STUDIES ON THE INFLUENCE OF ROOT SYSTEM OF ABELMOSCHUS ESCULENTUS AND SORGHUM VULGARE ON THE RHIZOSPHERE AND RHIZOPLANE MYCOFLORA OF CAJANUS CAJAN

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Influence of roots of three crops grown in earthen pots in sequence of *Abelmoschus* esculentus (L.) Moench. Sorghum vulgare Pers and Cajanus cajan (L) Millps. on rhizosphere effect was studied. The results indicated that the fungal population in the rhizosphere and non-rhizosphere soil varied with crop and age. The positive rhizosphere effect was found with *A. esculentus* and *C. cajan* while negative effect recorded in case of *S. vulgare*. A total number of 53 fungal species isolated from rhizosphere and non-rhizosphere soils during this study. However, 35, 40 and 36 fungi were reported from *A. esculentus*, *S. vulgare* and *C. Cajan*, respectively against 38 species when *C. cajan* grown alone (control). Hence it is suggested that the rhizosphere of *C. cajan* was influenced by other two crops.

Keywords : Crop sequence; Mycoflora; Rhizosphere; Non-rhizosphere; Rhizoplane.

### Introduction

Soil supports a very complex microbial community living in a state of dynamic equilibrium. Rhizosphere is a metabolically active region with conspicuous variation on its mycoflora depending upon the root exudates, genus, species, varieties, age and phase of growth of plants, soil conditions and environmental condition.

Age of the plant has an important bearing on rhizosphere population as it alters the soil mycoflora (Mishra and Kamal 1970; Deoray and Bhide, 1981; Dubey and Dwivedi 1988). Menon and Williams (1957) obtained differences in the mycoflora of soils after successive cropping of alfalfa, corn, oats or wheat. There is a little information regarding the different patterns of crop sequence (Prasad, 1968; Wajidkhan et al, 1973). Cultivation of crops favours the build up of some soil borne pathogens which are either pathoge. nic on specific crop or aggressive sccondary invaders. Therefore the deals with the paper present

rhizosphere ecology and rhizoplane mycoflora of *C. cajan* under the influence of root system of two crops.

## **Material and Methods**

Three crops viz., Abelmoschus esculentus (L) Moench., Sorghum vulgare Pers. and Cajanus cajan (L) Millsp. were selected for the present study. The healthy seeds of all the selected surface sterilized with crops were 0.1% mercuric chloride and washed with distilled water throughly. A. esculentus seeds were sown in earthen pots (10" X 12") as the first crop and each pot watered at regular intervals of three days. Care was taken to see at no time the water leaked out of the pots. Samples of the rhizosphere, non-rhizosphere and rhizoplane were collected from 10th. 20th, 30th, 40th and 50th day plants. After completing the collection of data on rhizosphere, non-rhizosphere and rhizoplane, the same soils (pots) were used for the second crop. S. vulgare. Similarly the third crop C. caian was raised in the same pots. In all cases collection of the data was similar as in the case of A esculentus.

Two media namely Czapek-Dox Agar (CZA) medium and Potato Dextrose Agar (PDA) medium were used. For rhizosphere studies, the soil adhering to the root system was removed using a sterile brush under asceptic conditions. To obtain soil suspension basically the method of

Timonin (1940) and Agnihothrudu (1953) were followed. Serial dilu tions were prepared by transferring one ml of the solution to the test tube containing 9 ml of sterile distilled water and obtained dilution up to 1:10,000 as this dilution was found satisfactory. Dilution plates were prepared by transferring 1 ml of soil suspension (1:10,000) asceptically into sterile petriplates containing 10 to 15 ml of CZA and PDA media and left for solidification. Later the plates were incubated at 25+2°C. A total of ten replicates were maintained for each sample (5 replicates for each medium). The plates thus prepared were observed uptil 7 days for mycoflora and the fungi appeared were isolated and maintained them for further study.

