



RESEARCH ARTICLE

Mycopathological Studies on *Vigna radiata* (L.) Wilczek (Green gram) from Patur, Dist. Akola (MS), India.

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ABSTRACT

Study of fungi infection from infected green gram plant was carried out in present identify. Various fungal pathogens were identified from green gram plants with respect to different localities and varieties at field condition. Selected samples were collected from regions of studied area. Total ten and eleven fungi were identified from two variety of green gram AKM-9911 and AKM-9904 respectively. Green gram (*Vigna radiata* (L.) Wilczek) is one of the most widely used pulse crop of India. It is widely cultivated in Maharashtra, Uttar Pradesh, Gujarat, Tamil Nadu, Andhra Pradesh and Bihar. It is cultivated in many tropical and subtropical regions of the world including India and was also cultivated in ancient Indian.

Keywords: Mycopathological Studies, *Vigna radiata* (L.) Wilczek, Green gram, AKM-9911 and AKM-9904, Patur

INTRODUCTION

India is an agricultural country where economy is based on agriculture. Till recent past, before independence our agriculture depended on rains. The farmers used to grow their own material for cropping but after the technological evolution since the commence of 21th century the scenario has changed as many hybrid and genetically engineered varieties of agricultural crop plants are commercialized and that are also used to grow by farmers. However, it has been observed that these varieties also showed susceptibility to variety of pathogens resulting into lesser yield. The amount of disease is generally referred to as disease intensity (Teng, 1985). On the other hand disease severity is determined by a function of degree of affection, colonization and damage of host tissue. Measurement of severity based on lesion number or lesion area may be less related to yield, but more to disease progress. Measurement of both actual and visible disease (Rouse, 1988) in terms of percent tissue affected and the green leaf area duration respectively, gives more precise conclusion on disease and yield loss in the presence of a pathogen. Fungi causes different diseases to crop plants. viz, Leaf spots, rust, smut, powdery mildew, downy mildew and wilt diseases are the common examples of fungal diseases. It has been observed that there are about 6000 rust fungi that attack wide range of seed plants and caused destructive diseases. Some common examples of fungal diseases are leaf spots of Mung bean.

Green gram botanically named as *Vigna radiata* (L.) Wilczek is one of the major pulse crop of India and cheapest source of plant protein. It is a short duration legume crop and cultivated worldwide for its dry seeds. Annual Mung bean production worldwide is around 2.5 to 3.0 million tones, harvested from about 5.0 million ha (Poehlman, 1991). India contributes about 45 % of the production. It is widely cultivated in Maharashtra, Uttar Pradesh, Gujrat, Tamil Nadu, Andhra Pradesh and Bihar. The dried seeds of green gram contain 23.6% protein, 58.2% carbohydrates, 1.2% fats, 3.3% % fibers and 4% minerals.

Green gram is one of the most important pulse crop. It is grown in almost all parts of the country and belongs to family Fabaceae. Mung bean is an excellent source of high quality protein. It is consumed in different ways as dal, halva, snack and so many other preparations. Ascorbic acid (Vitamin-C) is synthesized in sprouted seeds of mung bean. The leguminous crops have the capacity to fix-atmospheric nitrogen through symbiotic nitrogen fixation. It is also used as green manure crop. It is grown in summer and kharif season in northern India and in southern India (Yadava, 1992). The major fungal diseases which infect the Mung bean are Root rot - *Macrophomina phaseolina* (Tassi.) Goid.,

Web blight, *Rhizoctonia solani* Khun. - *Thanatephorus cucumeris*, Powdery mildew *Erysiphe polygoni* DC., *Cercospora* Leaf spot- *Cercospora canescens* and Anthracnose- *Colletotrichum dematium* and *C. lindemuthianum* (Grewal, 1988). Moreover, seed infection of *Rhizoctonia bataticola* on *M. phaseolina* ranges from 2.2-15.7% which causes 10.8% in grain yield and 12.3% in protein content of seed in Mung bean showed that, due to changing climatic conditions, the pathogen attack on the Mung bean also differs (Kaushik *et al*, 1998). Mung bean is more susceptible to *Cercospora* leaf spot disease in humid conditions and the percentage infection may reach up to 60% in locally cultivated varieties leading to nearly 50-60% yield loss. Chilkuri *et al*, (2012) studied the seed borne fungi in Green gram and subsequently determined their effect on seed germination.

