Airborne biological materials: Their identification, origin and impact on environment

M.N. ABUBACKER AND A. AMATUSSALAM

Asian Journal of Environmental Science, (June, 2011) Vol. 6 No. 1 : 1 -11

SUMMARY

authors' affiliations Correspondence to :

See end of the article for

M.N.ABUBACKER Department of Botany, National College, TIRUCHIR APPALLI, (T.N.)INDIA abubacker_nct@yahoo. com

Key words :

Biological

grains

Received:

Accepted :

Allergy, Asthma,

materials, Fungal

spores, Pollen

Identification, origin and impact on environment of airborne fungal spores, pollen grains and other biological materials of Tiruchirappalli, Tamilnadu was studied for a period of one year from January to December 2009 at 10 m height from the ground level using vertical cylinders as trap. Alternaria padwickii, Aspergillus niger, Cladosporium herbarum, Curvularia lunata, Helminthosporium oryzae and Nigrospora oryzae were major concentrated fungal spores. Azadirachta indica, Casuarina equisetifolia, Cocos nucifera, Eucalyptus globules, Grass spp. including Oryza sativa, Parthenium hysterophorus, Typha anguastata were the major contributors of pollen types. The other biological materials included, epidermal cells, epidermal hairs, protozoan cyst, mites and thrips. The major concentration of fungal spores, pollen and mites will lead to allergy and trigger attacks of asthma.

M.N. Abubacker and Amatussalam, A. (2011). Airborne biological materials: Their identification, origin and impact on environment. Asian J. Environ. Sci., 6(1): 1-11.

C tudy of airborne biological materials is Nown as aerobiology (Meier *et al.*, 1933). This field is related to the study of fungal spores, pollen grains, bacteria and other biological materials present in the air. The scope of it is now well known to carry a heterogenous population of an array of bio-particles (Singh et al., 2005). These bio-particles vary in origin, size and structural complexity and were called to constitute airspora (Studenkin and Sokolova, 1977; Nilsson, 1992).

Several aerobiological studies were conducted by many workers (Gregory, 1973, 1983; Norman and Lichtenstein, 1986; Rao et al., 1995; Singh, 1998; Dales et al., 2000; Potdar et al., 2000; Anderson et al., 2001; Sears et al., 2006; Amato et al., 2007; Emberlin, 2008). The purpose of this report is to provide a comprehensive picture of airspora for the clinical aspects because the inhalation of spores of different species results in different health effects from allergy to Aspergillosis (Tobin et al., 1987; Miller, 1990).

MATERIALS AND METHODS

December, 2010 January, 2011

Vertical cylinders of 0.5 cm diameter (Fig. 1) were used as trap (Ramalingam, 1968). A cellophane strip is stuck on the cylinder and coated with glycerine jelly. Exposed it daily round the clock for a period of one year at 10 m height at three different places in Tiruchirappalli to know the atmospheric concentration of fungal spores, pollen grains and other biological materials and their average was expressed in number (cm³). The exposed cellophane strips (2x2 cm) on the cylinders were prepared for microscopy (Ramalingam, 1968). An area of 0.15 cm² was scanned from each exposure for the biological material counts. The fungal spore types were identified and confirmed with the literature of eminent biologists (Ellis, 1971; Subramanian, 1971; Ainsworth et al., 1973; Gregory, 1973; Gilman, 1975; Tilak, 1982). The pollen types were identified and confirmed with the help of literature of some other biologists (Erdman, 1969; Tilak, 1982; Nair et al., 1986). The identification was confirmed with the help of reference slides of pollen collected directory from the plants. The pollen grains were mounted in safranin stained glycerine jelly.

