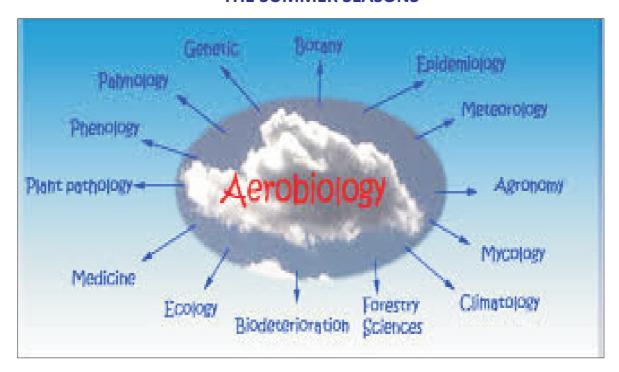
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# STUDIES IN PATHOGENIC AIRSPORA AND EPIDEMIOLOGY OF GROUNDNUT CROP AT SOLAPUR DURING THE SUMMER SEASONS







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#### **ABSTRACT:**

Aarchis hypogaea. Linn. is an important oil seed crop of Maharashtra. It is cultivated in both summer and kharif season in Solapur District. The bunchy variety SB-11 of groundnut was selected for aerobiological studies and the site selected for the study was a village Dadpur of Solapur District. For the occurrence of Pathogenic fungal spores and the incidence of disease, the aerial survey was carried out over the groundnut fields during summer seasons of the year 1991 and 1992 with the help of Volumetric Tilak Air Sampler. The meteorological data for the concerned period was obtained from Mahatma Phule Agriculture College, Solapur. The total number of biocomponents trapped by the sampler was 76 out of which 45 were pathogenic fungal spores. Other types included 31 types like saprophytic fungal spores, hyphal fragments, Pollen grains, nematode cysts and insect parts. In all, 45 different types of plant pathogenic fungal spores were trapped of which 3 types belonged to the class Phycomycetes, 11 to Ascomycetes, 2 to Basidiomycetes and 29 to Deuteromycetes. The spores like

Cladosporium. sp, Alternaria. sp, Helminthosporium.sp. Aspergillus. sp, Nigrospora. sp, Drechslera. sp, Geotrichum. sp, Fusarium. sp, Curvularia. sp, were the dominant types recorded from the air during two summer seasons. The epidemiological study was carried out during the period of investigation in which, the pathogenic spores like Alternaria. sp, Puccinia. sp, Cercospora. sp, Aspergillus. sp, Fusarium.sp, produced the diseases in crop.

**KEY WORDS:** Aerobiology, SB-11. Meteorology, Dadpur.

### **INTRODUCTION:**

Groundnut (*Arachis hypogaea*. L-family Fabaceae) is one of the important oil seed crops of Maharashtra and is grown in both summer and kharif seasons. The crop is generally infected by *Puccinia arachidis* speg. which is responsible for pod yield losses to the tune of 50% one or more itself or combined with other Pathogens. The fungal spores like *Cercospora personata* Ell and Ev, *Cercospora arachidicola* Hori, *Alternaria arachidis* Kulkarni, *Aspergillus niger Van* Tiegh, *Fusarium solani* (Mart) Sacc, (Maillaiah.1982) infected the crop and deceased the yield. All these spores are thought to be the only source of inoculums and development of the fungal diseases.

The out break of the diseases is mainly governed by the environmental factors. Any kind of aerobiological work was not carried out by anybody from this region. Hence it was felt necessary to carry out the present research work.

### **MATERIALS AND METHODS**

Aerobiological study was undertaken during the summer seasons of the year 1991-92 over the groundnut fields with the help of Tilak Air Sampler (Tilak.S.T.1988). The sampling site (Dadpur village) selected was west to and 14 kms away from Solapur city. Groundnut variety (early) SB-11(bunchy) was sown (40kg/acre) in the field during summer seasons of the year 1991 and 1992. Volumetric Tilak Air Sampler was installed on a table in the middle of groundnut field at the height 2.5 ft above the crop pointing the orifice towards the east. The fungal spores were trapped on the cello tape which was fixed on the rotating disc inside the Tilak air sampler. Air sampling was carried out about 123-125 days during each summer season. Sampling was started one week before the sowing and continued up to one week after harvesting the crop. The crop was harvested after about 110 days in each season. During the sampling period, the cello tape was removed once a week from the sampler and cut in to 8 equal parts, each part representing 24 hours spore trap. Each part was further divided in tom two equal parts i.e. one part representing day spora and other night spora. Each part was further divided into 12 equal divisions; each division representing one hours spore trap. During the period of investigation, the Permanent slides were prepared after every 8 days using glycerin jelly. Each slide was scanned for the pathogenic spores and other types.(saprophytic fungal spores,hyphal fragments,pollengrains, nematode cysts, insect parts) (Meshram. 1990, Sankaye, 1990).

