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## Atmospheric Fungi of Karachi City

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**Abstract:** Altogether 53 species of fungi belonging to 21 genera present up to a height of 10m were trapped in the atmosphere of Karachi city. Species of *Aspergillus*, *Alternaria*, *Penicillium* and *Cladosporium* were the most frequently encountered. A data on the prevalence of atmospheric fungi isolated during 1998-1999 is presented.

**Key words:** Atmospheric fungi, Karachi, fungi, fungi in Karachi, mycoflora in Karachi

### Introduction

Mycoflora present in the atmosphere of a city or country is studied with a view to identify fungal spores responsible for causing plant and human diseases (Kim, 1982; Chapman, 1999; Baratawidjaja *et al.*, 1999). For the purpose of plant disease forecasting, atmospheric fungi is generally studied on seasonal basis. Necessary spraying measures are carried out, if plant pathogenic spores are found in the atmosphere. Fungicidal spraying do not allow the fungal spores to survive, germinate and cause diseases on plants. No such effort for the study of fungal spores present in the atmosphere has been made in Pakistan. Plant disease forecasting and plant disease diagnostic service is needed in Pakistan. There is a long list of plant diseases in Pakistan (Ghafoor and Khan, 1976) which demand attention. Spraying of pesticides is carried at times under the auspices of FAO for the control of locust in South Asia on regional basis, which includes Pakistan.

The city of Karachi and its suburbs is inhabited by 15 million people and there has been a breakdown of conservancy services which has made it one of the dirtiest cities of the world. The soil, water and air of Karachi city have become highly contaminated and polluted resulting in unhygienic condition. This appalling condition has increased the incidence of water and airborne diseases.

Fungi and bacteria contaminate water and pollute air and cause many human diseases (Gregory, 1973). Spores of fungi present in the air also cause a number of allergic reactions (Al-Doory, 1984 and Sanches *et al.*, 1999). Ebner *et al.* (1992) reported that for effective treatment of allergies, knowledge of the airborne fungal spores are of great importance. It is reported that each country has its own fungal groups which cause allergy. Chapman (1999) put emphasis on air sampling data of air spora to determine the cause of allergy. In different regions of the world research works have been carried out on airborne fungi causing fungal allergy (Angullo Romero *et al.*, 1999; Munuera *et al.*, 1998; Vanderbergh *et al.*, 1999).

Fungi cause mycoses (deCastro *et al.*, 1999; Elewski, Hirschmann and Raugi, 2000). Both in India and Pakistan people suffer from bronchitis, rhinitis, dermatitis, urticaria, eczema etc. (Bapat, 1994). There is no data on the occurrence of fungal spores in the atmosphere of Karachi city. A large number of people are reported to be suffering from allergy in the city of Karachi and a record of the skin hospitals of Karachi would indicate gradual increase in the incidence of dermal diseases (Mughal, 2002).

In view of the importance of air spora in the causation of plant diseases and allergic reactions in human beings, it was considered worthwhile to study and obtain a data on fungi present in the atmosphere of Karachi city.

### Materials and Methods

For the study of airborne mycoflora glass slides (each of 6x2 cm<sup>2</sup>) smeared with glycerine or a drop of sterilized water only on one side were exposed at 1.5, 5 and 10m heights at the Karachi University Campus, Karachi during 1998-99. A series of replicate slides were exposed for 24hrs and brought back to the laboratory

for microscopic study. A drop of lactophenol was placed on exposed slides and was covered with cover slips for observation under microscope. The method used for trapping the fungal spores from the atmosphere is devised by Agashe and Alfadil (1989).

For studying the various fungal members present in the atmosphere, the identifying characters of the trapped fungal spores were taken into consideration and calculated on percent basis of the total occurrence of the spores at each altitude.

If on the basis of the morphological characters of the spores it was not possible to identify then replicate samples of such spores as water mounts were transferred and cultivated on potato dextrose agar (PDA) or Sabourad dextrose agar (SDA) media in Petri plates for germination and growth. Incubation for 3-5 days on either of the two media yielded identifiable fungal colonies and spores. For the identification of fungi, spores and colonies arising from spores on selective media were tallied with the description given in authoritative literature such as Manual of Penicillia (Raper & Thom, 1949), Manual of Aspergilli (Thom and Raper, 1945) and Dematiaceous Hyphomycetes (Ellis, 1971).

