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Research article

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Synthesis anti-bacterial and anti-fungal activity of curcumin and its derivatives

Deepthi, P.Meena, V.Kartheek, R.Swapna, D. Ravali Reddy

¹OPJS University, Churu, Rajgarh, Churu, Rajasthan 331303, ²Maharishi Markandeshwar University, MullanaAmbala Haryana-133207 *Corresponding Author: Deepthi Email id: meenapuppala96@gmail.com

ABSTRACT

Three curcumin derivatives having modification in active methylene group (1,3) and keto groups (2) where successfully synthesized. Derivative-1 synthesized from curcumin with 2-hydroxy benzaldehyde using EDTA and Chloroform. Derivative-2 synthesized from curcumin with 4-aminophenol using pyridine and ethanol. Derivative-3 synthesized from curcumin with N,N-dimethylamino benzaldehyde using EDTA and chloroform. The substitution on the active methylene site of curcumin increases the anti-bacterial and anti- fungal behaviour. While comparing compound 2 which have more potent bacterial activity with compound-1 and compound-3, the former shows higher scavenging activity. Finally the yield was found to be 63%.

INTRODUCTION

Curcumin is a bright yellow colour chemical produced by some plants. It is a principal curcuminoid of turmeric, a member of ginger family. It is used as herbal supplement, cosmetics ingredient, food flavouring and food coloring. Curcumin is a component of the Indian spice turmeric (curcumin longa), a type of ginger. It acts as powerful antioxidant and anti-inflammatory with highest bio availability[1-5].

PHYSICAL PROPERTIES OF CURCUMIN Source

The main source of curcumin is zingiberaceae family plant curmin longa, curcumin zedoaria.

Colour

Yellow crystalline powder

Odour

Slightly bitter taste

Solubility

Insoluble in water and ether, soluble in in ethanol, glacial acetic acid, propylene glycol and alkali solution.

CHEMICAL PROPERTIES

MEDICINAL USES OF CURCUMIN

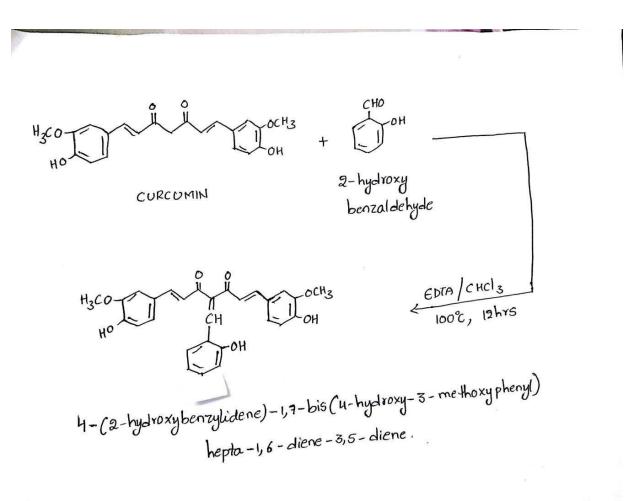
It is medicinally used as:

- 1. Anti-inflammatory drug
- 2. Anti-depressants
- 3. Chemotheraphy
- 4. Anti cogulant
- 5. Pain killer
- 6. Diabetic drug
- 7. Arthritis medication
- 8. Inflammatory bowel disease drug
- 9. Cholestrol drug
- 10. Steriods

ISOLATION OF CURCUMIN

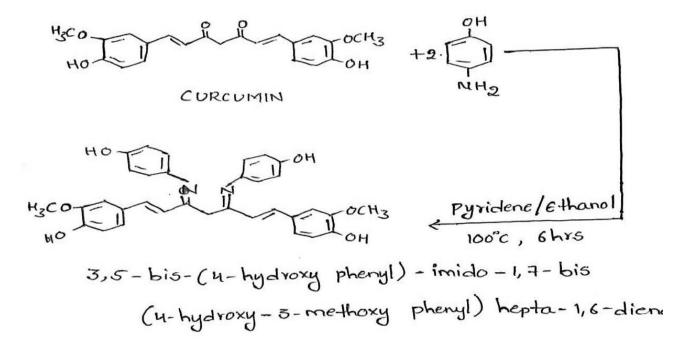
Curcumin is isolated from the turmeric powder using ethanol by extraction process.10grams of curcumin and 150ml of ethanol was taken in a round bottomed flask and stirred for 2 minutes and refluxed for 90minutes. Then the product is filtered and organic layer was separated. Then the collected organic layer was kept for extraction until the solid product was obtained and finally the extracted curcumin was collected and weighed [6. 7].