RESULTS AND DISCUSSION

Table 1, 2 and 3 illustrate the list of atmospheric concentration of fungal spores, pollen grains and other biological materials respectively. Altogether 24 fungal spores, 15 pollen types and other biological materials like

ి జుతు తో : ్లాజితలాయి అొ జింారికు పారాగిత	333	2.00 002 C		- y 2.009	ിരതേ സംഗ	. 2009								
	10 0000 M		NET.	Vian	N.E.Y			ACE	çõg	06.	AO/			Dencem 220
Aldermanta kongipes	320	.26	175	08,	.12	. 20	08	82,	60.	./8	09	99	997.	2.119
Alternaria padvickái	/ 80	290	016	2120	260	07.	200	. 60	202	22.0	2:7	32.0	3062	
Vscosjons	. 2.0	00	£	16	20	26	12.	10/	2.6	61.	ary .	097	93/	
Aspensilaus nager	20	82.0	087.	620	612.	520	126	180	62.0	126	080;	000.	9052	1.5.9.
Camptonneris albiviae	12.	20		0,	0	0		26		30		18	dillo	0.5.3
Cercospora personala	. 26	00 20 20	616	26.	98 8		20	16	30	24	25	0	928	211.
Chaeiomaiam globosum		12.	20	0	0	0	0	18	62,	.32	97.	18.		. 395
Cladosporium herbarum	. 620	.880	0.2.	8/8	812	612	686	92.0	. 683.	001.	. 82.0	087.	. 5378	28.836
Curvularia lunata	620	182	366	125	017		61.1	680	612	221.	1 66	5.0	6328	398.1
Drechslera xeicola	12		26	32,	12.		22.	12.	07		62	15	156	0.855
ห่านระเทราสหล อวสุระทุรอาหาสหล	83,	62,	00	12.	32,	99	200			1.9	20	52,	632,	\$22.
li leplosporella so.	98.	2:0	82	Q.,	16	11	1875	. 20	82,	06	97.	21.	3.10	2,569
llebminthosportum oryzae	12.0	360	280	12 J	00.		. 82	.52	.26	2/0	3/6	1 82.	3088	067.9
l ledminthosportum carbonam	26	12.	28	0,	28	52	21	22.		28	32.	36	398	01/10
hAyoosphaerelka muzicoka	32,	÷.	2.2.	22.	14 v.	0	G		26	12.	32.	26	2.9.	0.575
Nigerospora oryzae		370	2.82.	312	000	.62.	. 82	0°	2/0	2.82.	320	.82,	2,992.	5.6.0
^{kr} ericonia circinala	2.2	16.		2.20	. 7.	160	2:0	. 87	97.	1.28	<i></i>	2.0	2005	3.752.
Plerospora herbarum	1		26						ŝ	20			. S¢	0.281
Pringskennia 55.	22		¢,	225				26				16.	1 22	" Sol + W
P-seudotoruka %3.	15	61	19,	20	1.5	32				36		12	/ 03	55\$. M
Sphaeropsis tumefascience	32,	99	t.				99	32	2.6	12	20	26		0,598
Sporormiella megalospora	2				ф.,		er.		0	4	25	20	56.	0.23/
T'etrapola s.z.				20	20		18 N			4 AN			<i>(</i> 9.	1.02.0
1 richtottis paävickii	36	3.0	312	7.87	32/	120	. 26	2.	.02	66	83	6	2355	
```D`.E.`	63.11	5612	1592	359"	3116	32/8	2.8/0	3367	135	5059	5526	52,67	53333	ሰብሰና ግጥ " ህህዝ ህህ ፣

[Asian J. Environ. Sci. (June, 2011) Vol. 6 (1) ]

•HIND INSTITUTE OF SCIENCE AND TECHNOLOGY•

ົ. ຮັງ ດ 2: ີດເຈັດດູດເວັດ ທີ່ ຮັດກາງເປັນຄ	~~~~~~~~		E.N 2009		- 2,000									
		. co.	W.g.	A 37	W.EY			Aug	Sco	0%	NOV.			
Ailonahaas excelso	60	21:	52,	0°.	0	9	()) ())	0	0		3	12.	901	
Amaramihus spinosus	63	6% %	12.	16	0		36	S./.,	200.	V0 120	00	66	669	2,303
Aradirachta indica	0/	28	8112	. 226	160	. 20	20	0	0	Ó	63	0	2.821	9.36.
Casuarina equistifolia	. 82	620	22	Ő	0	0	9	0	0	6) 20	52	120	2.85	8161.
Cocos nucifera	125	380	3/0	022	210	2.0	120	530	3/0	02 :	0/. ·	280	1 095	3.511
lincalyptus globulus	\$2		2,62,	0°		0		ŝ	20	16			501	2.986
Mamerifera indica	386		2.0	~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~	18	0	0	0	0	Ô	15	91.	186.	156
IPoa saliva	0/97	@Z	820	119	280	60	00° .	326	130	61.1	980 780	210	1632	25.198
)/° arthrenitann		586	320	2.82.	2.52.	0.1	230	520	911	336	12.	636	5.35	:1.:85
hysterophoras										9 19			1	9711 W
Primecolobram dulce	67	0£	mr.	(M)	3	$n_{\mu}$	3	(N)-	5	20%	20 M	10	1.1.1	0.11 2.
Prosopis juliflora		00			30	32	01		0	0	£.'		503	