### Results and Discussion

Spores of fifty-three fungal species belonging to 21 genera were trapped and identified from the atmosphere of Karachi city at Karachi University Campus up to 10 m altitude. The relative frequency of each fungal species as a percentage of its contribution to total counts at each altitude during 1998-99 is presented in Table 1.

The spore counts were taken at low altitude (1.5m above ground level) for *Aspergillus niger* (17.61), *A. flavus* (15.22) and for *Alternaria tenuissima* (8.955%). Similarly the highest count at medium altitude (5 m above ground level) were recorded for *A. niger* (19.14), *A. flavus* (15.978) and for *Alternaria alternata* - 6.88%. *Alternaria brassicae*, *A. brassicola*, *Aspergillus sulphureus*, *A. ustus*, *Cladosporium oxysporum*, *Fusarium culmorum*, *F. equiseti*, *F. longipes*, *Penicillium* sp., *Rhizopus nigricans* and *Ulocladium* sp., were not recorded from low altitude. *Aspergillus sydowii*, *A. ustus*, *Chaetomium globosum*, *Drechslera nodulosus*, *Fusarium culmorum*, *F. equiseti*, *F. semitectum*, *Mycelia sterilia*, *Penicillium* sp. and *Rhizopus nigricans* were not found between the low and high altitude. Spores of *Alternaria brassicae*, *A. brassicola*, *Aspergillus candidus*, *A. glaucus*, *A. nidulans*, *A. niveus*, *Aspergillus sulphureus*, *A. sydowii*, *Chaetomium globosum*, *Drechslera dematioides*, *Drechslera* state of *Cochliobolus nodulosus*, *Mycelia sterilia*, *Phoma* sp. and *Stemphylium* sp., were not recorded from high altitude (10m height). A good deal of variation is shown in occurrence of fungal species to the extent that some of the fungal species were less than 1% of the total findings at different altitudes (Table 1).

On an overall basis, 36.22% of fungi were recorded from low altitude (1.5m above ground), 39.9% from the medium altitude (5m height) and 23.86% spores were recorded from the top altitude (10m height). The species of *Aspergillus* were the most prevalent fungus in the atmosphere of Karachi and was represented by thirteen species, mainly *Aspergillus niger*, *A. flavus* and *A. fumigatus* (Table 1).