Identification of fungal spores was done on the basis of spore morphology and reference slides prepared from the known fungi. (Subramanian, 1971; Barnett and Hunter; 1972, Ainsworth. et.al. 1973). The crop was also examined throughout the growing seasons at different growth stages for the disease incidence and associated pathogenic fungi. (Mukerji. et.al. 1986). Prior to the research work, meteorological data for the concerned period was obtained from Mahatma Phule Agriculture College, Ravivar peth, Solapur (Mulegaon farm).

Airspora concentration and percentage contribution of dominant pathogenic spores type, month wise and group wise seasonal spore concentration, epidemiological study of groundnut crop

was carried out during summer seasons. Prior to the epidemiological studies, Infectivity lindex (I.I.) was calculated (Sharma, P.D.2001) using the following formula. For+ the calculation of Infectivity Index (II), 200 diseased plant parts were taken.

II = <u>Sum of all disease ratings</u> X 100 Total number of ratings X Maximum disease grade

#### **RESULTS AND DISCUSSIONS**

During the summer seasons, 45 different types of pathogenic spores were encountered, along with other types. Some dominant spore types in both summer seasons were recorded as Aspergillus.sp, Puccinia.sp, Alternaria.sp, Cladosporium .sp,Cercospora.sp, Corynespora.sp, Curvularia.sp, Drechslera.sp, Fusarium.sp, Geotrichum.sp, Haplosporella.sp, Heliminthosporium.sp, Nigrospora.sp, Periconia.sp,. During the summer seasons, maximum spore concentration and percentage contribution was recorded of Cladosporium sp i.e 113582(S1) and 101276(S2)/m<sup>3</sup> of air and percentage contribution 31.43(S1) and 27.38(S2) followed by Alternaria sp as 34020(S1) and 33054(S2)/m<sup>3</sup> of air and percentage contribution 9.42(S1) and 8.94(S2), Helminthosporium sp as 26530(S1) and 31136(S2)/m<sup>3</sup> of air and percentage contribution 7.34(S1) and 8.42(S2), Aspergillus as 20502(S1) and 15806(S2)/m<sup>3</sup> of air and percentage contribution 5.69(S1) and 4.27(S2) and remaining followed by Nigrospora, Corynespora, Curvularia etc. (Table-I). The total spore concentration in Summer first(S1) was249388/m<sup>3</sup> of air and in second summer (S2)was246708/m3 of air. The total percentage contribution of pathogenic spores recorded in summer first (S1) was 69.04 where as it was 66.70 in summer second (S2). The mean percentage contribution of pathogenic spores in both the seasons was 67.87(Table-I) Cladosporium.sp, spores were dominant in both summer seasons (29.40%). The spore concentration of *Cladosporium.sp* was recorded as 113582 and 101276/m³ of the air. The mean percentage contribution of pathogenic airspora was recorded as 69.04% and 66.70% respectively during summer 1991 and 1992 season with the mean of both the season as 67.87% (Table-I) (Singh, et.al 1989). The spores like Rhizopus were not recorded during summer first where as the spores of; Pestalotia, Deightoniella, and Stigmina were not recorded from the air during summer 1992 season. (Bansal, et.al.1989) The low temp. high R.H. and scanty rains have positive effect on sporulation and discharge of spores. (Mishra, et.al 1971)

Table I

Total Airspora Concentration and Percentage Contribution of some dominant Pathogenic spore types during first summer (S1) and second summer (S2) season over the groundnut fields.

S1:27-01-1991 to 30-05-1991

S2:28-01-1992 to 30-05-1992

Spore types		entration of m <sup>3</sup> of air	Percentage of the total	Mean percentage	
	S1	S2	S1	S2	contribution
Aspergillus.sp	20502	15806	05.69	04.27	04.98
Puccinia.sp	3206	2632	00.89	00.71	00.80
Alternaria.sp	34020	33054	09.42	08.94	09.18
Corynespora.sp	7896	13118	02.18	03.55	02.86
Cercospora .sp	3206	3556	00.89	00.96	00.92
Cladosporium.sp	113582	101276	31.43	27.38	29.40
Drechslera.sp	6552	10066	01.82	02.72	02.27
Fusarium.sp	11606	7658	03.21	02.07	02.64
Helminthosporium.sp	26530	31136	07.34	08.42	07.88
Curvularia .sp	7378	13244	02.04	03.58	02.81
Nigrospora .sp	14910	15162	04.13	04.10	04.11
Total spores and % contribution	249388	246708	69.04	66.70	67.87