Ricinas communis		20	66			20	3	0	ŝ	10	00 90 , :		061.	2,336
11 annarindus indicus	0/:	. 20	0	0	0	01	\$2,	ŝ	0		00	\$\$ \$\$ \$\$ \$\$ \$\$	513	
Terminalia calappa	0	0		12	98 M	0	0	17 17	0	23	30	26	23.	0,165
71 spina americata	516	626	1 20	320	00 	0	0	69	0	0	285	210	1.1.9%	8,873
	5328	1520	3682	3/38	600	265	200.	997.	. 338	289.	20.2	3191	30.10	(TATAT "ATAT")
ి జూల 3: ెడిడలాలు ది జి.ాంకాలు	20:30 0.C 3	2	S CYS2 2			S 2 2	y 2009 -	0000-000	2,009					
	a barrow	. C.D.	W.E.	N.S.	N.E.Y			Aug	eas	. DC.	NOX.	)cc:	'0'.£.'	Teresen age
lipičarmel adls (monosol)	80	ζ.,	16.	. 20	61.	200	2.0	06.		100	36	20	188	008.
. Inclamment act & (also )	26	31	30	26	12.	36	18	201	20		22	26	358	151
Southerness (mensues in the s	4.	10° -			200	16	20		м. У		5.		587	2.54
A and the second of the second s	16	140	26	1 3	241	08	13			30		30	1.51	
	20			2.6	12.	60			22.		20		300	50-1
and comment while (S. S. S. S. S.		5.	\$	12		2.8	3		20		20	20		3.79
insoot seelo (moed)	13.	$\mathcal{C}_{\mathcal{C}}$	00	15	23.7		000	61.	0°	00	30	11	Y	- 12 CV -
iiiisoo( sosio (iiiosi)		82	. 82	91.	2.0	. 32		0	82	0/2	00	92	. / 88	20.09
3.00/02 cysts	28	12.	36	20 77	.61.	80	12	16	32.			22.	112	1.5.5
Dermanyasaus s.o.	32.	20	16	131.	0	62,	12	36	20	12.	36	20	208	6.86
Cievendare en		197 - S.		S.							0°.		57.	

[Asian J. Environ. Sci. (June, 2011) Vol. 6 (1) ]

•HIND INSTITUTE OF SCIENCE AND TECHNOLOGY•

trichomes, epidermal peelings, mites thrips and cysts were identified from the exposed vertical cylinders (Histograms 1-3).

#### **Fungal spores:**

The maximum concentration of fungal spores of

28.83% was recorded for *Cladosporium herbarum* followed by *Aspergillus niger* 16.97%, *Curvularia lunata* 11.86%, *Helminthosporium oryzae* 5.79%, *Alternaria padwickii* 5.74%, *Nigrospora oryzae* 5.61%, *Trichoc*onis padwickii 4.41%, and Periconia circinata 3.75%. All these spores were found throughout the year





[Asian J. Environ. Sci. (June, 2011) Vol. 6 (1) ]





and all these spores showed seasonal distribution peaks. The concentration of spores in air are quite variable according to the source of spores and infiltration rate at the time of sampling (Anonymous, 1988; Hunter *et al.* 1988). C. herbarum exhibited peak during September to March and A. niger exhibited peak during October to January. All types of spores except Camptomeris albizziae, Chaetomium globosum, Mycosphaerella musicola and Sporormiella megalospora occured throughout the year. The fungal spore in maximum concentrations dispersed in microdroplets of aerosols penetrate deep into the lungs which will lead to allergy and trigger attacks of asthma (Dales *et al.*, 2000; Anderson, 2001; Geiser *et al.*, 2000).

## Identification features of fungal spores

Alternaria longipes:

Conidiophores solitrary, simple, septate, olivaceous brown, smooth. Conidia ovoid, short, cylindrical beak, smooth, with 2 or 3 transverse and longitudinal septa, 40-50 x 30-35  $\mu$ m. It originates from damp ceiling paper, cardboard and textiles (Fig. 2a: a). causes leaf spot disease in tobacco.

#### Alternaria padwickii:

Conidia ovoid, cylindrical beak with scars, occurs in chain, 4 or 5 transverse and longitudinal septa, 75-100 15-25  $\mu$ m. Originates from decaying vegetable and plant debris (Fig. 2a: b). causes leaf spot disease in paddy.