The average yield of Green gram crop is very low mainly due to low inherent yield potential and susceptibility of the crop to disease (Thakur *et al*, 1977). Leaf spot disease caused by *Cercospora canescens* is a serious disease in the Green gram growing areas during the season (Bashir, 1985). *Cercospora* leaf spot is one of the important disease that causes serious losses to Green gram crop and 23% losses in yield have been reported (Quebral and Cagampang, 1970). Maximum loss of 61% was observed in case of grain yield (Iqbal *et al*, 1995). Several workers had reported the effective control of the disease with the application of fungicides (Singh and Naik, 1977; Singh and Singh, 1978).

MATERIAL AND METHODS

a) Isolation of fungi from infected plant parts of green gram

Infected plant parts of two Green gram varieties viz. AKM-9911 and AKM-9904 were collected from the region of Patur Taluka, Dist. Akola. The infected plant materials were collected from the agricultural field. Small pieces measuring 2 mm², each of infected tissue were cut off from infected green gram plants with the help of sterile sharp knife. Pieces of diseased fruit were washed with tap water and surface sterilized with 1% Sodium hypochloride solution for 2 min and washed twice with sterilized distilled water and then dried using sterile filter paper. The infected plant parts were separately transferred to sterilized petri-dishes containing Potato dextrose agar (PDA) medium and incubated at 25°C for 10 days. Petri-dishes were observed daily and colonies of fungi were chosen. The isolated fungi were purified using single spore technique and then kept in a refrigerator on PDA medium (Gams *et al*, 1998). Pure colonies of fungal isolates were identified according to Ellis (1971). Symptoms were confirmed by Koch's postulates.

b) Study of symptoms of fungal diseases on green gram

Study of Symptoms were carried out on the different varieties of Green gram ie. AKM-9911 and AKM-9902

c) Study of Pathogenicity

Healthy plant part from each sample were surface sterilized with 90% ethanol and incision were made on them using a sterile 4 mm cork borer, similar sterile cork borer was used to cut pellets of agar containing the cultures of fungal mycelia of the isolates. These fungi were then inoculated into the hole created on the healthy plant parts of green gram respectively in a laminar air flow chamber. The controls with incision but not inoculated by fungal pathogens were established. The inoculated seed and the controls were placed in a clean polythene bag each moistened with wet balls of absorbent cotton wool to create a humid environment and incubated at 28°C for 5 days. After 72 hours, the inoculated infected plant parts were observed for symptom development. The causal agents were re-isolated from the infected fruits and compared with the original isolates (Akinmusire, 2011).

d) Method of Petri plate exposure

The Petri dishes (9cm- diameters) containing 20ml selected Potato Dextrose Agar (PDA) medium were exposed at 1 meter height from ground level in every cases for 15 min. over green gram field every week. The exposed Petri plates were brought to the laboratory and incubated in an inverted position at 28 °C ±1 °C for 7 to 8 days. The colonies developed were examined regularly, counted and identified. The fungal colonies were enumerated after their growth on the petri plates.

e) Identification of Fungus

Identification of fungal colonies were made by visual and microscopic examinations. Identification up to generic level was done with the help of standard mycological books and manuals (Ellis, 1971; Gams et al 1998). Details regarding the qualitative nature of the mycoflora, their incidence, abundance and percentage contribution were recorded. Identified fungal species photographed and presented in Plate I.

OBSERVATIONS AND RESULTS

In order to study the fungi associated with the green gram, the samples of infected green gram parts were collected from different agricultural field of Patur Taluka of Akola district. These infected samples were brought to the laboratory. Infected tissues of the plant were cut and

sterilized with 0.1% Mercuric chloride, washed thrice with sterile distilled water and placed on PDA medium. After incubation period of seven days grown fungi were isolated and identified and results are noted in Table No.1

Table 1: Isolation of fungi from different varieties of Green gram

Fungi	Green gram varieties	
	AKM-9911	AKM-9904
<i>Alternaria solani</i> (Ell. and Mart.) J. & G.	-	+
<i>Aspergillus flavus</i> Link.	+	+
<i>Aspergillus niger</i> Tiegh.	+	+
<i>Cercospora canescens</i> Ellis & G. Martin	+	+
<i>Chaetomium globosum</i> Kunze ex Fr.	-	+
<i>Cladosporium</i> sp.	+	+
<i>Colletotrichum lindemuthianum</i> Scrib.	-	+
<i>Curvularia lunata</i> (Wakker) Boedjin	+	+
<i>Erysiphe polygoni</i> DC.	+	+
<i>Fusarium oxysporum</i> f. sp.	+	+
<i>Fusarium solani</i> (Mart.) Sacc.	-	+
<i>Helminthosporium</i> sp.	+	-
<i>Phytophthora nicotianae</i> . Breda de Haan.	+	+
<i>Rhizopus stolonifer</i> (Ehrenb.) Vuill.	+	-

