SYNTHESIS & CHARACTERIZATION OF A NOVEL SCHIFF BASE FOR ITS BIOLOGICAL ACTIVITY

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ABSTRACT

Schiff base synthesis is usually acid-catalyzed and usually require refluxing the mixture of aldehydes (or ketone) and amine in organic medium. However, assistance of stirring for synthesis of Schiff base is introduced. In the present study stirring promoted condensation reaction of vanillin and m-toluidine are displayed. The method was also compared with conventional method for determination of production efficiency and production economic. Characterization of these Schiff base were done by TLC and IR spectra. From the study, it was concluded that this method is very rapid, reliable and economic method for production of schiff base.

Keywords: -Vanillin, m-toluidine, Ethyl alcohol, Glacial Acetic acid.

Introduction

- Ugo (Hugo) Joseph Schiff ,one of the founders of modern chemistry, was born in Frankfurt on the 26 April 1834, into a wealthy Jewish family of merchants, Joseph Moses Schiff(1784–1852) and Henriette Trier (1798–1888).
- He studied chemistry and physics in Frankfurt with Professors Böetteger and Löwe, and continued his studies in Göttingen, where he got his

degree in 1857 under the supervision of professorWölher.

- A Schiff base is a nitrogen analog of an aldehyde or ketone in which the C=O group is replaced by C=N-R group.
- It is usually formed by condensation of an aldehyde or ketone with a primary amine according to the following scheme:

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Primary amine Aldehyde or ketone

Schiff bas

Where R, may be an alkyl or an aryl group.

 Schiff bases that contain aryl substituents are substantially more stable and more readily synthesized, while those which contain alkyl substituents are relatively unstable.





Experimental Methods

All the chemicals were purchased from Loba company in high quality standard & purity.

- The product was recrystallised & melting point was noted using basic melting point apparatus.
- Then characterization was done by IR spectroscopy.

Procedure for preparation of Schiff base:-

The Schiff base was prepared using two ways:-

- 1. Microwave Irradiation.
- 2. Manual Stirring.
- 1. Microwave Irradiation:-
- Take 1gm of Vanillin & 1ml of mtoluidine in a clean dry 50ml conical flask. Add about 1.5ml of ethyl alcohol to the flask.
- Then irradiate this in a microwave with 360 watts for 10 mins. Then pour this solution to a ice cold water.
- Filter it &dry.Determine its melting point.
- 2. Manual stirring:-
- Take 7 gm of Vanillin & 7ml of mtoluidine in a clean beaker.
- Add about 10 ml of ethyl alcohol to the solution then stir it manually for 5-10 mins.
- A sticky substance is observed.Add a drop of glacial acetic acid to it.
- Then orangish yellow substance is obtained.
- Reaction:-

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Schiff base

The antibacterial activity of newly synthesized Schiff'sbase was conducted against Gram positive bacteria i.e

<u>Staphylococcus</u> <u>aureus</u>, <u>Corynebacterium</u> <u>diphtheric</u> andGram negative bacteria i.e. <u>Escherichia coli</u>, <u>Salmonella</u>

typhibyusingDITCHPLATETECHNIQUE.Septran tablet was employedasreferencestandardtocomparetheresults.STERILEMUELLERHINTONAGAR was used forthe screening methods.

Antibacterial activity

ANTIMICROBIAL ACITIVITY

All the synthesized test compound were screened for their anti-bacterial activity by Ditch Plate Technique Using (Gram positive) organism Such as Staphylococcus aureus and Coryrobacteriu diphtherie (Gram negative) Escherichia coli And Salmonellatyphi. Media used is Sterile Mucller Hinton Agar and Inhibition condition is 37^{0} C For 24 Hours

Results were recorded as zone of inhibition as shown in table.

Comple						
Sample	Amount	Escherichia	Salmonella	Staphylococcus	Coryrobacterium	Conclusion
	added	coli	typhi	aureus	diphtherie	
Culture used	added	COII	typin	durcus	aipitaiene	
	to ditch					
	in gm					
	0					
Positive	0.05	No Growth on	No Growth on	No Growth on test	No Growth on test	
control	0.05	test culture	test culture	culture	culture	Compound has
control		test culture	test culture	culture	culture	Compound has
						antibacterial
(Septran	0.1	No Growth on	No Growth on	No Growth on test	No Growth on test	activity and is
tablet)	0.1	test culture	test culture	culture	culture	
						active against
Vanillina		No Crowth on	No Crowth on	No Crowth on tost	No Crowth on tost	both gram
vannne	0.05	No Growin on	No Growin on	No Growin on test	No Growin on test	positive as well
		test culture	test culture	culture	culture	positive as well
+ m-						as gram negative
Toludine	0.1	No Growth on	No Growth on	No Growth on test	No Growth on test	bacteria
				•		
		test culture	test culture	culture	culture	

1. Growth on ditch

2. Growth around ditch

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Sample Culture used	Amount added to ditch	Escherichia coli	Salmonella typhi	Staphylococcus aureus	Coryrobacterium diphtherie	Conclusion
Dogitivo	in gin	No Crowth on	No Crowth on	No Crowth on tost	No Crowth on tost	
control	0.05	No Growth on test culture	No Growth on test culture	No Growth on test culture	No Growth on test culture	Compound has
(Septran tablet)	0.1	No Growth on test culture	No Growth on test culture	No Growth on test culture	No Growth on test culture	activity and is active against
Vanilline + m- Toludine	0.05	No Growth on test culture	Growth on test culture	No Growth on test culture	No Growth on test culture	both gram positive and gram negative
Totalit	0.1	No Growth on test culture	No Growth on test culture	No Growth on test culture	No Growth on test culture	Jaciena

RESULT AND DISCUSSION

The melting point observed was 85°C of recrystallised product & yield was about 75-80%.



- The sample was characterized by IR & also for biological activity.
- ≻ <u>By IR</u>:-
- The Infra red spectra of the synthesised compounds were scanned in the range of 4000-600cm⁻¹ in KBr pellet.
- A narrow band at 2260-2210cm⁻¹ is attributed to C=N stretch.
- A band at 1370-1350cm⁻¹ is attributed to C-H bend.
 - A band at 1335-1250 cm⁻¹ is attributed to C-N stretch.

CONCLUSION:-

Biological Activity:-

- At 0.05 concentration it showed partial solubility in water.
- The compound showed antibacterial activity & is active against Gram positive as well as Gram negative bacteria at both the concentrations.

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