#### Ascospores:

Spores are globose, unicellular, occur in groups, spore wall smooth 10-12  $\mu$ m. Originate from decaying vegetable and fruits (Fig. 2a: c). The fungus producing such type spores causes decay and degradation of vegetables and fruits.

#### Aspergillus niger:

Spores are very small, round, unicellular, echinate 3-5  $\mu$ m. Originates from decaying food stuff of different kinds and decaying paper and wood (Fig. 2a: d). Causes allergy and Aspergillosis on excessive amount of inhalation.

#### Camptomeris albizziae:

Conidia long, cylindrical, septa 4 or 5, cylindrical beak



with scar.  $180-200 \times 30-40 \,\mu\text{m}$ . Originates from decaying plant debris (Fig. 2a: e) causes leaf spot disease in crop plants.

#### Cerospora personata:

Conidia long, cylindrical, septa present 5 or 6, cylindrical beak with scar.  $200-210 \times 25-35 \mu m$ . Originates from burnt soil, infected leaf debris (Fig. 2a: f), causes leaf spot disease in groundnut plant.

#### Chaetomium globosum:

Conidia oval, unicellular 15-20 mm. Originates from

decaying fruits (Fig. 2a: g). Occurs on damp paper, clothes, decaying materials.

#### Cladosporium herbarum:

Conidia oblong with prominent scar, occur in branched chains, some conidia are septate, 8-12  $\mu$ m, occurs on damp paper, cloth, decaying fruits and vegetable (Fig. 2a: h). Causes deacay and degradation of fruits and vegetable.

#### Curvularia lunata:

Conidia broad, septate 2-4, second septa broad, conidia typically bent with prominent scar,  $80-90 \times 40-50 \mu m$ . Originates from decaying food stuff, soil debris (Fig. 2a: i). Causes leaf spot infection in many crop plants.

#### Drechslera zeicola:

Conidia, long, cylindrical, pseudosepta present 6-10,  $140-150 \times 35-45$  mm, originates from infected paddy straw and decaying soil debris (Fig. 2a: j). Causes leaf spot infection in paddy and allied crops.

#### Fusarium oxysporum:

Macroconidia fusiform, septate, 3-6, 30-40 x 5-8  $\mu$ m, originates from soil, infected leaf debris, decaying fruits (Fig. 2a: k). Causes wilt disease in many crops including banana.

#### Heplosporella sp.:

Conidia oval, single celled with prominent scar. 25-30 x 10-12  $\mu$ m. Originates from decaying vegetable (Fig. 2a: 1). Causes decay and degradation of vegetable and crop residues.

#### Helminthosporium oryzae:

Conidia long and cylindrical, beak smooth, septa 6-8, with basal scar 320-340 x 50-60  $\mu$ m. Originates from soil, infected plant debris like grass, paddy and maize (Fig. 2b: m). Causes leaf spot disease in paddy and allied crops.

#### Helminthosporium carbonum:

Conidia long slightly bent, cylindrical, beak smooth, septa 7-9, with basal scar  $360-380 \times 60-70 \,\mu\text{m}$ . Originates from soil, infected plant debris like grass, paddy and maize (Fig. 2b: n). Causes leaf spot in paddy, maize and allied crops.

#### Mycosphaerella musicola:

Conidia oblong, septate, 3 celled, middle cell larger than other two cells, with basal scar  $70-80 \times 30-40 \ \mu m$ . Originates from decaying fruits (Fig. 2b: o). Causes



degradation of organic debris.

#### Nigrospora oryzae:

Conidia, single celled oval to globose, opaque, black, 40-50  $\mu$ m. Originates from soil and infected grass and paddy (Fig. 2b: p). Causes leaf spot infection in paddy and other grasses.

