**/	Management Of The Post- Harvest Fungi By Fruit Extracts Of Wrightia Tinctoria Shamal P. Mali, S.S.Kamble, Dr.Bhagwan, M.Waghmare, G.P.Shendge.	194-197				
48/	Enumeration Of Micro-Flora On The Library Books Dr. Ayesha G Siddiqui, Miss Minakshi B, Bondge	198-201				
49	Study of Improvement in Behavior of College Students Through Sahaja Yoga Meditation Prof Rasika Beohar					
50	A Study Of Factor Affecting Job Satisfaction Employees In A Public Sector Banks Niranjana Uttamrao Machewad.					
51	Assessing The Impact Of The Mode Of Learning On Wellbeing Of College Students: A Students' Perspective. Mrs. Aditi V. Yadav	211-216				
52	Changes In The Literacy Levels Of Both Genders In Urban And Rural India Submitted By Prasanta Mujrai					
53	Grammar Competence And Academic Self Concept Of Secondary School Students Neetha V T, Prof. (Dr.) Bindu R L					
54	Socio- Economic Impact of Rural- Urban Migration- A Case Study of Bhaini Badshah Pur, Hisar District, Haryana Puja Devi, Dr. Narendra Kumar Bishnoi	231-238				
55	Human Health risks from Heavy Metals in fresh Water fish in Paschim Medinipur district of West Bengal Tanmoy Basak	239-242				
56	A study on rural people's usage levels of online banking especially in pandemic period Delma Pulikkottil	243-251				
57	Temperature Dependence Thickness, Grain size And Optical Properties of ZnO Thin Films By Spray Pyrolysis Techniques P.M. Devshette	252-255				
58	Growth and Optoelectronic properties of ZnO Thin films by Chemical Spray Pyrolysis Deposition Technique P. M. Devshette	256-258				

Management Of The Post- Harvest Fungi By Fruit Extracts Of Wrightia Tinctoria Shamal P. Mali¹ S.S.Kamble² Dr.Bhagwan³ M.Waghmare⁴ G.P.Shendge.⁵

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Abstract: Bio-pesticides and their use in the management of plant diseases has become recent trend in agriculture to be given alternate traditional plant disease management which is termed as Biological control. Management of various pathogenic micro-organisms by using the different extracts of Botanicals has been attempted and the results are found effective. Botanicals highly potential, economical, safe, biodegradable and eco-friendly for management of fungi. The concentration of the extracts, such as 0.0625, 0.125, 0.25, 0.5, 1.0 and 2.0 mg/ well of the methanol, petroleum ether and ethyl acetate fruit extract of W.tintcoria were utilized to evaluate the minimum inhibitory concentration (MIC) against fourpathogenic The results are very much positive to maximum inhibition of tested fungi. The MIC value is varied according the fungi for that standard drug were used as Amphotericin. Fruit extract of W. tincoriahave been attempted against thespecies of Neurospora, Aspergilus, Fusarium and Cladosporium for determine minimum inhibitory concentration by using different solvents system has benefitedfor management of tested fungi.

Key wards; Post harvest Pathogens, Wrightiatinctoria.

Introduction

Fungicides are widely used for management of pathogens. In this scenario increasing demand to develop,eco-friendly, bio-based, bio-control agents. It need to be focused for systemic management of fungi significantly and thereby improving the health of food grains. The Botanical constitute a promising alternative to hazards fungicides in the field of integrated disease management. In vitro activity of different concentration of Botanicals were screened for their antifungal ability against species of Fusarium, Aspegillus, Cladosporium and the results were observed complete inhibition in mycelia growth of the tested fungi which were isolated from common food grains. Isolation of fungal pathogens as per (ISTA 1966), Neergard 1977, Agarwal 1981, and identification was made with the help of different keys and as per literature by Dodge 1928, Wollen et al. 1935, Bessey 1950, Joseph Gilman 1960, Ramnath et.al. 1970, Booth 1971, Gerlach et.al. 1982, Dube 1990, Mukadam1997, Keith et.al.2002and Mukdamet.al. 2006.

Materials and Methods

Antifungalactivity: The agar diffusion method was used to evaluate the antifungal activity. Fungi which were cultured overnight at 28 °C for 72 hrs. In potato dextrose broth (PDB) used as inoculum. A final inoculum, using 100 ml suspension containing 104 spore/ml of fungus spread on potato dextrose agar (PDA) medium, respectively.

Wells of 6 mm diameter were prepared in the solid agar and Amphotericin 25, 50,100,400, 800 mg/well was used as positive controls. The test samples 0.0625, 0.125, 0.25, 0.5, 1.0, 2.0 mg/well) were applied and plates were at 28° C for 72 hours for fungal incubation for a visible growth. The antifungal activity has been expressed as diameter immm of inhibition zone and measured by using standard scale. Triplicate set were taken for each test.

