(Research Article)

E-ISSN: 0975-8232; P-ISSN: 2320-5148



INTERNATIONAL JOURNAL OF PHARMACEUTICAL SCIENCES AND RESEARCH



Received on 13 July, 2017; received in revised form, 27 November, 2017; accepted, 13 December, 2017; published 01 April, 2018

ANTIMICROBIAL ACTIVITY OF COCONUT SHELL OIL

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Keywords:

Coconut shell oil, Anti-bacterial activity, Anti-fungal activity

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ABSTRACT: Coconut shell oil was extracted from raw coconut shells, collected from Thiruvallur District. These were extracted using various solvents based on the polarity which includes ethanol, chloroform, acetone, petroleum ether and aqueous. The antibacterial activity against the growth of Staphylococcus aureus, Enterococcus faecalis, Escherichia coli, Klebsiella pneumonia, Pseudomonas aeruginosa and Salmonella typhi was done. The antifungal activity against the growth of organisms such as Epidermophyton floccosum, Aspergillus niger, Penicillium, Microsporum canis, Candida albicans and Aspergillus flavus were tested. The extracts were compared with standards like novobiocin, amoxillin and ketoconazole for antibacterial and antifungal activity respectively. The results indicate that zone of inhibition increased when increasing the concentration of the extracts. Among the five extracts of coconut shell oil the ethanolic extract exhibited maximum antibacterial activity against six bacterial strains. Ethanol and petroleum ether extract showed maximum inhibitory activity against *Epidermophyton* and *Candida albicans*. This study showed that coconut shell oil possesses antimicrobial activity.

INTRODUCTION: Cocos nucifera Linn. (family Arecaceae) commonly known as coconut is an important fruit crop in tropical countries ¹ and produced in 92 countries worldwide taking 10 million hectares of land. It is grown for its several beneficial products such as fiber, timber, functional food and oil. Coconut is mainly cultivated as an oil crop that is rich in lauric acid with a wide variety of other uses in addition to commercial oil production ². Coconut water and kernel were used to treat abscesses, asthma, bronchitis, burns, constipation, dysentery, earache, flu ³, diarrhoea, hemostat, pediculicide ⁴. In India, the inflorescence infusions of coconut are used for the oral treatment of menstrual cycle disorders ⁵.



DOI:

10.13040/IJPSR.0975-8232.9(4).1628-31

Article can be accessed online on: www.ijpsr.com

DOI link: http://dx.doi.org/10.13040/IJPSR.0975-8232.9(4).1628-31

White flesh of the coconut fruit is used to treat fever and malaria ⁶, it is also used to relieve skin rashes caused by HIV infection ⁷. Coconut Shell oil had higher number of secondary metabolites and they reported to have many biological and therapeutic properties ⁸.

MATERIALS AND METHODS: The samples of coconut shells were collected from the local market in Thiruvallur district, Tamil Nadu, India. They were sundried for few days to remove the moisture content. Then it is broken into small pieces and ground into course powder. Ground coconut shells (250 g) were heated in an earthen pot for a span of 3 hours giving a yield of 25 cc of oil. The oil was extracted with ethanol, chloroform, acetone, petroleum ether and aqueous (v/v) separately and preserved in airtight container for further studies. The bacterial and fungal cultures were procured from the Post Graduate and Research Department of Microbiology, Madras Christian College, Tambaram, Chennai, Tamil Nadu, India and subcultured.

