

(Thermal, Antimicrobial, Solubility) - Studying of Diazepine Compounds

Dr. Nagham Mahmood Aljamali^{*1}, Zaid Noor Obaid Al-Husseini², Sabreen Ali Abdalrahman³

^{1*}Department of Chemistry, Faculty of Education for Girls, Iraq.

²Ministry of Education, Babylon Directorate of Education, Yahya Bin Zaid Preparatory School, Iraq.

³B.Sc in Chemistry, Chemistry Department, Education College for Girls, Iraq.

*Corresponding Author E-mail: dr.nagham_mj@yahoo.com

Article Received: 30 April 2018

Article Accepted: 29 July 2018

Article Published: 27 August 2018

ABSTRACT

In the present paper we complete first work (first paper-in first reference) through studying of thermal behavior of aldamine compound, oxazepine compound, diazepine compounds which prepared and identified in first paper with variety techniques methods in organic chemistry field, then we studied microbial assay in this paper on types of bacteria, studying of solubility our compounds in types of solvent, some physical properties. Diazepine compounds have high spectrum of pharmaceutical drugs and medical fields.

Keywords: Diazepine, oxazepine, Schiff base, antimicrobial, pericyclic.

1. INTRODUCTION

Pericyclic reactions include changes of bond in a cyclic system of atoms ,bonds are formatted but other bond broken in a cyclic transition state (T.S) of compounds, which means that there are no intermediates formed in the course of the reaction⁽¹⁻³⁾.



Fig (1): Mechanism of Oxazepine and Diazepine in Pericyclic Reaction

Pericyclic reactions act an important type of concerted process including pi-systems; rearrangement of the electrons due to (sigma and pi-bonds) to simultaneously break hen form which gives stereo control of the products⁽⁴⁻¹⁷⁾. This type of reaction (pericyclic reaction) involved three types of reactions: electrocyclic reaction, cycloaddition reaction and sigmataropic rearrangement. The transition state (cyclic) go to an arrangement of components throughout the course of the reaction⁽¹⁸⁻²⁴⁾. The diazepine compounds have



Indo-Iranian Journal of Scientific Research (IIJSR)

(Peer Reviewed International Journal), Volume 2, Issue 3, Pages 64-73, July-September 2018

many applications⁽²⁵⁻³⁴⁾ in medical drugs, and in synthesis of pharmaceutical and biochemical compounds⁽³⁵⁻⁴⁶⁾.

2. EXPERIMENTAL PART:

In the past studying⁽¹⁾ seven compounds from (aldamine, oxazepine, diazepine were formatted, now we completed our work by studying effect of compounds on classes of bacteria represented through scanning of three concentrations ,which explained in results according to data in table (1).



Scheme.1: Prepared Compounds { 1-7 }

Biological Assay

Screening of Microbial activity for imine, oxazepine, diazepine derivatives have been tested with agar through past procedures $^{(33-36)}$. The antimicrobial activities have done at three concentrations (3, 5, 7 micro gram) concentrations in (DMSO) solvent with class of bacteria (**bacteria** - *Salmonella .typhi*},) and (*bacteria K. Pneuomona*). These bacterial strains were incubated for 24 hrs at 37°C.



Indo-Iranian Journal of Scientific Research (IIJSR) (Peer Reviewed International Journal), Volume 2, Issue 3, Pages 64-73, July-September 2018



Picture.(1): bacteria Salmonella.typhi



Picture.(2); bacteria K.Pneuomona

3. RESULTS AND DISCUSSION

In past work¹, we produced aldamine, oxazepine, diazepine derivatives while in the present work completed the second part from our paper, we will study Thermal analysis then Activity against types of microbes, and solubility of compounds in a variety solvents with some physical properties.

