

Studies on Biological Control of Fungi

Tugaonkar S.G.

Department of Botany, Indira Gandhi Senior College, CIDCO, Nanded (M. S.)

Abstract— The poor incidence of *Aspergillus rubrer* was found in different concentrations of seed extracts of *Datura stramonium* and *Ipomea carnea*. Minimum incidence was found in 10 and 25 percent extracts of *Argemone mexicana* and *Cassia tora*, In general 50 percent seed extracts of all the tested medicinal plants were beneficial to control incidence of fungi. The maximum incidence of *Alternaria alternata* *Aspergillus flavus*, *A. niger*, *A. ustus*, *Curvularia lunata*, *Fusarium roseum* and *F. moniliforme* was observed on Soybean in presence of all the test percent extracts of medicinal plants.

Keywords— Sunflower, Soybean, Fungi, Medicinal plants

I. INTRODUCTION

Seeds are the main sources for contamination of oil seeds in field. Handling and packaging causes additional contamination during storage. The seeds are collected by simple methods and are commonly exposed to many contaminants before being dry enough and at storage in low temperature as per ([1],[2],[3]) Numerous studies revealed a wide spectrum prospects of using extracts of plants for biological control of pathogenic fungi ([4],[5],[6]). As a result, the management of fungi was carried out by using different seed extracts of medicinal plants.

II. MATERIALS AND METHODS

A) Collection of Seed Samples

The collection of Seed samples was as per method described by [7]. The seed samples were collected from fields, store houses, market places and seed companies and composite sample was prepared by mixing the individual samples together. The samples preserved in cloth bags in laboratory at room temperature. The seeds of *Annona squamosa*, *Argemone mexicana*, *Cassia tora*, *Datura stramonium* and *Ipomoea carnea*, were collected from study region. The seeds were shade dried at room temperature and were ground into fine powder and stored in airtight containers at room temperature till extraction.

B) Preparation of Acetone Extract

The method for acetone extraction was adopted as per [8] with slight modification in the preparation of concentrations. The powder of seeds extracted with organic solvent acetone, based on order of polarity using soxhlet apparatus. The extracts obtained were subsequently concentrated under reduced pressure to get their corresponding residues. Seeds of test spices were soaked in 10, 25 and 50 percent concentrations of 50 ml. acetone seed extracts in 100ml Erlenmeyer flask covered with parafin. The flasks were kept overnight at 25 to 30°C in darkness as per [9]

Seeds were tested [10] with slight modifications. The seeds soaked in plant extracts were taken and ten seeds plated in each petriplates on three layers of moistened blotter papers. The seeds of the same

samples soaked in distilled water served as control. Seeds were incubated at 25 to 30°C for ten days. After ten days, seeds were examined under a stereomicroscope for growth of fungi and the data was recorded.

C) Identification of Seed Borne Fungi.

The fungi occurring on each and every seed on the plates were identified preliminary on the basis of sporulation characters like sexual or asexual spores with the help of stereoscopic binocular microscope. The identification and further confirmation of seed-borne fungi was made by preparing slides of the fungal growth and observing them under compound microscope. Pure cultures of these fungi were prepared and maintained on glucose nitrate agar (GNA) slants. The identification was made with the help of manuals, as per ([11], [12])

III. EXPERIMENTAL RESULTS

As per the results of table number 1, maximum appearance of *Alternaria alternata*, *Aspergillus flavus*, *A. glaucus*, *A. niger* and *Cladosporium cladosporidies* was noticed in 10 and 25 percent testd seed extracts. The *Aspergillus rubrer* did not appear in the extracts of *Datura stramonium*, *Ipomea carnea* and *Annona squamosa*. Least incidence of fungi was found in 10 and 25 percent extracts of *Argemone mexicana* and *Cassia tora*. The seed extracts of the tested medicinal plants were more beneficial to control their appearance on spices. *Botrytis cineria*, *Curvularia tetramera*, *Fusarium oxysporum*, *Fusarium roseum*, were inhibited in all the concentrations of test extracts of medicinal plants.

The poor incidence of *Aspergillus rubrer* was found in different concentrations of seed extracts of *Datura stramonium* and *Ipomea carnea*. Minimum incidence was found in 10 and 25 percent extracts of *Argemone mexicana* and *Cassia tora*, whereas 50 percent seed extracts was effective in their incidence in case of *Aspergillus rubrer*. Different concentrations of seed extracts of *Annona squamosa* proved very much effective to control fungal incidence. The *Helminthosporium tetramera* did not respond to test concentrations of seed extracts of *Argemone mexicana* and *Cassia tora* except *Datura stramonium*, *Ipomea carnea* and *Annona squamosa*. In general 50 percent seed extracts of all the tested medicinal plants were beneficial to control incidence of fungi.

As per the results of table 2, least number of fungi were observed with their dominance on Soybean as compared to Sunflower. Seven fungi appeared on Soybean in presence of different seed extracts of medicinal plants. The maximum incidence of *Alternaria alternata* *Aspergillus flavus*, *A. niger*, *A. ustus*, *Curvularia lunata*, *Fusarium roseum* and *F. moniliforme* was observed on Soybean in presence of all the test percent extracts of medicinal plants. In general 50 percent seed extracts were highly beneficial to control the appearance of fungi.

