International Journal of Current Microbiology and Applied Sciences ISSN: 2319-7706 Volume 4 Number 8 (2015) pp. 88-92 http://www.ijcmas.com



Original Research Article

Inhibitory Activity of Dopamine HCl and Codeine Phosphate on Aspergillus Species

A. W. Yenkar* and L. P. Dalal

P. G. Department of Botany, J. B. College of Science, Wardha (MS)-442001, India *Corresponding author

*Corresponding author	

ABSTRACT

Keywords

Dopamine HCL, Codeine Phosphate, Inhibitory activity Inhibitory activity of two drugs i.e. Dopamine HCL & Codeine Phosphate against two plants pathogenic fungi namely *Aspergillus niger* van Tieghem and *A. flavus* Link was studied. Effect of these drugs on morphological characters like colour, shape and thickness, radial or diametric growth, and conidial germination in drugs as well as in water of all these fungi was investigated. The result showed that 0.5% concentrations of these two drugs were inhibitory for *A. niger* van Tieghem and *A. flavus*.

Introduction

Aspergillus is a saprophytic fungus occurring sometimes as a parasites, Many species of Aspergillus are pathogenic and cause several diseases in plants as well as in animals (Martinelli and Kinghorn, 1994). Among plants, it causes diseases mainly on cereals which happens to be the largest source of cause of these fungi. Dopamine HCL and Codeine phosphate is alkaloid and are use in medicines (Bagul and Patel, 2001; Bhownik and Chowdhary, 1982; Shivpuri *et al.*, 1988).

The present study reports the antifungal activity of Dopamine HCL and Codeine phosphate against two plant pathogenic fungi (*Aspergillus niger* van Tieghem *and A. flavus* Link). *Aspergillus niger* and *A. flavus* is ubiquitous and its air borne spores can cause respiratory diseases known as Aspergillosis.

Materials and Methods

The culture of *Aspergillus niger* van Tieghem *and A. flavus* Link were isolated from the soil and cultured in the laboratory. Pure cultures were developed by serial dilution and by spore culture method, fungal mycelium and fungal cells. Then they were stained with cotton blue viewed and mounted in lactophenol. Microscopic observation of spore and mycelium were done by using standard method of micrometry at 400x (Irobi and Darmola, 1993).

In vitro, inhibitory activity of Dopamine HCL and Codeine Phosphate was determined with different parameters viz. Germination percentage, radial or diametric growth of the colony, colour of colony and conidia. The solution of Dopamine HCL and Codeine Phosphate was redissolved in distilled prepare water to various concentrations like 0.01%, 0.1%, 0.5%. The mention solution (conc.) was above employed for individual culture tubes containing sterile nutrient media. Potato dextrose agar (PDA) media was used for Aspergillus niger and Czapek-dox agar (CzA) media for Aspergillus flavus. These tubes were inoculated with assay disc of 2mm diameter (taken out with the help of cork borer) of pure culture of each test fungi and incubated at 37°C.

Growth of fungus was observed on 3rd, 6th and 9th days. Tubes containing drugs free medium were used as control and simultaneously a Dopamine HCL and Codeine Phosphate medium was also maintained. Three replicates of each concentration were maintained and experiment was repeated thrice.

It was observed that higher concentration of these drugs that did not permit any growth of fungus and were recorded as MIC (Maximum Inhibitory Concentration). MFC (Maximum Fungicidal Concentration) was determined by inoculating treated fungi on Codeine Phosphate and Dopamine HCL and incubating at 37°C. From the results, it was concluded that the MFC were the maximum concentration, which prevented the growth of any fungal colony on Dopamine HCL and Codeine Phosphate medium.

Result and Discussion

Tables 1 & 2 elucidate the MIC as well as effect of Dopamine HCL and Codeine growth (drugs) Phosphate on and germination percentage of Aspergillus niger and A. flavus. Luxuriant growth was observed in control. At 0.5% concentration of Codeine Phosphate and Dopamine HCL inoculum remain unchanged i.e. growth was inhibited. Therefore completely this concentration was taken as MIC (Max. Inhibitory Concentration) (Histogram I & II).

Name of	Concentration	Diameter in mm			Thickness	Colour of the	
Drugs		Petri Plate A	Petri Plate B	Average		colony	
	Control	70	70.7	70.8	+ + + + +	Carbon black	
Codeine Phosphate	0.01	34.8	34.3	34.5	+ + + +	Blackish	
	0.1	18.5	18.2	18.3	+ +	Carbon black with white shade	
	0.5	2.4	2.3	2.3	+	Faint black	
	Control	70	70.3	70.15	+ + + + +	Carbon black	
Dopamine HCL	0.01	30.5	30.3	30.8	+ + +	Carbon black with white shade	
	0.1	12.6	12.7	12.6	++	faint black	
	0.5	2.5	2.4	2.4	+	Light brownish	

Table.1 Effect of various concentration of Dopamine HCL and codeine phosphate on morphology of Aspergillus niger