+ = Presence of fungus, - = Absence of fungus

The variety AKM-9911 gave fungi viz, *Rhizopus stolonifer*, *Erysiphe polygoni*, *Aspergillus niger*, *Aspergillus flavus*, *Cercospora canescens*, *Cladosporium* sp., *Curvularia lunata*, *Fusarium solani*, *Helminthosporium* sp., *Phytophthora parasitica*. The AKM- 9904 variety of green gram shown fungi such as *Chaetomium* sp., *Erysiphe polygoni*, *Alternaria solani*, *Aspergillus niger*, *Cercospora canescens*, *Cladosporium* sp., *Colletotrichum lindemuthianum*, *Curvularia lunata*, *Fusarium solani*, *Fusarium oxysporum*, *Phytophthora parasitica*. (Table No.1)The diseases caused by fungal pathogens were found on both varieties of green gram.

DISCUSSION

The present report suggests variations in the composition of mycoflora in different varieties of green gram reflect the variable degree of resistance and susceptibility for the establishment of particular group of fungi in the varieties yielding maximum number of fungal species. On the

contrary, the varieties with poor incidence of fungi represent their resistant capacity.

The presence of *Fusarium* in air over test field might be due to the prevalence leaf spot and top Necrosis disease in the field of green gram caused by *Fusarium* sp. Similar observations were reported by (Sreeramalu and Ramlingam, 1966; Mallaiiah and Rao, 1980 and Singh, 1992). The prevalence of *Colletotrichum* in the infected parts of the crops in the present study might be due to the infection to the crops by *Colletotrichum lindemuthianum*. It caused Anthracnose disease to Green gram crops. Hartung et al., (1981) and Singh (1992) reported these spores from the air from the abroad and India respectively. It is clear from results that *Curvularia lunata* and *Phytophthora parasitica* filtrate showed minimum green gram seed germination. Similarly *Curvularia lunata* and *Phytophthora parasitica* drastically hampered the Green gram seed germination at 100% concentration. These results are agreed with results given by Haikal (2008). The presence of *Cercospora* in the present study might be due to the infection of green gram crop by *Cercospora canescens* caused the *Cercospora* leaf spot. *Cercospora* leaf spot is a devastating disease that causes qualitative and quantitative losses to the crop (Sivaprakasam, 1983). Several scientists have screened the number of fungicides against the fungal growth. Efficacy of Daconil has been reported against *Cercospora* leaf spot on Mung bean (Iqbal et al., 1990) and mung bean anthracnose (Bashir et al., 1985).

CONCLUSION

It can be concluded that the farmers should be scientifically acquainted and trained about disease cycle, Identification of symptoms of disease at an early stage, Field sanitary practices, Crop rotation systems, methods of intercropping, proper use of water to a specific interval etc. Are the aspects whose knowledge would be useful to farmers to avoid problems of yield loss of agricultural crops.

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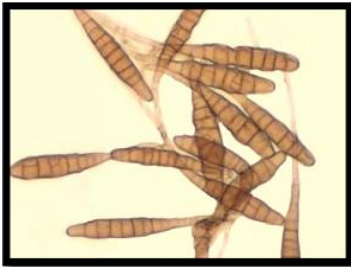
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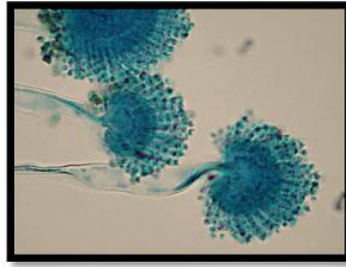
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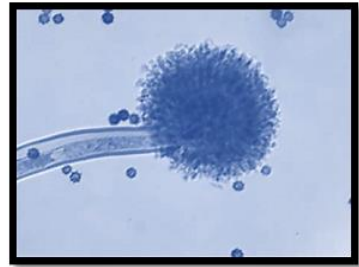
PLATE I



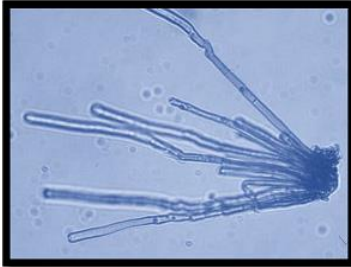
Alternaria solani (Ell. and Mart.) J. & G.