The spores of Deuteromycetes class were recorded maximum 80080/m³ of air in the month of February (1992) and79492/m³ of air March (1991) and minimum 14364/m³ of air in the month of January 1991 and18648/m³ of air in the month of January 1992. The spores of Asomycetes class were recorded maximum16660/m³ of air in the month of March (1992) and12572/m³ of air May (1991) and minimum 5278/m³ of air in the month of January 1992 and5656/m³ of air in the month of March1991. The spores of Basidiomycetes class were recorded maximum1862/m³ of air in the month of April (1992) and1316/m³ of air May (1991) and minimum 294/m³ of air in the month of January 1991 and196/m³ of air in the month of January1992. The spores of Phycomycetes class were recorded maximum1134/m³ of air in the month of February (1991) and700/m³ of air February (1992) and minimum 98/m³ of air in the month of January 1991 and210 /m³ of air in the month of March1992. The pathogenic spores were recorded maximum89950/m³ of air in the month of February (1992) and786058/m³ of air March (1991) and minimum 22708/m³ of air in the month of January 1991 and24416/m³ of air in the month of April1992. The spore group deuteromycetes was dominant in both the summer seasons followed by ascomycetes, other types, basidiomycetes and phycomycetes. (Table-II).

The other types were recorded maximum9632/m³ of air in the month of May1991 and11578/m³ of air May (1992) and minimum 2142/m³ of air in the month of January 1991 and2940/m3 of air in the month of January1992. The maximum spore types were recorded as 97678/m³ of air in the month of February 1992 and93968/m³ of air March 1991 and minimum 24850/m³ of air in the month of January 1991 and27356/m³ of air in the month of January1992. Maximum percentage contribution of pathogenic spores to the total air spora was recorded as recorded as 92. 08 % in the month of February 1992 and 91.58% in the month of March 1991 and minimum 88.38% in the month of April 1991 and86.63% in the month of May 1992. This spore concentration is mostly related to the earlier work. (Sahu, S.K.1988). The fungal spores are not only pathogenic but also occur as biopollutants in the atmosphere with the genera belonging to the spore group deuteromycetes as dominant. (Sinha Anupama et.al.1998) (Table-II).

#### Table- II

Month wise seasonal Pathogenic spore concentration in number/m<sup>3</sup> of air of each spore group and percentage contribution of spores in first (S1) and second (S2) summer season over the groundnut field.

S1:27-01-1991 to 30-05-1991

S2:28-01-1992 to 30-05-1992

Sr.	Spore group/total	January		February		March		April		May	
No.	spores/% contribution.	S1	S2	S1	S2	S1	S2	S1	S2	S1	S2
1	Phycomycetes	98	294	1134	700	532	210	196	350	224	238
2	Ascomycetes	7952	5278	9128	8554	5656	16660	11900	10388	12572	8946
3	Basidiomycetes	294	196	1008	616	378	868	784	1862	1316	1162
4	Deuteromycetes	14364	18648	52220	80080	79492	56406	58268	51968	66570	64708
5	Other types	2142	2940	8302	7728	7910	10892	9352	8610	9632	11578
6	Total pathogenic	22708	24416	63490	89950	86058	74144	71148	64568	80682	75054
	Spores										
7	Percentage contribution of pathogenic spores	91.38	89.25	88.44	92.08	91.58	87.19	88.38	88.23	89.33	86.63
8	Total	24850	27356	71792	97678	93968	85036	80500	73178	90314	86632

So far the distribution of fungal spores based on taxonomic spore group is concerned, deuteromycetes spore group was dominant with the maximum spore trap in the month of March 1991 (79492/m3of air) and February 1992 (80080/m³ of air), followed by ascomycetes as12572/m³ of air in May 1991 and16660/m³ of air in March 1992, basiodiomycetes as1316/m³ in May 1991 and1162/m³ of air in May1992, other types as 9632/m³ of air in May 1991 and 11578/m³ of air in May1992 respectively. The highest spore concentration during summer season was recorded in and March 1992 (10892/m³ of air) during the summer seasons the month of March 1991 (86054/m³ of air) and February 1992) (Table-II)