Determination of Minimum Inhibitory Concentration (MIC)

Minimum inhibitory concentration (MIC) values were studied against fungi, which were determined as sensitive to the extracts in agar diffusion assay. Agar wells 6 mm diameter were filled with solution containing 0.0625, 0.125, 0.25, 0.5, 1.0, 2.0 mg/well of test extract. Minimum inhibitory concentration was defined as the lowest concentration of extracts that inhibited visible fungal growth on agar.

In order to investigate the antifungal activity of fruit extracts of Results and Discussion: Wrightiatinctoria R.Br which were utilized against four pathogenic fungi for their management. The fungi such as Neurosporacrassa, cladosporium oxysporum, Aspergillus flavusand Aspergillus niger were employed. Antifungal activity and MIC values (mg/well) of fruit extract of W. tinctoria R. Br. against different pathogenic fungi which were isolated from common food grain.

Antifungal activity and MIC values mg/well of fruit extracts of W. tinctoria R. Br. against A. flavus The fruit extracts of Wrightiatinctoria R.Br wereused againstAspergillusflavus for their growth management. Themethanol, petroleum ether and ethyl acetate extracts with their concentrations 0.0625 to 2 mg/well were used for the observation and calculation of antifungal activity and MIC The results are very beneficial to the control of tested fungi which were summarized in the table 1. As per the results the maximum inhibition zone was observed in presence of petroleum ether and ethyl acetate extracts of Wrightiatinctoria R.Br at the concentration of 2 mg/well. The minimum inhibitory concentration (MIC) was determined against Aspergillus flavus in presence of all the tested extracts of fruit of Wrightiatinctoria R.Br and minimum inhibition was found in ethyl acetate extracts at the concentration of 0.25mg/well.

Table No.1. Antifungal activity and MIC values mg/well of fruit extracts of W. tinctoria R. Br.

Sl. No.	Compounds Name	against A. flavus Concentration of Compound						
		0.0625 mg	0.125 mg	0.25 mg	0.5 mg	1 mg	2 mg	MIC mg
1	Methanol extract of Wrightiatinctoria R.Br	0	0	0	0	0.1	0.3	1
2	Petroleum ether extract of Wrightiatinctoria R.Br	0	0	0	0	0.1	0.6	1
3	Ethyl acetate extract of Wrightiatinctoria R.Br	0	0	0.1	0.3	0.5	0.6	0.25
4	Standard drug (Amphotericin)	25 μg	50 μg	100 μg	200 μg	400 μg	800 μg	MIC μg
5	Readings	0	0	0	0	0.7	1	400

Antifungal activity and MIC values mg/well of fruit extracts of W. tinctoria R. Br. against Cladosporiumoxysporum

The fruit extracts Wrightiatinctoria R.Br were used against Cladosporiumoxysporum for inhibition of their growth. The methanol, petroleum ether and ethyl acetate extracts with their concentrations 0.0625 to 2mg/well were employed for the determination and observation of antifungal and MIC

The results, the maximum zone of inhibition was exhibit in the presence of methanol extracts followed by ethyl acetate and petroleum ether fruit extracts of *W. tinctoria* R. Br.at the concentration of 2 mf/well. Table No.2, data of the results given in table No.2. The MIC was recorded against *C. oxysporium* in the presence of the fruit of *W. tinctoria* R. Br. and minimum inhibition was found in methanol and ethyl acetate extracts at the concentration of 1 mg/well

.Table No.2: Antifungal activity and MIC values mg/well of fruit extracts of W. tinctoria R.

Br. against Cladosporium Oxysporum Concentration of compound MIC 2 Sr. 1 Compounds Name 0.125mg 0.25mg 0.5mg 0.0625mg NO mg mg mg Methanol extract of W. tinctoria 1 0 0.3 0.5 0 0 0 R. Br. Petrolium ether extract of W. 2 0.3 0 0 0 0 0 2 tinctoria R. Br. Ethyl acetate extract of W. 0.4 1 0 0.1 0 0 0 3 tinctoria R. Br. 800 MIC 400 200 μg 100 μg 50 µg 25 µg Standard drug Amphotericin) 4 μg μg μg 50 0.9 1.5 0.7 1.3 0 0.2 Readings 5

Antifungal activity and MICvalues (mg/well) of fruit extracts of Wrightiatinctoria R.Br against Neurosporacrassa

The different concentration of fruit extracts of Wrightiatinctoria R.Br were used againstNeurosporacrassa for determination of their inhibition of growth. Themethanol, petroleum ether and ethyl acetate extract with their concentrations 0.0625 to 2 mg/well were used for the observation and determination of antifungal and (MIC)

The results were summarized in the table 3. The results the maximum inhibition zone was exhibited in presence of ethyl acetate fruit extract of *Wrightiatinctoria R.Br* followed by petroleum ether extracts at the concentration of 2 mg/well. It is interesting to note that the, methanol extracts did not shown any zone of inhibition up to 2 mg/well concentration. The (MIC) was recorded against *Neurosporacrassa* in presence of all the tested extracts of fruit of *Wrightiatinctoria R.Br* and minimum inhibition was found only in ethyl acetate extracts at the concentration of 1 mg/well.