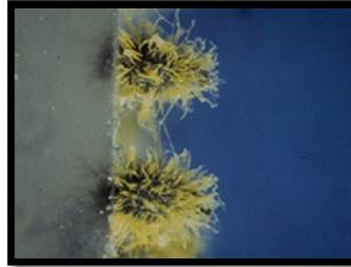
Aspergillus flavus Link



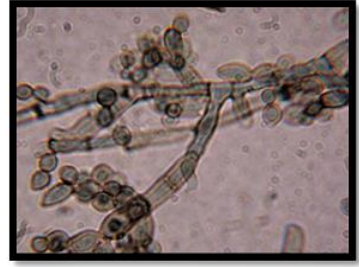
Aspergillus niger Tiegh



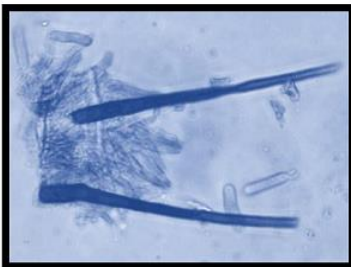
Cercospora canescens Ellis & G. Martin



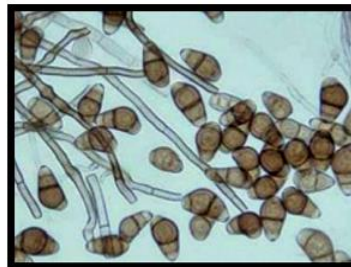
Chaetomium globosum Kunze ex Fr.



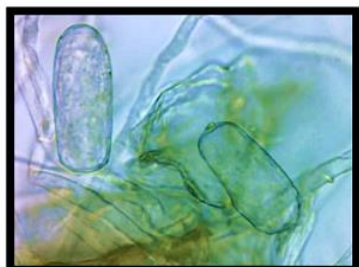
Cladosporium sp



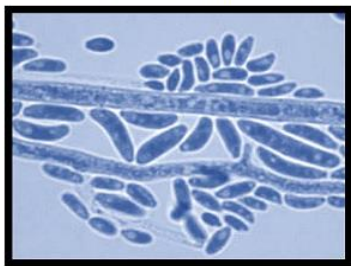
Colletotrichum lindemuthianum Scrib.



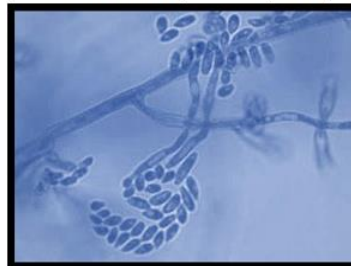
Curvularia lunata (Wakker) Boedjin



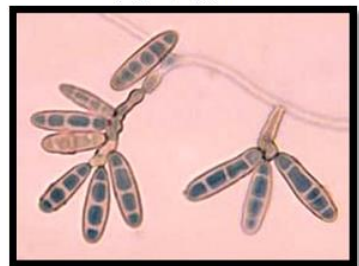
Erysiphe polygoni DC.



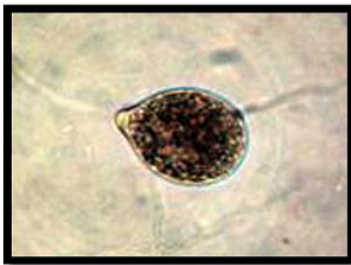
Fusarium oxysporum f.sp.



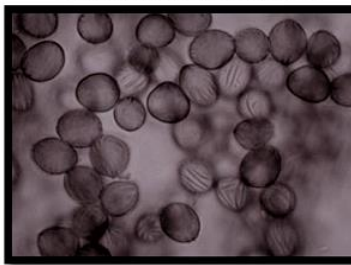
Fusarium solani (Mart.) Sacc.



Helminthosporium sp.



Phytophthora nicotianae. Breda de Haan.



Rhizopus stolonifer (Ehrenb.) Vuill.



Infected plants of Green gram at field