Int.J.Curr.Microbiol.App.Sci (2015) 4(8): 88-92

		Diameter in mm					
Name of Drugs	Concentration	Petri Plate A	Petri Plate B	Average	Thickness	Colour of the colony	
	Control	88.5	88	88.2	+ + + + +	Dark green	
Codeine Phosphate	0.01	40.3	40.5	40.4	+ + + +	Light green	
	0.1	25.1	25.5	25.3	+ +	Greenish with yellow spot	
	0.5	2.6	2.5	2.5	+	Patches of yellow colour	
Dopamine HCL	Control	87	88	87.5	+ + + + +	Dark green	
	0.01	35.6	35.6	35.6	++++	Light green with white spot	
	0.1	11.6	11.5	11.5	+ + Greenish with whit patches		
	0.5	2.7	2.5	2.4	+	whitish green	

Table.2 Effect of various concentration of Dopamine HCL and Codeine Phosphate on morphology of Aspergillus flavus

Table.3 Effect of the duration of the treatment with Dopamine HCL on conidial germination of Aspergillus niger & Aspergillus flavus

		A	Aspergillus ni	ger	Aspergillus flavus			
Sr.	Concent	Con	idial germina	ation	Conidial germination			
No.	ration	2 Hrs.	2 Hrs. 4 Hrs.		2 Hrs.	4 Hrs.	8 Hrs.	
		Germinat	minat Germinati Germ		Germinatio	Germinatio	Germinati	
		ion %	on %	ion %	n %	n %	on %	
1	Control	90	90	90	90.5	90.5	90.5	
2	0.01	85	75	65	80	70	60	
3	0.1	50	45	40	50	43	38	
4	0.5	50	05		13	7		

Table.4 Effect of the duration of the treatment with codeine phosphate on conidial germination of Aspergillus niger & Aspergillus flavus

	Aspergillus niger				Aspergillus flavus			
Sr.	Concent	C	onidial germinat	tion	Conidial germination			
No.	ration	2 Hrs. 4 Hrs.		8 Hrs.	2 Hrs.	4 Hrs.	8 Hrs.	
		Germination	Germination	Germination	Germination	Germination	Germination	
		%	%	%	%	%	%	
1	Control	90	90	90	90.5	90.5	90.5	
2	0.01	80	70	63	85	80	70	
3	0.1	50	42	40	65	50	40	
4	0.5	14	10		12	7		





Microscopic examination of the mycelium viz., width, cytoplasmic content of Aspergillus niger and A. flavus appeared shrunken at MIC as compared to control. As evident in tables 3 & 4, reduction in germination percentage and reduction in spore size were observed at 0.5%. Present investigation revealed a significant decrease in growth and germination percentage of Aspergillus niger and A. flavus following treatment with drugs i.e. Codeine Phosphate and Dopamine HCL. The drugs showed fungicidal activity against Aspergillus niger and A. flavus The results are in conformity with the results of Sharma et al. (2007), Jain and Sharma (2007), Galagan (2005), Bagul and Patel (2001), Soni et al. (1992), Naseem and Lanjewar (1989) and Bhownik and Chowdhary (1982).

References

- Anju Sharma, Amita Dass, Paul, M.S. 2007. Antifungal effect of Neem extract on some common phytopathogenic fungi. *Ad. Plant Sci. (II)*, Pp. 357–358.
- Bagul, M.M., Patel, J.G. 2001. Evaluation of botanicals for fungi toxicity against Alternaria alternate in vitro. J. Mycil. Pl. Pathol., 31: 105–106
- Bhownik, B.H., Chowdhary, B.K. 1982.
 Antifungal activity of leaf extracts of medicinal plants of *Alternaria alternate* (Fr) Keissler. *Indian Bot. Reptr.*, 1: 2–3.
- Irobi, O.N., Darmola, S.S. 1993. Antifungal activities of crude extracts of *Mitracarpus villosus* (Rubiaceae). J. *Etanophamacol.*, 40: 137–140.
- Jain, T., Sharma, K. 2007. Antifungal potential of *Polyalthia longifolia* Benth. & Hook leaves. *Proc. Nat. Acad. Sci. India*, 77(B), I.
- Martinelli, S.D., Kinghorn, J.R. 1994. Aspergillus: 50 years on. Elsevier.
- Naseem, M., Lanjewar, R.D. 1989. Studies

on the influence of neem oil on *Aspergillus niger*. The storage fungus associated with two rice cultivars. *Indian Phytopathol.*, 42: 28–289.

- Shivpuri, A., Sharma, O.P., Jhamaria, S.L. 1988. Fungitoxic properties of plant extracts against pathogenic fungi. J. Mycol. Pl. Pathol., 27: 29–31.
- Soni, G.L., Sedha, R.K., Khana, P.K., Garcha, H.S. 1992. Growth inhibition of *Fusarium oxysporum* by phenolics compounds. *Indian J. Microbial.*, 32(1): 45–49.