Synthesis of 3,5-bis-(4-hydrox y-phenyl)-imido-1,7-bis(4hydroxy-3-methoxy phenyl)hepta-1,6-diene SCHEME

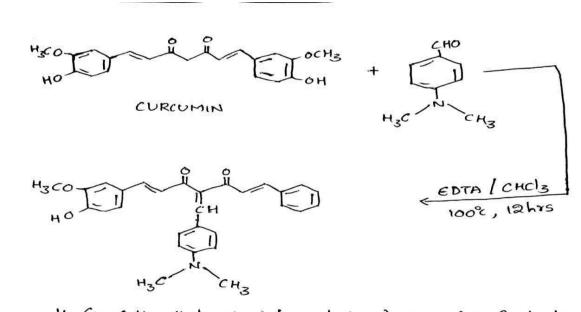




SCHEME







4- (4- (dimethylamino) benzylidene) - 1,7- bis (4-hydroxy-3-methoxyphenyl) hepta-1,6-diene-3,5-diene.

RESULTS

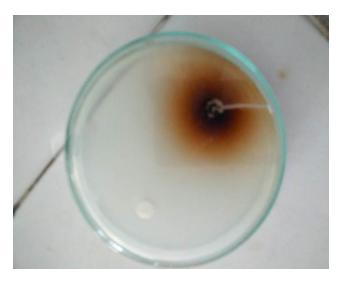
Anti-Bacterial Activity of Curcumin Derivatives						
S.No	Bacterial Zone of Inhibition					
1	SAMPLE	DERIVATIVE-1	DERIVATIVE-2	DERIVATIVE-3		
2	AMOXACILLIN	0.1	1.5	1.1		
3	SAMPLE PRODUCT	0.1	0.7	1.1		

	Anti-Fungal Activity of Curcumin Derivatives							
S.No Fungal Zone Of Inhibition								
1	SAMPLE	DERIVATIVE-1	DERIVATIVE-2	DERIVATIVE-3				
2	KETOCONAZOLE	2.2	1.4	0.9				
3	SAMPLE PRODUCT	1.1	1.1	0.2				

CONCLUSION

The 2nd derivative (3, 5-bis-(4-hydroxy-phenylimido-1, 7-bis (4-hydroxy-3-methoxyphenyl) hepta-1,6-

diene) has more potent anti- bacterial activity when compared to derivative 1 and 3.



The 3rd derivative (4-(4-dimrthylamino)benzylidene)-1,7-bis(4-hydroxy-3-methoxyphenyl)hepta-1,6-

diene3,5diene) has more potent anti-fungal activity when compared to derivative 1 and 2.

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REFERENCES

- [1]. D.Shahwar, U.Sana and N.Ahmad, Turk. J.Chem, 37, 2013, 262.
- [2]. S.Shishodha, G.Sethi and B.B.aggarwal, Ann.N.YAcad. Sci., 1056, 2005, 206.
- [3]. S. Mishra, U.Narian, R. Mishra and K.Mishra Bioorg. Med.Chem, 13, 2005, 1477.
- [4]. S.Sumathi, P. Tharmarj, C.D.Sheela and R. Ebenzer, J.Coord., 65, 2012, 506.
- [5]. J.Lal, S.K.Gupta, D.Thavaselvam and D.D.A garwal, Eur.J.med.chem, 64, 2013, 579.
- [6]. M.Cousins, J. Adelberg, F.Chen and J.Rieck, Ind. Crops prod., 25, 2007, 129.
- [7]. D.Yasudha, K. Takahashi, T.Ohe, S. Nakamura and T.Mashino, bioorg.Med.chem, 21, 2013, 7709.
- [8]. A.R.Garrett, E.G.Weagel, A.D.Martinez, M.Heaton, R.A.Robbison and K.L.O Neil, Food chem., 158, 2014, 490.
- [9]. M.Kelkel, C.Jacob, M.Dicato and .Diederich, Molecules, 15, 2010, 7035.
- [10]. K.I.Priyadarshini, D.K.Maity, G.H.Naik, M.S.Kumar, M.K.Unnikrishnan, J.G.Satav and H.Mohan, Free radic. Boil.Med., 35, 2003, 475.