For the isolation of rhizoplane mycoflora "root washing" technique of Harley and Waid (1955) was adopted. After excess of water was removed from roots with sterile filter paper, they were then cut into small pieces (2mm) and each segment was placed in petriplates containing CZA and PDA media with streptopenicillin. Ten replicates with 5 segments per plate were maintained. Plates were incubated at  $25\pm2^{\circ}$ C and observed for mycoflora uptil 7 days.

# **Results and Discussion**

Quantitative analysis – The positive rhizosphere effect observed in A. esculentus gradually decreased from

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Table 1 : Influence of Root System of Three Crops on Rhizosphere and Nonrhizosphere Soil Mycoflora	

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	2 U			1.212	1.095	1.126	1.194	1.198	1.550	1.300	1.290	1.110	1.110	1.236	0.942	0.990	0.815	0.868	1.120	1.019	1.024	1.215	1.134	
Cajanus cajan	Number of fungi in nonrhizosphere soil	(.c.n.o 6/= 01)		13 20	13.60	6.30	14.90	15.60	8.00	12.20	13 10	16.10	14.00	23.30	39.80	25.40	22.80	19.00	11.00	30.50	28.30	9.50	24.60	
Crop sequence : Abelmoschus esculentus, Sorghum vulgare and Cajanus cajan	Number of fungi in rhizo- sphere soil (10 <sup>-4</sup> /g 0.D.S.)			16.00	14.90	7.10	17.80	18.70	12.40	15.90	16.90	18.00	15.55	28 80	37.50	25.20	18.60	16.50	13.00	31.10	29.00	11.55	27.90	0.D.S : Oven dried soil
Crop sequence : Abelmoschus	Age of the Crop plant (in	days)	Crimin orign (Control)	Cajanus cujun (connor)	20	30	00	50	11-1-1-00-00-000 10	Abelmoschus 20	escurentus == 30	40	50		Sorgnum vuigure		40	50		Cajanus cujun 20	30	40	20	0.D.S ; Ove

J. Phytol. Res. 4 (1)

10th to 15th-day. More fungi species were recorded in the rhizosphere than the non-rhizosphere up to the 40th day (Table 1).

S. vulgare grown in the same pots used for A. esculentus recorded a negative rhizosphere effect except on 10th-day. The maximum and minimum fungal populations were recorded on 20th and 50th-day of both rhizosphere and non-rhizosphere soils, respectively and as regard to number of species, the same was recorded on 20th and 40th-day of rhizosphere and 10thand 20th-day of non-rhizosphere soils, respectively (Table 1) C. cajan, grown in the same set of pots showed a positive rhizosphere effect and it increased gradually from the 20th-to 40th-day which later decreased (Table 1).

Control C. cajan rhizosphere also showed a positive rhizosphere effect at all stages of plant growth. The maximum rhizosphere effect was observed on 10th day which later decreased on 20th-day. However the rhizosphere effect gradually increased with the increase of the plant growth from 20th to 50th-day. But C. cajan grown in this sequence of crops expressed maximum and minimum rhizosphere effect on 10th-day and 40th-day, respectively. As regards to the number of species the maximum was recorded on the 10th. day in both rhizosphere and non-

rhizosphere soils of control C. cajan where as minimum number of species was observed on the 50th-day rhizosphere and 20th and 50th-day of nonrhizosphere soils (Table 1). Both rhizosphere and non-rhizosphere soils recorded maximum number of fungal population on 20th-day where as minimum was recorded on the 40th day of C. cajan in the sequence. However when C. cajan grown alone (control) recorded maximum fungal population on 50th day and minimum fungal population on 30th day rhizosphere and nonrhizosphere soils.

Qualitative analysis-A total number of 53 fungi were isolated from the rhizosphere and nonrhizosphere soils of 3 crops grown in a sequence. But 35, 40 and 36 fungal species were observed from A. esculentus, S. vuland C. cajan, respectively gare (Table 2). It is interesting to note that Actinomucor elegans, Bipolaris tetramera, Drechslera halodes, Oedocephalum sp. and Penicillium expansum which were isolated from A. esculentus were absent in both soils of S. vulgare. However A. mellus, Cylindrocarpon sp., Rhizopus arrhizus and Torula herbarum were exclusively present in both soils of S. vulgare, and not reported with A. esculentus (Table 2).