#### Periconia circinata:

Conidia globose, 1 celled, dark brown echinulate 20-25 mm. Originates from soil, decaying vegetable and fruits (Fig. 2b: q). Causes degradation of organic materials.

#### Pleospora herbarum:

Conidia oblong, septata 2-4, cylindrical beak with basal scar. 130-150 65-75  $\mu$ m. Originates from soil, decaying plant debris (Fig. 2b: r). Causes decay and degradation of plant debris.

#### Pringshemia sp.:

Conidia oblong, pseudosepta present 2-3, cylindrical beak with prominent scar 140-150 x 50-60  $\mu$ m. Originates from soil and decaying plant debris (Fig. 2b: s). Causes decay and degradation of organic debris.

#### Pseudotorulla sp.:

Conidia 4 celled to 8 celled chains, branched, each conidia is globose 90-100 x 10-12  $\mu$ m (8 celled chain). Originates from decaying fruits (Fig. 2b: t). Causes decay of organic debris.

#### Sphaeropsis tumefascience:

Conidia large, 1 celled oval to globose  $60-80 \times 50-60$  µm. Originates from soil and decaying plant debris (Fig. 2b: u). Causes degradation of organic matter.

#### Sporomiella megalospora:

Ascospores very dark, 3-septate, much constricted at septa  $300-350 \times 50-60 \,\mu\text{m}$ , middle cells cylindrical, end cells slightly longer and somewhat conical. Originates from decaying vegetable and horse dung (Fig. 2b: v). Bring about degradation process of organic materials.

#### Tetrapola sp.:

Conidia cylindrical with 4 radiating septate appendage. Conidia septate 4 celled,  $110-120 \times 60-70 \,\mu\text{m}$  appendage 90-100 mm long. Originates from decaying aquatic plants (Fig. 2b: w). Causes decay of aquatic plant debris.

#### Trichoconis padwickii:

Conidia 6-9 celled with a prominent long trichome like back. Apical cell somewhat conical second and third cells are broader than other cells,  $220-240 \times 20-60 \mu m$ , trichome 180-220  $\mu m$  long. Originates from soil, decaying paddy straw (Fig. 2b: *x*). Causes leaf spot infection to paddy and allied crops.

#### **Pollen grains:**

The maximum concentration of pollen grains of 25.49% was recorded for *Poa sativa*, followed by 17.18% for *Parthenium hysterophorus*, 13.57% for *Cocos nucifera*, 9.36% for *Azadirachta indica*, 8.87% for *Typha angustata*, 7.24% for *Casuarina equisetifolia*,

and 4.15% for *Mangifera indica*. *C. nucifera*, *P. sativa* and *P. hysterophorus* pollen grains were abundant and found throughout the year and all these pollen showed seasonal distribution peaks which attributed to the effects of climate (Amato *et al.*, 2007), whereas low amount of pollen source are *Ailanthus excels*, *Amaranthus spinosus*, *Eucalyptus globulus*, *Pithecolobium dulce*, *Prosopis juliflora*, *Ricinus communis*, *Tamarindus indicus* and *Terminalia catappa*. The abundant pollen such as *P. sativa*, *P. hysterophorus*, *C. nucifera*, *A. indica*, *T. angustata* and *equisetifolia* derived from dispersed microdroplets represent a possible mechanism by which allergy/pollinosis can trigger attacks of asthma (Menz *et al.*, 2006; Pham *et al.*, 2006; Amato *et al.*, 2007).

#### Identification features of pollen grains:

Ailanthus excelsa (Simaroubaceae): Pollen grains 3-zonocolporate, oblate spheroidal. Grain size 90-95  $\mu$ m. Colpus long, streak-like, exine thicker at poles. Exine surface finely reticulate (Fig. 3: a).