Biological Studying⁽⁴⁴⁻⁴⁶⁾:

The scanning of microbial activity of the bacteria were involved work on classes of bacteria to study the microbial activity according to procedures⁽⁴⁴⁻⁴⁶⁾ for {(bacteria- *K. Pneuomona*) and (bacteria *Salmonella* .*typhi* }, Table (1) appeared the level of inhibition for measured compounds in (mm) towards the selected



Indo-Iranian Journal of Scientific Research (IIJSR)

(Peer Reviewed International Journal), Volume 2, Issue 3, Pages 64-73, July-September 2018

bacteria. The antibacterial results are summarized in table (1) gave good activity against classes of bacteria, which improved evidence for the results that the activity of all diazepine compounds have high microbial activity than aldamine and oxazepine compounds which inhibit the growth of bacteria. The prepared compounds [7, 4] have higher activity than other compounds which due to presence of sulfur atoms in their structures which represented by (sulphadiazine compound, thiazole ring) in structure ^(34, 36) which appeared in high inhibition ⁽³³⁻³⁷⁾ in wall of cell of bacteria.

	(average of three Measurements)	(average of three Measurements		
Compounds	Salmonella .typhi	K. Pneuomona		
[1]	4<	4<		
[2]	4<	4		
[3]	4	6		
[4]	8	10		
[5]	8	10		
[6]	6	8		
[7]	10	12		

Table(1): Antimicrobial Assay of Compounds (Inhibition Zone in (mm))as average of three Concentrations (3, 5, 7 micro gram)



Photo.(1): Inhibition zone on *bacteria K. Pneuomona*





Photo.(2): Inhibition zone on Salmonella.typhi

Thermal – Analysis of Compounds:

The measurements of compounds showed information of stability for compounds against high temperatures in figures (2- 6):



Fig (2): Thermal Curve of Compound {2}



Indo-Iranian Journal of Scientific Research (IIJSR)

(Peer Reviewed International Journal), Volume 2, Issue 3, Pages 64-73, July-September 2018













Fig (5): Thermal Curve of Compound $\{5\}$



Fig (6): Thermal Curve of Compound {6}



Indo-Iranian Journal of Scientific Research (IIJSR) (Peer Reviewed International Journal), Volume 2, Issue 3, Pages 64-73, July-September 2018



Fig (7): Thermal Curve of Compound {7}

Solubility of Compounds in Chemical Solvents:

All compounds in this paper tested with series of chemical solvents according to (nature of solvent, polarity of solvents, activity of functional groups in compounds) in our compounds in this study, the results summarized in Table (2).

Reagents	Solvents						
	C ₂ H ₅ OH	Methanol	Dioxane	Hexane	Benzene	Toluene	
(1)	+	+	+	-	-	-	
(2)	+	+	+	-	-	-	
(3)	+	+	+	+	+	+	
(4)	+	+	+	+	+	+	
(5)	+	+	+	+	+	+	
(6)	+	+	+	+	+	+	
(7)	+	+	+	+	+	+	

Table (2): Behavior of Reagents in Many Solvents

REFERENCES

[1]. Nagham Mahmood Aljamali, "Synthesis and characterization of new 1,3-oxazepine ., diazepine "., Asian J. Research Chem. ,7, 12,1067-1104, 2014.

[2]. Leach M. R., "Pericyclic Reaction Chemistry", page.1 (2007).



[3]. Alwan, A.S; Abou, S.Z., "IRAQI Drug Guide. First edition national", board for the selection of drugs. IRAQ., (1990).

[4]. Wilson C. O. and O. Givold, "Text book of Organic Medicinal and pharmaceutical Chemistry", 5th Ed., Pitman Medical Publishing Co. LTD, London copy right. C by. J. B. Lippin Cott Company (1966).

[5]. Marry B.A.; "Organic Reaction Mechanism", Ch. 1, Jon Willey, sons, (2005).

[6]. Clark, W.G.; Brater, D.C.; Johnson, A.R. Medical pharmacology Goths. Introduction to chemotherapy mechanisms of antibacterial. International edition., (1992).

[7]. Cruickshank, R.; Duguid, J.P.; Matmion B.P.,"A Guide to the laboratory diagnosis and control of infections. 12Edition . London Vol:1, 1973.