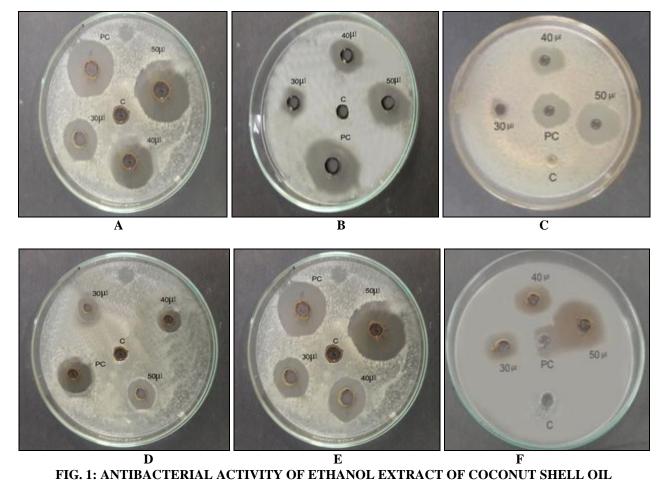
E-ISSN: 0975-8232; P-ISSN: 2320-5148

Antibacterial Susceptibility Test: The well diffusion method on Muller Hinton Agar (MHA) medium was used to screen the antibacterial activity ⁹. The MHA plates were prepared by pouring media into sterile Petri plates and bacterial culture inoculated by spread plate technique. Wells of 6 mm diameter was bored and extracts were applied to different wells in serially increasing volumes of 30 μl, 40 μl and 50 μl. Dimethyl sulphoxide (DMSO) was used as negative control, novobiocin and amoxicillin were used as positive control for Gram (+) and Gram (-) bacteria respectively. The plates were incubated at 37 °C for 24 h ¹⁰. The diameter of inhibition zones was measured and tabulated.

Antifungal Susceptibility Test: All the extracts were screened for antifungal activity against pathogenic fungal strains such as *Epidermophyton floccosum*, *A. niger*, *Penicillin*, *Microsporum canis Candida albicans* and *A. flavus*. The fungi were

grown on Potato Dextrose Agar (PDA) medium. The fungal mycelial suspension was spread on PDA plates. After solidification 6mm diameter wells were made and extracts applied in volumes of 30 μ l, 40 μ l and 50 μ l. DMSO served as negative control and ketoconazole (10 μ g) used as positive control. The plates were incubated at 28 °C for 48-72 hours. The diameter of the zone of inhibition was measured and the results were recorded.

RESULTS AND DISCUSSION: From the well diffusion method it is observed that the coconut shell extract exhibited higher antimicrobial activity against all the tested bacteria (**Table 1**). The highest zone of inhibition against *S. aureus, Enterococcus faecalis, E.coli, Pseudomonas* and *Salmonella* formed in ethanol extract and their zone of inhibition was higher than the control shown in **Fig. 1**. The petroleum ether extract showed more inhibition zone against *Klebsiella*.



A. Staphylococcus aureus, B. Enterococcus faecalis, C. Escherichia coli, D. Klebsiella pneumonia, E. Pseudomonas aeruginosa, F. Salmonella typhi,

TABLE 1: ANTI-BACTERIAL ACTIVITY OF COCONUT SHELL OIL

Test	Zone of inhibition in mm																			
organisms		Eth	nanol		Chloroform					Ace		Р. е	ther		Aqueous					
	C 30 40 50			С	30	40	50	С	30	40	50	С	30	40	50	С	30	40	50	
		μl	μl	μl		μl	μl	μl		μl	μl	μl		μl	μl	μl		μl	μl	μl
S. aureus	30	21	27	34	30	14	20	22	30	12	20	29	30	20	22	28	30	-	10	13
Enterococcus	26	12	15	19	26	-	15	17	26	11	14	18	26	-	12	13	26	12	13	16
E. coli	24	22	26	32	24	11	14	16	24	12	13	19	24	10	14	17	24	11	13	16
Klebsiella	28	14	20	25	28	-	-	9	28	14	19	22	28	21	27	30	28	9	13	16
Pseudomonas	25	24	27	34	25	10	17	22	25	11	19	22	25	18	27	31	25	9	14	16
Salmonella	30	20	26	29	30	17	22	24	30	14	18	21	30	-	10	13	30	-	-	12