#### Amaranthus spinosus (Amaranthaceae):

Pollen grains multiporate, spherical. Grain size 80-90 mm. Pore membrane crustate. Exine surface granulate (Fig. 3: b).

#### Azadirachta indica (Meliaceae):

Pollen grains, 4-zonocolporate, prolate spheroidal. Grain size 135-145  $\mu$ m. Exine thick, ectexine thicker than endexine. Exine tectate, surface psilate and punctate (Fig. 3: c).

#### Casuarina equisetifolia (Casurinaceae):

Pollen grains 3-zonoporate, oblate, grain size 100-110  $\mu$ m, amb triangular with slightly convex sides. Pores aspidate. Exine surface psilate (Fig. 3: d).

#### Cocos nucifera (Arecaceae):

Pollen grains ellipsoidal, 1-colpate. Exine thick and conspicuously folded, surface psilate. Grain size 185-195 x 70-75  $\mu$ m (Fig. 3: e).

#### Eucalyptus globuslus (Myrtaceae):

Pollen grains amb triangular, 3-zynocolporate, exine surface granulate. Grain size 90-100  $\mu$ m (Fig. 3: f).

#### Mangifera indica (Anacardiaceae):

Pollen grains 3-zonocolporate, suboblate. Grains size 85-95  $\mu$ m. Endocolpium elongate. Exine surface striate (Fig. 3: g).



#### Poa sativa (Poaceae):

Pollen grains monoporate, pore circular, annulate, spheroidal. Grain size 185-195  $\mu$ m. Exine surface psilate (Fig. 3: h).

#### Parthenium hysterophorus (Asteraceae):

Pollen grains 3-zonocolporate, suboblate. Exine surface spinate tip acute. Grain size 40-50  $\mu$ m (Fig. 3: i).

#### Pithecolobium dulce (Mimosoideaceae):

Individual pollen grains. Unite to form a polyad, spheroidal, polyad 16 celled. Exine thicker at outer margin thinner towards inner margin. Exine surface granulate. Grain size 170-190  $\mu$ m (Fig. 3: j).

#### Prosopis juliflora (Mimosoideaceae):

Pollen grains 3-zonocolporate, Endocolpium lalongate. Exine surface striate. Grain size 150-160 m (Fig. 3: k).

8

#### Ricinus communis (Euphorbiaceae):

Pollen grains 3-zonocolporate, suboblate, endocolpium la-longate, colpus ends pointed, exine surface finely reticulate. Grain size 190-200  $\mu$ m (Fig. 3: 1).

#### Tamarindus indica (Caesalpiniaceae):

Pollen grains 3-zonocolporate, endocolpium lalongate, exine surface regulate. Grain size 100-110  $\mu$ m (Fig. 3: m).

#### Terminalia catappa (Combretaceae):

Pollen grains 3-zonocolporate with 3 alternating pseudocolpi, oblate spheroidal, endocolpium circular. Grain size. 140-150  $\mu$ m (Fig. 3: n).

#### Typha angustata (Typhaceae):

Pollen grains 1-porate, spheroidal, exine surface reticulate. Grain size 130-140 mm (Fig. 3: 0).

# Epidermal cells, stigma, epidermal hairs, insect scales and protozoan cysts:

Epidermal cells of monocot, dicot plants, stigmatic part of flower, epidermal hairs, trichome, stellate hairs of plant origin, insect scales of various size and shape and protozoan cysts, mites and thrips were of common occurrence in air (Fig. 4 a-i). These will lead to the relative risks of development of allergy and asthma (Sears *et al.*, 2006).

#### Characteristic features of mites and thrips:

Dermanyssus sp. (Dermanyssidae):

The chicken mites, pest of poultry, long oval body the cephalothorax long, oval body, require blood meal and quite resistant to starvation. 450-500  $\mu$ m in length 120-130  $\mu$ m broad (Fig. 4: j) (Singh and Sachan, 2004).

