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COMPARATIVE STUDY OF ANTIMICROBIAL ACTIVITY OF EUGENIA CARYOPHYLLUS AND CLEOME VISCOSA

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ABSTRACT

The present study was aimed at detecting and evaluating antimicrobial activities of Eugenia caryophyllus and Cleome viscosa known for their medicinal properties in folk medicine. The Acetone and Ethanol extract of Eugenia caryophyllus cloves shows good activity against some bacterial strains. In case of Cleome viscosa in all the extract ethanol extract showed the highest activity against all microorganism. In case of Eugenia caryophyllus, Acetone and Ethanol extract also shows good antifungal activity against A.niger, and P.carysogenum. In case of antifungal activity of Cleome viscosa ethanol extract showed highest antifungal activity.

KEYWORDS: Eugenia caryophyllus, Cleome viscosa, Antimicrobial activity, Studies.

INTRODUCTION

Eugenia caryophyllus. a plant belonging to family Myrtaceae. [1] The clove tree is an evergreen which grows to a height ranging from 10-20m, having large oval clove and crimson flower in numerous groups of terminal clusters. the flower buds are at first of a pale color and gradually become green, after which they develop into a bright red, when they are ready for collecting, cloves are harvested when 1.5-2 cm long and consist of a long calyx, terminating in four spreading sepals and four unopened petals which form a small ball in the centre.

Cleome viscosa is a plant belonging to family Capparaceae. It is a weed distributed throughout the tropics of the world and the plains of india. The plant is an annual; sticky herb with a strong penetrating odour, seed. It is known as Hurhur(Hindi), hurhuria(Bengali), Nayikkadugu (Tamil) in indian traditional medicin. Leaves are digitately compound, with 3-5 leaflets. Fruit 30-75 mm long, 3-5 mm broad, linear-oblong, erect, obliquely striated, tapering at both ends, glandular-pubescent, slender; style 2-5 mm long; seeds many, 1-1.4 mm in diam., glabrous with longitudinal striations and transverse ridges, dark brown. Cleome viscosa is highly effective in widely spectrum of disease and reported to posses antidiarrhoeal, analgesic, pharmacological, antimicrobial properties including in vitro Helicobacter pylori and wound healing activity. [3-4]

MATERIALS AND METHODS

1. Collection of clove of Eugenia caryophyllus and Seeds of Cleome viscosa.

Clove of Eugenia caryophyllus and Seeds of Cleome viscosa were collected from area around kargi chock, Dehradun during the month of Oct to Dec.The collected plant material was washed with water to remove mud and other undesirable material and dried under shade.

2. Extraction of clove of Eugenia caryophyllus and Seeds of Cleome viscose.

The collected plant Material was washed with water to removed other undesirable material and dried under shade. The air-dried clove (300 gm) of Eugenia caryophylluswere crushed. The crushed clove extracted with different solvents of increasing polarity viz. acetone, ethanol at room temperature. The extract was evaporated till dryness to obtain residue. These extracts were concentrated under reduced pressure. The extract was used for used for antimicrobial activity.

The collected plant Material was washed with water to removed other undesirable material and dried under shade. The air-dried seed (300 gm) of Cleome viscosa were crushed. The crushed seed extracted with methanol at room temperature. The extract was evaporated till dryness to obtain residue. These extracts were concentrated under reduced pressure. The extract was used for used for antimicrobial activity.

4. Anti-microbial activity

The **anti-microbial activity** of the clove of Eugenia caryophyllus and Seeds of Cleome viscosa was carried out. The clove and seeds extract were screened for anti bacterial and anti fungal activities.

Arti et al.

Anti bacterial activity of clove and seeds extract

In this study, the anti bacterial activity was studied against the micro organism and the

bacterial cultures used in the study were.