C. cajan grown as a third crop recorded the presence of 16 fungal species from the rhizosphere soils whereas 6 fungi were recorded from the nonrhizosphere soils. Fungi such

as A. restrictus, Helminthosporium sp., Mortierella sp., Rhizopus stolonifer, Sporothrix sp. and Trichothecium roseum were restricted to the rhizosphere soils of C. cajan and not found with A. esculentus and S. vulgare whereas Phoma sp., was isolated from the nonrhizosphere soils only. It is interesting to not that Absidia gluaca, Actinomucor elegans, Alternaria sp., Aspergillus candidus, A. chevalieri, A. oryzae, A. phoenicis, A. sclerotiorum, A. terreus, Bipolaris tetramera, Curvularia lunata, Mucorales member, Oedocephalum sp., Paecilomyces sp. and Torula herbarum present in the two crops studied were found absent in both soils of C. cajan. (Table 2).

In all 38 fungi were recorded in control C. cajan, of which Absidia gluaca Acremonium sp., Aspergillus unguis, Bipolaris specifera, Chaetomium sp., Fusarium poae, Penicillium nigricans, P. rubrum, Rhizopus oryzae, Torula sp., Bipolaris tetramera, Cunninghamella echinulata were recorded exclusively and not reported with the crop grown in the sequence (Table 2).

Rhizoplane studies— The rhizoplane fungi comprised of 34, 23 and 29 fungal forms in case of *A. esculentus*, *S. vulgare* and *C. cajan*, respectively. A total of 44 fungal species were recorded from the rhizoplane of 3 crops studied in a sequence. In this

crop sequence A. esculentus rhizoplane represented by Absidia gluaca, Aspergillus alutaceus, A. chevalieri, A. japonicus, A. oryzae, Cylindrocarpon sp., Mucor sp., Phoma sp., and Rhizopus nigricans and were not recorded from the rhizoplane of S. vulgare and C. cajan. The fungal species present in the rhizoplane of A. esculentus and S. vulgare but absent in C. cajan were Acrophialophora SD., brown sterile mycelium. Drechslera halodes, Fusarium oxysporum and Penicillium citrinum. A. flavipes, Curvularia lunata, Verticillum sp. and white sterile mycelium present in the rhizoplane of A. esculentus and C. cajan were absent with S. vulgare rhizoplane. The rhizoplane fungi of control C. cajan differed by having Aspergillus alutaceus, A. phoenicis, A. ustus, Bipolaris tetramera, Chrysosporium sp. Cylindrocarpon sp. Fusarium oxysporum, F. semitectium, Oedocephalum sp., muscorales member, Paecilomyces sp, Phoma sp., Penicillium citrinum, P. digitatum and Rhizopus nigricans. This observation indicated that those species are specific to C. cajan but not occurred on account of the influence of other crops grown earlier. Hence the root system of each crop might have its specificity in controlling the fungi. This may be due to differences in the chemical nature of root surface that influence the occurrence of fungal flora and hence the variation (Table 3)

However this needs further study in this direction.

Tiwari and Mehrotra (1968) and Daval and Srivastava (1973) reported the abundance of Aspergilli and Penicillia in the rhizosphere of various crops. Later Dubey and Dwivedi (1988) also reported that Aspergilli were dominant followed by Penicillia and Fusaria in the non-rhizosphere and rhizosphere regions of Soybean. In the present study species of Mucor, Rhizopus, Syncephalastrum, Alternaria, Aspergillus and Fusarium were found to be the most common fungi in rhizosphere and nonrhizosphere soils of 3 crops studied. However the dominant fungi in both rhizosphere and nonrhizosphere soils of 3 crops include Aspergillus, Penicillium and Fusarium. Of these Aspergillius was recorded in greater abandance followed by Fusarium. Similar observations were made by Patil and Thite (1987) who isolated Aspergillus, Penicillium, Fusarium, and Trichoderma from rhizosphere and nonrhizosphere soils of S. vulgare. In the present study increase in the rhizosphere mycoflora with the age of the plant was observed. It seems that the increase was due to various factors like increased exudation. decomposition of morbund root hairs, epidermal cells and cortex accumulation of cell materials as observed by earlier workers (Rovira, 1965 a, b Tiwari and Mehrotra, 1968) In the present investigation roots in early stages of plant growth were chiefly colonized by mucorales, Hyphomycetes and sterile mycelia. Ascomycetes started colonizing the roots only during the flowering of plant. This pattern of colonization of different fungi on the plant roots in succession corroborate the findings of previous workers (Parkinson and Clark, 1961; Dickinson and Pugh, 1965).