#### Glyciphagus sp.:

The flour mite, cylindrical cephalothorx and broad abdomen  $360-380 \times 90-110 \,\mu\text{m}$  (Fig. 4: *k*) (Wallis, 2002).

#### Pyemotes sp. (Pediculoididae):

The straw itch mite, causative agent of hay or grain itch, attack human in hot weather and causes intense itch, long tubular body. 210-220 x 65-85  $\mu$ m (Fig. 4:1) (Singh and Sachan, 2004).

#### Tetranychus sp. (Tetranychidae):

The two-spotted spider mite, suck the plant sap. oval body 130-140 x 80-90  $\mu$ m (Fig. 4: m) (Singh and Sachan, 2004).



#### Thrips sp.:

The flower thrips with distinct head, thorax and abdomen with narrow end.  $230-240 \times 70-80 \ \mu m$  (Fig. 4: n) (Wallis, 2002).

#### Tyroglyphus sp.:

The common cheese mite, shows a division into two parts the cephalothorax and abdomen measures 330-350 mm  $\times$  120-140  $\mu$ m (Fig. 4: o) (Wallis, 2002).

#### **Conclusions:**

The results of this study demonstrate that differences in ambient concentration of biological materials and meteorological conditions appear to influence allergic rhinitis, conjunctivitis and allergic asthma to the population.

#### Acknowledgements:

The authors thank DST-FIST, Government of India, New Delhi for providing the infrastructure facilities for the Departments of Botany and Chemistry, National College, Tiruchirappalli. The authors also thank Sh. K. Ragunathan, Secretary and Dr. K. Anbarasu, Principal, National College for their encouragement.

#### Authors' affiliations:

A. AMATUSSALAM, Department of Chemistry, National College, TIRUCHIRAPALLI (T.N.) INDIA

#### References

**Ainsworth, G. C.**, Sparrow, Frederick K. and Sussman, Alfred, S. (1973). *The Fungi: An advanced treatise*. Academic Press, New York.

Amato, G. D., Spicksma, F., Th. M., Liccardi, G., Jager, S., Russo, M., Konton-Fili, K., Nikkels, H., Wathrich, B. and Bonini, S. (2007). Pollen-related allergy in Europe. *EAACI*., **53**: 567-578.

Anderson, W., Prescott, G.J., Packham, S., Mullins, J., Brookes, M. and Seaton, A. (2001). Asthma admissions and thunderstorms: A study of pollen, fungal spores, rainfall and ozone. *Q. J. Med.*, **94**: 429-433.

**Anonymous** (1988). *Determination of fungal propagules in indoor air*. Canada Mortgage and Housing Corporation, Ottawa, Canada.

**Dales, R. E.**, Cakmak, S., Burnett, R. T., Judek, S., Coates, F. and Brook, J.R. (2000). Influence of ambient fungal spores on emergency visits for asthma to a Regional Children's Hospital. *AJRCCM*, **162**: 2087-2090.

**Ellis, M.B.** (1971). *Dematiaceous hypomycetes*. Commonwealth Mycological Institute, Kew, England.

Emberlin, J. (2008). The effects of patterns in climate and pollen abundance on allergy. *EAACI*, **49**: 15-20.

Erdman, G. (1969). *Handbook of Palynology: An introduction to the study of pollen grains and spores.* New York.

**Geiser, M.,** Leupin, N., Maye, I., Vinzenz Hof, I. M. and Gehr, P. (2000). Interaction of fungal spores with the lungs - Distribution and retention of inhaled puff ball (*Calvatia excipuliformis*) spores. *J. Allergy & Clinical Immunology*, **106**: 92-100.

Gilman, J. C. (1975). *A manual of soil fungi*. Revised Ed., Oxford IBH Publ. Co., New Delhi.