The epidemiological study of groundnut diseases was carried out during the summer seasons of the year 1991 and 1992. During the summer seasons (S1&S2) *Alternaria*.sp, spores were recorded in the last week of January, leaf blight disease appeared (I.I21.6%) after 30 (S1) and 31 (S2) days in the crop when the temp. was  $23.7^{\circ}$ C and  $21.0^{\circ}$ C and R.H. was 42.5% and 24%, wind velocity was 4.9 and 4.4 km/h respectively during summer first and second season when the crop was at vegetative growth stage.(Table-III)

Aspergillus.sp, spores were recorded in the last week of January (S1 & S2), pod rot disease appeared after 61 (S1) and 62(S2) days in the crop (I.I 27.5%) when the temp was  $29.3^{\circ}$ c and  $19.4^{\circ}$ c, R.H. was 56.5% and 35.5%, wind velocity was 5.5 & 6.9 km/h respectively during summer first and second season when the crop was at pod formation and maturation stage. .(Table-III)

Cercospora.sp, spores were trapped in the last week of February (S1) and first week of March (S2), after 34 (S1) and 47 (S2) days, the tikka disease appeared in the crop (I.I29.4%) when the temp. was  $30.0^{\circ}$ C and  $20.6^{\circ}$ C, R.H. was 33.5% and 34.5% and wind velocity. was 7.9 & 4.1km/h respectively during summer first and second season when the crop was at vegetative-flowering stage. .(Table-III)

Spores of *Fusarium*.sp, were recorded from the air in the last week of January (S1) and first week of February (S2), the wilt/root rot disease appeared in the crop (I.I.14.0%) after 20(S1) and 41(S2), days

when the temp. was 21.3°C and 17.6°C, R.H. was 45.0% and 39.5% wind velocity. was 4.6 & 4.6 km/h respectively during the summer first and second season when the crop was at seedling stage. .(Table-III) *Uredospores of Puccinia*.sp, were trapped from the air in the last week of January (S1 & S2), the rust disease appeared in the crop (I.130.50%) after 86(S1) and 81(S2) days when the temp. was 19.5°C and 20.9°C, R.H. was 44.0% and 27.0% wind velocity. was 3.4 & 4.0 km/h respectively during summer first and summer second season when the crop was at pod elongation and maturation stage. .(Table-III) The crop remained heavily infected by rust disease as it was also shown by earlier worker (*Subramanyam*.et.al 1983). The pathogenic spores like *Colletotrichum, Curvularia, Dipoldia, Myrothecium and Pestalotiopsis*.sp were recorded from the air but did not produce the diseases in this crop.. This work correlates the earlier work carried out for the other crop.(Bansal.et.al.1988) Epidemiological study indicated that the Alternaria arachiidis caused leaf blight, Aspergillus niger caused pot rot, *Cercospora arachidicola* caused tikka, Fusarium solani caused wilt and Puccinia arachidis caused rust disease in the groundnut crop in both summer seasons. The disease intensity was comparatively low as compared to the Infectivity Index of kharif crop. (Mali *et.al.*2011) This is because of the meteorological Parameters of summer seasons have somewhat adverse effects on the

Table-III

Epidemiology of groundnut crop during summer first (S1) and summer second (S2)

S1:27-01-1991 to 30-05-1991

S2:28-01-1992 to 30-05-1992

sporulation and spore discharge. So it can be better to grow the crop in summer season

Spore Types	Date of spore incidence		I		Infectivity Temperature Index ( °c)		Relative Humidity(RH) (%)		Wind velocity (Km/h)		Crop stage	
	S1	S2	S1	S2	(mean)	S1	S2	S1	S2	S1	S2	
Alternaria	27 Jan1991	29 Jan1992	25 Feb.1991	28 feb.1992	21.6%	23.7° C	21.0° C	42.5	24.0	4.9	4.4	Vegetative
Aspergillus	26 Jan1991	27 Jan1992	26 Mar 1991	28 Mar 1992	27.5%	29.3° C	19.4 <sup>0</sup> C	56.5	35.5	5.5	6.9	Pod formation and maturation
Cercospora	28 feb1991	07Mar 1992	03 Apr 1991	23 Apr 1992	29.4%	30.0° C	20.6° C	33.5	34.5	7.9	4.1	Vegetative and flowering
Fusarium	23 Jan1991	02 Feb 1992	12 Mar 1991	14 Apr 1992	14%	21.3° C	17.6° C	45.0	39.5	4.6	4.6	Seedling stage
Puccinia	26 Jan1991	28 Jan 1992	21 Apr 1991	16 Apr 1992	30.50	19.5° C	20.9° C	44.0	27.0	3.4	4.0	Pod elongation and maturation

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