1	-	-	-	1	2	1	4	3	2	15	2.10	37.5		
<i>Phoma</i>	2	1	1	-	2	2	1	2	5	3	7	7	4	0.56	10	
<i>Periconia</i>	2	1	1	-	-	-	-	3	4	1	2	33	4.63	82.5		
<i>Pringsheimia</i>	2	1	1	-	-	-	-	3	4	1	2	17	2.38	42.5		
<i>Rhizopus</i>	3	4	2	1	1	4	3	3	4	10	12	6	53	7.43	132.5	
<i>Stemphyllum</i>	2	1	3	1	-	-	-	-	3	4	7	7	20	2.81	50	
<i>Trichoderma</i>	2	1	1	-	-	-	-	2	7	4	3	20	2.81	50		
<i>Unclassified</i>	3	4	2	1	1	4	3	3	4	10	12	6	53	7.43	132.5	
Total	21	22	28	26	25	18	46	48	108	133	117	121	713	100	1782.5	

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