# Afzal and Mehdi: Fungal air spora

Table 1: Percent occurrence of fungal spores in the atmosphere

Fungal species	Altitude (m)		
	1.5	5	10
<i>Alternaria alternata</i>	5.67	6.88	1.66
<i>A. brassicae</i>	0.00	0.215	0.00
<i>A. brassicola</i> (Schw.) Wiltshire	0.00	1.705	0.00
<i>A. citri</i> (Ellis and Pierce) Pierce	4.179	3.44	1.25
<i>A. solani</i> Soroner	7.164	7.526	2.5
<i>A. tenuissima</i> (Kunz ex pers.) Wiltshire	8.955	9.03	4.583
<i>Aspergillus candidus</i> (Link) Thom & Church	1.492	0.215	0.00
<i>A. flavus</i> Link	15.22	13.978	7.08
<i>A. fumigatus</i> Fres	4.477	3.44	1.666
<i>A. glaucus</i> Link	0.298	0.215	0.00
<i>A. nidulans</i> (Eidum) Went.	0.597	0.430	0.00
<i>A. niger</i> Von Tiegh	17.61	19.14	7.916
<i>A. niveus</i> (Var.) Blochwitz	0.298	1.075	0.00
<i>A. ochraceus</i> Wilhelm	0.298	0.215	0.00
<i>Aspergillus</i> sp.	0.597	0.430	0.416
<i>A. sulphureus</i> (Fres) Thom and Church	0.00	0.215	0.00
<i>A. sydowi</i> (Bain and Sart) Thom and Church	0.597	0.00	0.00
<i>A. terreus</i> Thom	1.492	0.430	0.416
<i>A. ustus</i>	0.00	0.00	0.833
<i>Aureobasidium pullulans</i> (Debary) Arnaud	0.895	0.215	0.833
<i>Chaetomium globosum</i> Kunze	0.298	0.00	0.00
<i>Cladosporium herbarum</i> Pers Link ex S.F.Gray	0.597	0.860	6.666
<i>C. oxysporum</i> Burk and Curd	0.00	0.215	0.833
<i>C. sphaerospermum</i> Penz	0.895	1.720	10.41
<i>Curvularia clavata</i> Jain	2.686	2.58	1.25
<i>C. lunata</i> (Wakker) Baedign	0.895	2.795	0.833
<i>Drechslera dematioides</i> Bubuk and Wroblewski	0.298	0.430	0.00
<i>Drechslera</i> state of <i>Cochliobolus nodulosus</i>	0.597	0.00	0.00
<i>D. poae</i> (Baudys) Shoemaker	2.985	1.720	1.25
<i>Drechslera</i> state of <i>Cochliobolus spicifer</i> Nelson	2.0.89	2.365	0.416
<i>Fusarium culmorum</i> (Smith) Sacc.	0.00	0.00	0.833
<i>F. equiseti</i> (Corda) Sacc.	0.00	0.00	0.833
<i>Fusarium longipes</i>	0.00	0.215	0.00
<i>F. oxysporum</i> Schlecht emend Snyder and Hans	0.298	0.215	1.250
<i>F. semitectum</i> Berk and Rav.	0.00	0.00	0.416
<i>F. solani</i> (Mart) Appel and Wollenw emend Snyder and Hans	0.597	1.505	9.58
<i>Mucor mucedo</i> Michelo ex Fries	0.895	3.655	2.50
<i>Mucor</i> sp.	1.194	0.430	1.666
<i>Mycelia sterilia</i>	1.194	0.00	0.00
<i>Penicillium notatum</i> Westling	2.89	1.075	11.25
<i>P. brefeldianum</i> Dodge	1.79	1.290	9.583
<i>Penicillium</i> sp.	0.00	0.00	0.416
<i>Phoma</i> sp.	0.895	0.215	0.00
<i>Puccinia recondita</i>	1.194	1.505	0.416
<i>Rhizopus nigricans</i> Ehrenb	0.00	0.215	1.25
<i>R. stolonifer</i> (Ehrenberg ex Fries) Vuillemin	1.49	3.22	1.25
<i>Saccharomyces cerevisiae</i> Meyer	1.791	1.935	3.333
<i>Scopulariopsis</i> sp.	0.895	1.075	0.416
<i>Stemphylium</i> sp.	1.194	0.215	0.00
<i>Syncephalastrum racemosum</i> Cohn ex Schroter	0.895	1.075	3.336
<i>Trichoderma viride</i> Pers.	0.597	0.860	0.00
<i>Ucladium</i> sp.	0.298	0.645	0.00
<i>Ustilago tritici</i>	1.194	0.430	0.416

The species of *Alternaria* was the second most widespread fungus especially at low and middle altitude. However, it was also recorded at high altitude relatively in low numbers. Species of *Penicillium* were 8.4% and species of *Cladosporium* were 8.3% among the total fungal count during the year 1998-99. The possible reason for the variation in occurrence of the fungal spores may be related to the weight of spores, gravitational force and the velocity of wind (Tan *et al.*, 1992). Interestingly, *Cladosporium* spp., *Fusarium* spp. and *Penicillium* spp. were comparatively more in spore counts at high altitude as compared with low and middle altitudes (Table 1). The most frequently reported fungus *Aspergillus* from different regions of the world was also the most prevalent fungus during our investigation. Tilak (1990) reported *Aspergillus* to be common in atmosphere. Tan *et al.* (1992) stated that *Aspergillus* is a common air-borne fungus at 40m height in Singapore. Ebner *et al.* (1992) found *Aspergillus* as the most occurring fungus from atmosphere on plates. Earlier Ebner (1989) had reported that *Aspergillus* may play a role in indoor and *Penicillium* in outdoor allergy causation.