[8]. Finkelstein,R;.Kassis,E.;Reinhertis,G.,"Community a quaired urinary tract infection in adult. Journal hospital infection, 38:193-202., 2005.

[9]. Brewster R. E., W. E. McEwen; "Organic Chemistry", Ch. 30ed Ed., p.638, (1971).

[10]. Gebe, S.S., "Asymptomatic bacteria. Ethiopia-medical Journal. 36(3):158-192., 2006.

[11]. Sykes P.; "Agide Book to Mechanism in Organic Chemistry", 5th Ed., Longman, (1974).

[12]. Guold, J.C .,"The comparative bacteriology of acute and chronic urinary tract infection. Urinary tract infection . 8th Edition . Nephrology series. England., 1973.

[13]. Joseph, L.H., "Chemotherapeutic drugs. Clinical pharmacy and thetapeutics"., 3th Edition., 1998.

[14]. Kafaf P.A.," Gentic study on antibiotic resistance of some gram- negative bacteria isolated from urinary tract infection . Thesis, M.Sc. College of science. Almustansiriya university., 2000.

[15]. Knight J.A.," Encyclopedia of genetics. Second edition". New York., 1999.

[16]. Mims H.M., "Antimicrobial agents and chemo therapy" .Medical microbiology Third edition., 1995.

[17]. Sachadev, K.N., "Examination of urine clinical pathology and bacteriology"., seventh edition. India., 1989.

[18]. George S., "Organic Chemistry" Mosby-Year Book., Chp.14, p. 589-649 (1995).

[19]. Bramhananda, N. R.; Venkataramudu, B.; Ravindranath, L. K.; Aleem, S. A. and Narendra, N. S., *Der Pharma Chemica*, 8,4,101-112., 2016,

[20]. Subbiah, R.; Tanmoy, G.; , Tanushree, S.; Puspita, R.; Benu, P.S.; Jayatri, N.; Avijit, D. and Tapan, K.M., *Der Pharma Chemica*, 8, 4, 446-452., 2016.

[21]. Sahar B. A. and Ammar, A. R., Der Pharma Chemica, 8,4, 63-66., 2016.

[22]. Filali B.Y.; Elmsellem, Y. H.; Kandri, R. H.; Steli, C.A.; Ouzidan, Y.F.; Ouazzani, C.; Sebba, N. K.; Essassi, E. M. and Hammouti, B., *Der Pharma Chemica*, 8,4,159-169., 2016.

[23]. Kiran M. K.; Sagar, A. J.; Pramod, B. P.; Vikas, R. D. and Shitalkumar, S. P., *Der Pharma Chemica*, 8, 4,1-5., 2016.

[24]. Chao J .; Huia, P. X .; Lia, Z ., "Synthesis and Antibacterial Activities of Novel Biphenyltetrazole Derivatives Bearing 1,3,4- Oxadiazole." *Journal of The Chinese Chemical Society*, 52, 539-544 539., 2005.

[25]. Srinivas, K.; Srinivas, U.; Bhanuprakash, K.; Harakishore, K. "Synthesis and antibacterial activity of various substituted s-triazines". *Eur J Med Chem*., 41, 1240-1246., 2006.



[26]. Woese, C.R.; Kandler, O.; Wheelis, *M.L.* "Towards a natural system of organisms: proposal for the domains Archaea, Bacteria, and Eucarya ". ,*Proceedings of the National Academy of Sciences of the United States of America.*, 87, 12, 4576–9., 1990.

[27]. Aatesh, È.; Kocabalkanli, A.; Cesur, N. "Synthesis and antimicrobial activity of some 5-aryl-2-[(N, N-disubstituted thiocarbamoylthio) acylamino]-1,3,4-oxadiazoles", *Farmaco*, 53, 541-544., 1998.

[28]. Montalbetti, C. A.; Falque, V. "Amide bond formation and peptide coupling". *Tetrahedron.*, 61,46, 10827–10852., 2005, doi:10.1016/j.tet. 2005.08.031.