1. Pseudomonas aerugisa

2. Klebsiella species

3. B.cereus

These bacterial cultures were maintained on nutrient agar slants at first being incubated at 37⁰

c for about 18-24 hours and then stored at 4⁰ c as stock for antibacterial activity. Fresh

cultures were obtained by transferring a loop full of cultures into nutrient broth and then

incubated at 37°c overnight. To test antibacterial activity, the well diffusion method used.

Culture media preparation

The microbiological media prepared as standard instruction provided by the HI-Media

Laboratories, Mumbai. The media used for anti-bacterial activity Muller- Hinton Agar

(MHA) and Nutrient broth (NB). They were prepared and sterilized at 121°C at 15 psi for 15-

30 minutes autoclave.

Plate preparations

25 ml of pre autoclaved Muller-Hinton agar (MHA) was poured into 90 mm diameter pre

sterilized petri-plates. These petri-plates were allowed to solidify at room temperature.

Well diffusion method

After the plated solidified the freshly prepared microbial growth culture suspension (about

20µl) was spread over the Muller – Hinton agar (MHA) media using L shaped sterilized glass

spreader separately under the aseptic condition using laminar air flow. Then well were made

in each plate with the help of borer of 8 mm diameter. In these well, about 100µl of each

leaves extracts individually was loaded. This method depend upon the diffusion of leaves

extracts from hole through the solidified agar layer of petri-dish to such an extent that the

growth of added micro organism is prevented entirely in a circular area or Zone around the

hole containing leaf extract.

Incubation: Petri plates were incubated for overnight at $37^{\circ}\text{C} \pm 0.5^{\circ}\text{C}$ in the incubator.

Inhibition Measurement of zone of inhibition

After incubation, the diameter of clear zone of incubation produced around the well or holes were measured in mm by ESR Tube and compared with standard drug.

RESULTS

Table-1: Antibacterial activity of the extract of Eugenia caryophyllus (clove).

Test organism	Acetone	Ethanol)
Pseudomonas	20	18
aerugisa	(36mm)	(36mm)
Klebsiella species	21	24
	(35mm)	(35mm)
B.cereus	20	20
	(-)	(-)

Table-2: Antibacterial activity of the extract of Cleome viscosa Seeds

Test organism	Acetone	Ethanol)
Pseudomonas aerugisa	9	15
Klebsiella species	12	14
B.cereus	10	17

Table- 3: Antifungal activity of the extract of Eugenia caryophyllus clove.

S. No.	Test organism	acetone	ethanol
1	A.niger	20	20
2	Penicillium chrysogenum	25	23

Table-4: Antibacterial activity of the extract of Cleome viscosa Seeds

S. No.	Test organism	acetone	ethanol
1	A.niger	8	13
2	Penicillium chrysogenum	10	12

DISCUSSION

In case of **Eugenia caryophyllus** Acetone and ethanol extract of clove showed similar activity against B.cereus. Ethanol extract of clove showed higher activity against Klebsiella. Acetone extract of clove showed higher activity against Pseudomonas aerugisa. In case of seeds of Cleome viscosa acetone and ethanol extract showed antibacterial activity against all tested microorganisms. Among all the extract ethanol extract showed the highest activity against all microorganisms.

In case of **Eugenia caryophyllus** Acetone and ethanol extract of clove showed similar activity against Aspergillus niger and P.chrysogenum. In case of Cleome viscosa antifungal activity ethanol extract showed highest antifungal activity against P.chrysogenum and Aspergillus niger.

REFERENCES

- 1. Sulekha M., Yadav S., Yadav S., Kumar N.R., J. Chem. Pharma. Res., 2009; 1(1): 102-
- 2. Joseph S, Peter KV. Econ Botany, 1985; 39(1): 68-73.
- 3. Bajpai A, Ojha J.K, Sant H.R. Medico botany of the Varanasi district.Int J Pharm, 1983; 33(2): 172-176.
- 4. Kokate C.K., Plant constituents: Practical Pharmacognosy, Delhi; Vallabh Prakashan: 2005; 107-111.