**Gregory, P. H.** (1973). *The microbiology of the atmosphere*. 2nd ed. Plant Science, Monograph, Leonard Hill Publ., London.

**Gregory, P. H.** (1983). *Aerobiology*. past, present and future Intl. Aerobiol. News Letter.

**Hunter, C. A.,** Grant, C., Flannigan, B. and Bravery, A. F. (1988). Mould in buildings: The air spora of domestic dwelling. *International Biodeterioration & Biodegradation*, **24**: 81-101. Meier, F. C., Stevenson, J. A. and Charles, V. K. (1933). Spores in the upper air. *Phytopathology*, **23**: 23.

Menz, G., Olecek, C. D., Scttonheit-Kenn, V., Ferreira, F., Moser, M., Schneider, T., Suter, M., Boltznitulescv, G., Ebner, C., Kraft, D. and Valenta, R. (2006). Serological and skin-test diagnosis of birch pollen allergy with recombinant Bet VI, The major birch pollen allergen. *Clinical & Experimental Allergy*, **26**: 50-60.

Miller, J.D. (1990). Fungi as contaminants in indoor air. *Pro. 5th Internat. Conf. on indoor air & climate, Toronto,* **5**: 51-64.

Nair, P. K. K., Joshi, A. P. and Gangal, S. V. (1986). *Airborne pollen, spores and other plant material of India:* A survey. C.S.I.R. Centre for Biochemicals, Delhi and National Botanical Research Institute, Lucknow.

Nilsson, S. (1992). Aerobiology: An interdisciplinary and limitless science. *Indian J. Aerobiol. (Spl. Vol.)*, pp. 23-27.

Norman, P. S. and Lichtenstein, L. M. (1986). The great debate: Immunology and asthma. *Clin. Allergy*, **16**: 269-271.

**Pham, N. H.**, Baldo, B. A. and Bass, D. J. (2006). Cypress pollen allergy. Identification of allergens and cross reactivity between divergent species. *Clinical & Experimental Allergy*, **24**: 558-565.

**Potdar, S. K.**, Nair, V. S., Chaudhary, S., Thomas, D. and Nair, L. N. (2000). In: *Vistas in Mycology and Plant Pathology* (Commemoration Volume), L. N. Nair (Ed.), Commonwealth Publishers, pp. 178-219.

**Ramalingam, A.** (1968). The construction and use of a simple air sample for routine aerobiological surveys. *Environ. Health*, **10**: 61-67.

**Rao, C.,** Burge, H. and Chang, J. (1995). Review of concentration, standards and guidelines for fungi in indoor air. In: Air and waste management association symposium on engineering solutions to indoor air quality problems. Raliegh, NC. Int.

Sears, M. R., Herbison, G. P., Holdaway, M. D., Hewitt, C. J., Flannery, E. M. and Silva, P. A. (2006). The relative risks of sensitivity to grass pollen, house dust mite and cat dander in the development of childhood asthma. *Clinical & Experimental Allergy*, **19**: 419-424.

Singh, Lokendra, Singh, V., Saima Perveen and Chopra, A. (2005). Isolation and identification of aeromycoflora of Mawana. *Biochem. Cell. Arch.*, **5**: 101-104.

**Singh, A. B.** (1998). Airborne fungi of allergenic significance in work environments. In: *Recent Trends in Mycoses*, pp. 9-17.

Singh, Rajendra and Sachan, G.C. (2004). *Elements of Entomology*. Rastogi Publications, Meerut, India.

**Studenkin, N.** and Sokolova, T. (1977). *Allergic disorders in children*. Mir Publishers, Moscow.

Subramanian, C. V. (1971). *Hyphomycetes*. Indian Council of Agricultural Research Publication, New Delhi.

Tilak, S.T. (1982). *Aerobiology*. Vaijayanthi Prakasan, Aurangabad.

**Tobin, R. S.,** Baranowski, E., Gilman, A. P., Kuiper-Goodman, T., Miller, J. D. and Giddings, M. (1987). Significance of fungi in indoor air. *CJPH*, **78**: S.1-32. **Wallis, T. E.** (2002). *Textbook of Pharmacognosy*. 5th ed., CBS Publishers and Distributors, New Delhi, India.

___