Srivastava and Daval (1982) found that A. fumigatus in rhizosphere through out from seedling to senescence of the plant though certain forms appear for short duration. In the present study it is clear that different stages of plant growth had marked influence on the fungi in the rhizosphere. It is of interest to note that in the present study there was a significant difference in the number of different fungi on rhizoplane at different stages of plant growth. Some fungi established themselves on the root surface where as others were confined to rhizosphere only. At the same some fungi were resticted to nonrhizosphere soil and were not isolated either from rhizosphere or rhizoplane. They include Aspergillus japonicus, A. parasiticus and Phoma sp. This behaviour may be attributed tentatively to the degree of sensitivity of fungi to their response to the amount of exudates in the soil from the sites of exudation along with roots as suggested by Dix (1967). Earlier workers have

Aspergillus alutaceus Actinomucor elegans Absidia gluaca Name of the fungi Table 2: Influence of Abeimoschus esculentus and Sorghum vulgare on the Rhizosphere Mycoflora of Cajanus cajan Alternaria sp. Acremonium sp. A. chevalieri A. carneus A. candidus niger nidulans oryzae erythrocephalus japonicus parasiticus melleus fumigatus flavus Havipes -D Co Z N 34 Ae D Z 10th day Sv C D Z 6 Z R 7 S 1 8 2 -J Co 20th day N Z ω 7 Ae 4 T R G Z Sv J 0 7 D Z Cc 00 R Co -D Z N ω Z 30th day D Ae 1 4 T G Z D Sv 6 1 1 Z D Cc 8 D -Z D Co N D ω 7 Z Ae 40th day 4 D S Z R Sv 6 D 1 z D Cc J 00 Z -D Co 1 N D ω Z Ae D 50th d 4 D

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C. lunata Curvularia geniculata Cunninghamella echinulata Chaetomium sp Brown sterilemycelium B. specifera **Bipolaris** tetramera A. versicolor A. sclerotiorum A. phoenicis 4. restrictus ustus terreus

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Table 3: Influence of Abelmoschus esculentus and Sorghum vulgare on the Rhizosplane Mycoflora of Cajanus cajan	esculer	itus e	s pui	orghi	a with	ulga	re on	the	Rhiz	ospla	ne N	lycofl	ora o	f Caj	anus c	cajan			
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shown that the composition of rhizosphere microflora differs both quantitatively and qualitatively from that in soil beyond the influence of the root (Krisilinkov 1963; Katznelson, 1965). Similar variations were recorded in the present study. Rangarao and Mukerji (1971) who studied four cultivars of wheat rhizosphere and rhizoplane observed that the peak population levels were attained at different stages of plant growth and rhizosphere and rhizoplane populations differed appreciably in four cultivars of wheat. Earnoticed workers restricted lier occurrence of rhizoplane fungal flora in groundnuts (Gangawane, 1972) Corchorus capsularis and Hibiscus 1978), gram cannabinus (Mishra, (Antique et al., 1982). They also stated that there was significant difference in the number of different fungi on the rhizoplane at different ages of plant growth and observations made in this study are in confirmity with the earlier workers. In the present investigation the three sequence also crops grown in a differed both qualitatively and quantitatively.

#### Acknowledgements

The autors gratefully acknowledge the financial assistance provided by UGC, New Delhi for carrying out this work.

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