Similarly Pasanen *et al.* (1991), Simeray *et al.* (1995), Farifax *et al.* (1999) and Sanches *et al.* (1999) reported *Aspergillus* as an airborne fungal allergen. *Alternaria* was found as the second largest spp., which has also been reported by Tan *et al.* (1992) as a next most abundant fungal genus (16.71%) from Singapore. Kumar (1982) reported *Alternaria alternata* as most prevalent species in India. In other reports, abundance of *Alternaria* spp., is the second to that of *Cladosporium* spp., (Frey and Durie, 1962; Meyer *et al.*, 1983; Palmas and Cosentino, 1990). Ebner *et al.* (1992) reported *Cladosporium* as the most dominant genus. Marchisio *et al.* (1992) reported *Penicillium* from Turin Italy as the most frequent genus. Simeray (1995) reported *Cladosporium*, *Penicillium* and *Aspergillus* as the most frequent fungi in France. Fungal spores are disseminated by air current and in general large sized spores were comparatively found at lower altitude and smaller sized spores generally occurred at high altitude. On an average, *A. alternata* spores are 20-63µm long and 9-18µm wide (Ellis, 1971) whereas *Aspergillus niger* spores are 5.5-8µm (Thom and Raper, 1945) and *Penicillium notatum* conidia is

3-3.5 $\mu$ m in length (Raper and Thom, 1949). This shows that size and weight of the spores are extraordinarily important in distribution at different altitudes. This view has been supported by Tan *et al.* (1992), who reported that conidia of *Penicillium lapidosum* are smaller (2.0-2.5 $\mu$ m) and capable to reach high altitude. Since spores of *Alternaria* usually are of larger size therefore it may not have been easily blown to a higher level of the atmosphere by air current.

Simeray *et al.* (1995) reported 40 genera of fungi from the atmosphere of bake houses in which the most frequently occurring fungi were species of *Cladosporium*, *Penicillium* and *Aspergillus*.

During winter season spores of *Puccinia recondita*, *Ustilago tritici* which cause rust and smut disease of wheat were respectively trapped as air-borne spores. The detection of rust and smut spores in the atmosphere at the time of wheat sowing can be used in plants disease forecasting system for the control of rust and smut diseases. Environment plays a key role on the prevalence and distribution at various heights of air spora. Relative humidity, temperature and wind velocity are among the environmental factors affecting the distribution of spores in the atmosphere.

The atmosphere of Karachi city is laden with particulate matters containing soil, sand and dust impregnated with toxic chemicals and toxic gases emitted from vehicles and industries and released from degrading effluents in addition to the air spora together causing health problems (allergies and skin diseases) which are on increase posing more risk to human health with the increase in pollution day by day. The impact of toxic chemicals and gases on human health is required to be estimated which is beyond the scope of this work, however, the air-spores in the atmosphere of Karachi city is being presented for the first time which may be of interest to doctors dealing with dermal and respiratory diseases caused by fungal allergens and to those engaged in the protection of plants from the wind blown spores of fungi.

In short, the fungal spores are important in allergy provoking factors, which always remain present in the atmosphere in more or less concentration at different altitudes. Although, their quantity and quality depend upon different factors and meteorological parameters, but concentration is greater at medium (5 m height) altitude as compared to others with reference to atmosphere of metropolitan city of Karachi.

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## References

Agashe, S.N. and Alfadil, 1989. Gravity sampling device. Grana, 28: 97-104.  
 Al-Doory, 1984. Air-borne fungi. In Mould Allergy (Eds.): Al-Doory & J.F. Domson, pp: 27-40.  
 Angullo-Romero, J., M.A. Mediavilla and V.E. Dominguez, 1999. Conidia of *Alternaria* in the atmosphere of Cordoba, Spain, in relation to meteorological parameters. Int. J. Biometeorol., 43: 45-49.  
 Bapat, B.N., 1994. Pragmatic approach to atrophy. Current Trends in Life Sciences. In Advances in Mycology and Aerobiology, pp: 355-65.  
 Baratawidjaja, I.R., P.P. Baratawidjaja, A. Darwis, H.I. Soo, T.C. Fook, W.L. Bee and K.G. Baratawidjaja, 1999. Prevalence of allergic sensitization to regional inhalants among allergic patients in Jakarta, Indonesia, Mar., 17: 9-12.  
 Chapman, J.A., 1999. Update on air-borne mould allergy. Allergy Asthma Proc., 20: 289-292.