Fungi	<i>Annona squamosa</i>			<i>Argemone mexicana</i>			<i>Cassia tora</i>			<i>Datura stramonium</i>			<i>Ipomoea carnea</i>			
	% of seed extract															
	C	10	25	50	10	25	50	10	25	50	10	25	50	10	25	50
	% incidence															
<i>Alternaria alternata</i>	15	20	10	10	15	10	10	15	10	10	20	15	05	15	10	05
<i>Aspergillus flavus</i>	25	20	15	--	20	10	10	15	15	15	25	20	10	15	15	10
<i>Aspergillus glaucus</i>	35	25	20	10	30	25	10	25	15	10	25	15	10	10	05	05
<i>Aspergillus niger</i>	20	--	--	--	15	05	--	15	10	10	20	20	05	10	--	--
<i>Aspergillus rubrer</i>	10	--	--	--	10	05	--	15	10	--	--	--	--	05	--	--
<i>Cladosporium cladosporidies</i>	25	25	15	05	20	10	05	30	20	50	20	10	--	10	--	--
<i>Curvularia tetramera</i>	10	--	--	--	10	--	--	05	--	--	--	--	--	--	--	--
<i>Fusarium oxysporium</i>	15	05	--	--	10	--	--	10	--	--	--	--	--	05	--	--
<i>Fusarium roseum</i>	10	--	--	--	10	--	--	10	--	--	--	--	--	--	--	--
<i>Helminthosporium tetramera</i>	15	10	10	05	15	10	--	05	--	--	15	10	10	10	15	05

C= Controle, -- = No incidence

Table 1. Effect of seed extracts of medicinal plants on mycoflora of Sunflower (Acetone)

Fungi	<i>Annona squamosa</i>			<i>Argemone mexicana</i>			<i>Cassia tora</i>			<i>Datura stramonium</i>			<i>Ipomoea carnea</i>			
	% seed extract															
	C	10	25	50	10	25	50	10	25	50	10	25	50	10	25	50
	% incidence															
<i>Alternaria alternata</i>	10	10	10	15	20	05	05	10	05	05	25	20	15	15	10	05
<i>Aspergillus flavus</i>	15	15	10	--	20	15	05	20	10	05	20	15	10	15	10	10
<i>Aspergillus niger</i>	20	20	20	10	15	10	05	20	15	10	30	30	25	25	10	10
<i>Aspergillus ustus</i>	15	15	10	05	20	10	05	20	20	15	20	20	15	20	20	10
<i>Curvularia lunata</i>	10	10	10	05	15	10	10	20	20	10	20	10	10	15	15	05
<i>Fusarium moniliforme</i>	20	20	15	05	20	15	10	20	15	10	20	10	10	20	05	--
<i>Fusarium roseum</i>	20	20	05	10	20	15	05	20	15	10	10	20	20	05	05	--

C= Controle, -- = No incidence

Table 2. Effect of seed extracts of medicinal plants on mycoflora of Soybean (Acetone)

REFERENCES

- [1] W. Kneifel and , E. Berger. "Microbial criteria and random samples of spices and herbs retailed on the Austrian market". *J. Food Prot.* vol. **57**: p.893-901,1994
- [2] S. Durakovic, J. Galic and P. Pajvoric, "Tokricni i kancerogenic metabolite gljiva 4 namirnicama i krimivima", *Hrana i Ishrama*.vol. **2**: pp.71-100. 1989
- [3] G. R Dimic, D.Suncica, T. Kocic, N. T.Alcksandra, L. V. Biserka, and M. S. Zdravko, "Mycopopulation of spices". *Acta Periodica Technologica*,.vol. 39 pp.1- 9 2008.
- [4] F. G..Braga, L. M. Maria, R. L. Bouzada, , M. O. Matos., F. O. Moreira, , E. Scio, and E. S. Coimbra, "Antileishmanial and antifungal activity of plants used in traditional medicine in Brazil". *J. Ethnopharmacol.* Vol.111 pp.396-402. 2007.
- [5] B. M. Waghmare., S.R. Shinde, V.T. Gorgile, R.S. Bajgire, and G.T. Sumanth, "Studies on novel bioproperties of botanicals against growth and sporulation on different species of *Fusarium*" National Conference, modern trends in plant sciences. Organised by Dept. of Botany Dr. Babasaheb Ambedkar Marathwada University, Aurangabad (M.S.), Abs. pp. 39. 2007
- [6] S. R. Shinde, "Studies on ethnobotanical and bioefficacy of some medicinal plants common in Kinwat forest." Ph. D. Thesis submitted to Swamy Ramanand Teerth Marathwada University, Nanded (Maharashtra). 2008
- [7] P. Neergaard, "Detection of seed borne pathogen by culture test seed". *Sci. and Tech.* vol. **1**, pp. 217-254. 1973
- [8] B. Varaprasad, K. Prasanth Kumar, K. Chandrasekhar Naidu and P. Somasekhar, "Antifungal activity of selected plant extracts against phytopathogenic fungi *Aspergillus niger* F2723". *Ind. J. Sci. Technol.* Vol.2(4) pp. 87- 90. 2009
- [9] M. Azher "Seed mycoflora of shisham (*Dalbergia sissoo* Roxb.) and their integrated management." Ph.D. Thesis submitted to Department of Plant Pathology, University Of Agriculture, Faisalabad (Pakistan). 2009
- [10] S. B. Mathur, and O. Kongsdal, "Common laboratory seed health testing methods for detecting fungi." 1st Edition ISTA. Bassersdorf, Switzerland. pp. 425. 2003
- [11] D. S. Mukadam, C. Ashok, M. S. Patil, and R. P. Anjali. The illustration of fungi. Saraswati Printing Press, Aurangabad. (M.S.) India. 2006
- [12] J. I. Pitt, and A. D. Hocking, "Fungi and food spoilage", 3rd Edn. Springer Sci. Buss. Media. pp. 53- 80. 2009