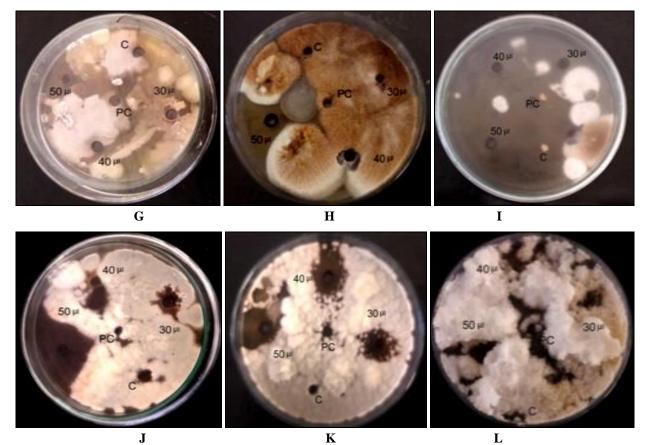


FIG. 2: ANTIFUNGAL ACTIVITY OF ETHANOL EXTRACT OF COCONUT SHELL OIL

G. Epidermophyton floccosum, H. Aspergillus niger, I. Penicillium, J. Microsporum canis, K. Candida albicans, L. Aspergillus flavus (C-Control, PC-Positive control)

TABLE 2: ANTIFUNGAL ACTIVITY OF COCONUT SHELL OIL

Test		Zone of inhibition in mm																		
organisms	rganisms Ethanol			Chloroform				Acetone					Р. е	ther		Aqueous				
	C	30	40	50	С	30	40	50	С	30	40	50	С	30	40	50	С	30	40	50
		μl	μl	μl		μl	μl	μl		μl	μl	μl		μl	μl	μl		μl	μl	μl
Epidermophyton	18	14	20	24	18	-	-	-	18	10	12	15	18	17	22	26	18	-	9	12
A. niger	23	10	21	23	23	21	28	32	23	-	-	-	23	-	-	-	23	14	16	20
Penecillium	22	-	10	12	22	-	-	-	22	16	21	24	22	19	23	28	22	-	-	10
Microsporum	20	-	-	12	20	-	-	-	20	-	-	10	20	19	21	28	20	-	-	12
C. albicans	21	12	15	16	21	15	21	23	21	15	18	21	21	16	19	21	21	11	15	18
A. flavus	24	-	-	8	24	-	-	-	24	-	13	16	24	14	18	24	24	-	-	10

The extracts exhibit inhibitory activity against all the tested fungal strains such as *Epidermophyton floccosum*, *A. niger*, *Penecillium*, *Microsporum canis*, *Candida albicans* and *A. flavus* shown in **Fig. 2** Inhibition zones of each fungal species is given in **Table 2**. The zone of inhibition of chloroform extract on *Candida albicans* and *A.*

niger were high at all concentrations. From this study it is concluded that the petroleum ether extract of coconut shell oil was more active against the fungal pathogen especially in *Epidermophyton floccosum* and *Candida albicans*. The results of the study clearly showed that the effectiveness of the

extract increases with increasing concentration as observed from the zone of inhibition.

CONCLUSION: Coconut shell oil regarded to have high therapeutic potential for drug preparation. The current study provided useful information concerning the various medicinal properties to protect humans against common skin diseases. Since the shells of *Cocos nucifera* and cashew nut discarded as waste - indeed comprising a major portion as agro waste in our country. Cashew Nut Shell oil has various phytochemical contents and it can be used as an antibacterial agent ¹¹. This study reveals that Coconut shell oil can be used to produce antibacterial and antifungal agents and also it could serve as a cheap source of antimicrobial compounds replacing the conventional chemical antibiotics which give side effects to the mankind.

ACKNOWLEDGEMENT: Nil

CONFLICT OF INTEREST: Nil

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How to cite this article:

Pritha SDSJ and Karpagam S: Antimicrobial activity of coconut shell oil. Int J Pharm Sci & Res 2018; 9(4): 1628-31. doi: 10.13040/IJPSR. 0975-8232.9(4).1628-31.

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