deCastro, C.C., G. Bernard, Y. Ygaki, M.Y. Shikanai and G.G. Cerri 1999. MRI of head and neck paracoccidioidomycosis. Br. J. Radiol., 72: 717-22.  
 Ebner, M.R., K. Haselwandter and A. Frank, 1989. Seasonal fluctuation of airborne fungal allergens. Mycol. Res., 91: 170-176.  
 Ebner, M.R., K. Haselwandter and A. Frank, 1992. Indoor and outdoor fungal allergens at low and high altitudes, alpine environment. Mycol. Res., 96: 117-124.  
 Elewski, B.E., 2000. *Tinea capitis*, a current perspective. J. Am. Acad. Dermatol., 42: 1-24.  
 Ellis, M.B., 1971. Dematiaceous Hyphomycetes. Commonwealth Mycol. Institute, Kew, Surrey, England.  
 Farifax, A.J., V. David and G. Douce, 1999. Laryngeal aspergillosis following high dose inhaled fluticasone therapy for asthma. Thorax, Sept., 54: 860-861.  
 Frey, D. and E.B. Durie, 1962. Estimation of air fungal spores: A comparison of slide and culture method. Mycopathologia, 16-17: 295-303.  
 Ghafoor, A. and S.A.J. Khan, 1976. List of diseases of economic plants in Pakistan. Govt. Publication, Ministry of Food and Agriculture, Islamabad.  
 Gregory, P.H., 1973. The Microbiology of the Atmosphere. Leonard Hill, London.  
 Hirschmann, J.V. and G.J. Raugi, 2000. Pustular tinea pedis. J. Am. Acad. Dermatol., 42: 132-3.  
 Kim, C.K., 1982. Improved method for rice blast forecasting. Korean J. Pl. Prot., 21: 19-22.  
 Kumar, R., 1982. Aerospora in pine forest in India. Grana, 21: 179-181.  
 Marchisio, V.F., C. Mozenzo and R. Caramello, 1992. Preliminary survey of airborne fungal propagules in Turin, Italy. Mycol. Res., 96: 535-541.  
 Meyer, G.H., H.E. Prince and W.J. Raymer, 1983. Airborne Fungi, a Survey. Ann. of Allergy, 51: 26-29.  
 Mughal, F.H., 2002. Impact of air pollutants. Dawn, Feb. 20, 2002, Karachi, Pakistan.  
 Munuera, M., M. Giner, G.J.S. Carrion and J. Selles, 1998. Incidence of *Alternaria* spores in the atmosphere of Murcia, (S.E. Spain), seasonal, monthly and intra diurnal variation. J. Investig. Allergol. Clin. Immunol., 8: 304-8.  
 Palmas, F. and S. Cosentino, 1990. Comparison between fungal air spores concentration at two different sites in the south of Sardinia. Grana, 29: 87-95.  
 Pasanen, A.I., P. Kallikowski, P. Pasanen, M.J. Jantunen, A. Levanen, 1991. Laboratory studies on the relation between fungal growth and atmospheric temperature and humidity. Env. Int., 17: 225-228.  
 Raper, K.B. and C. Thom, 1949. Manual of the Penicillia the William and Wilkins Co., Baltimore, USA.  
 Sanches, S.B., R.M. Gallardo, J.A. Navarro Chavarria and M.I. Cabrera Munoz, 1999. Allergic fungal sinusitis, recent developments. Rev. of Allergy, 46: 145-50.  
 Simeray, J., D. Mandin and J. Chaumont, 1995. Variation in distribution of fungal spores in the atmosphere of bake house. Impact on studies of allergies. Grana, 34: 269-274.  
 Thom, C. and K.B. Raper, 1945. Manual of Aspergilli. Williams and Wilkins Co., Baltimore, USA.  
 Tan, T.K., T.S. Teo, H. Tan, B.W. Lee and A. Chong, 1992. Variation in tropical air spora in Singapore. Mycol. Res., 96: 221-224.  
 Tilak, S.T., 1990. Airborne spores as bioindicators. Perspective in Mycological Research-2, Prof. G.P. Agarwall Festschrift, 227-236.  
 Vanderbergh, M.F., P.E. Verweij and A. Voss, 1999. Epidemiology of nosocomial fungal infection invasive aspergillosis and the environment. Diagn. Microbiol. Infect. Dis., 34: 221-7.