Statistical Studying of Biochemical Compounds on **Microbes**

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ABSTRACT: The statistical studying in our work to complete first work (first paper- in first reference), which includes statistical calculations for studying of microbial behavior on many bacteria and effect of our biochemical compounds against resistant of bacteria, effect of imine derivatives on DNA and wall of bacteria cell through diameter of inhibition (mm) as parameter of activity, effect of types of substitution which linked with ether, nitro, amide, imine, carboxyl, ketone groups in structures of compounds on bacteria.

Keywords: statistical, table, antibacterial, biological.

I.INTRODUCTION

The occurrence of new diseases and concomitant acquisition of microbial resistance to currently medical antibiotics and a drugs, therefore it becomes necessary to discover new pharmaceutical drugs and antimicrobial compounds⁽¹⁻⁷⁾. The microbial effects of anil or imine compounds are quite different and have been identified in several studies⁽⁸⁻¹⁴⁾. Imine are widely known as active group in any chemical compounds which give more bio - effects on their $activity^{(15-31)}$

The nature structure of imine (CH=N) group gave good properties for any chemical compounds which containing its derivatives medical properties as antimicrobial , anti-inflammatory , antihypertensive , antidepressant , antiviral and anticancer activities and other fields $^{(32-40)}$.



Fig(1): Imin group in Compound as A drug

Most of imine compounds have a wide spectrum of medical, industrial application in various fields⁽³¹⁻⁴⁰⁾, it used also as a starting materials in many chemical reactions⁽⁴¹⁻⁵¹⁾.



Fig(2): Imin group in Compound as antimicrobial

EXPERIMENTAL PART :

In the previously $work^{(1)}$ series of imine derivatives were prepared, now in this work we studied effect of our imine compounds on three types of bacteria represented by scanning via three concentrations, then all data of inhibition diameters calculated by statistical



Scheme.1: Prepared Compounds - Imine

Biological Procedure:

Assay of activity for prepared derivatives have been screened for their antibacterial activities by agar through biological procedures $^{16-19}$. The antimicrobial activities were done at three concentrations (1, 3, 5) micro gram) concentrations in (DMSO) solvent with types of bacteria (Staphylococcus aureu), (bacteria K. Pneuomona) and (E- Coli). These bacterial strains were incubated for 24 hr at 37°C.



Picture. (1): Staphylococcus arureus



Picture. (2) : E- Coli



Picture .(3); K. Pneuomona

RESULTS AND DISCUSSION

In past work¹, we formatted imine derivatives while now completed the second part from this work, we will study Activity against three types of microbes.

Bio – Experiments:

The test of the sensitivity of the bacterial isolates were positive for gram, which included work on two types of bacteria to measure the biological activity⁽⁹⁻⁵¹⁾ of certain compounds which bacteria positive for the dye Cram (bacteria *Staphylococcus aureu*) and negative gram (*bacteria K. Pneuomona*), and (*E- Coli*) Tables (1-7) show the diameter of inhibition zone for vehicles chemical measured in mm towards the species bacterial.

Statistical Calculations :

All statistical tables of bacterial inhibition were carried out by using (Statistical package social sciences)program :

Table.1: Statistical Calculations of Biological Activity (Inhibition Zone in ((mm)) in (Concentrations (1, 3, 5 μ gm) for Compound [8]

Bactria	(1) μ gm	(3) µ gm	(5) μ gm
Staphylococcus aureus	7.44±0.32b	12.33±0.32a	14.22±2.15a
K. Pneuomona	5.43±0.45b	14.01±1.12a	11. 13±0.31a
E- Coli	5.19±0.35b	13.11±1.23a	13.44±2.22a

Table.2: Statistical Calculations of Biological Activity (Inhibition Zone in ((mm)) in (Concentrations (1, 3, 5 μ gm) for Compound [9]

Bactria	(1) μ gm	(3) μ gm	(5) μ gm
Staphylococcus aureus	18.14\pm2.15^{°*}b	19.71±1.07^{0*}a	$28.66 \pm 2.789^{\circ*}a$
Bacillus subtilis	$20.87 \pm 1.12^{\circ *}$ b	24.16±2.91 ^{◊*} a	29.91 \pm 2.87 ^{\circ*} a
Streptococcus pyogenes	20.18±2.44 ^{\circ*} b	24.86±1.71 ^{◊*} a	$28.76\pm2.31^{\circ*}a$

Table.3: Statistical Calculations of Biological Activity (Inhibition Zone in ((mm)) in (Concentrations (1, 3, 5 μ gm) for Compound[10]

Bactria	(1) μ gm	(3) µ gm	(5) μ gm
Staphylococcus aureus	11.09±1.16b	11.09±1.51a	12.14±1.07a
K. Pneuomona	10.31±1.18b	10.11±1.03a	13.13±1.10a
E- Coli	7.23±1.22b	10.10±1.31a	11.19±1.17a

Table.4: Statistical Calculations of Biological Activity (Inhibition Zone in ((mm)) in (Concentrations (1, 3, 5 μ gm) for Compound [11]

Bactria	(1) μ gm	(3) μ gm	(5) μ gm
Staphylococcus aureus	24.11±1.19 $^{\circ*}$ b	$28.41\pm3.10^{\circ*}a$	28.97±2.02 ^{◊*} a
Bacillus subtilis	$20.31 \pm 1.11^{\circ *}$ b	$24.14\pm2.73^{\circ*}a$	28.93±1.17 ^{◊*} a
Streptococcus pyogenes	$24.12\pm2.21^{\circ^*}b$	30.54±3.8 ^{◊*} a	30.68±2.31 ^{◊*} a

Table.5: Statistical Calculations of Biological Activity (Inhibition Zone in ((mm)) in (Concentrations (1, 3, 5 μ gm) for Compound [12]

Bactria	(1) μ gm	(3) µ gm	(5) μ gm
Staphylococcus aureus	16.65±2.37^{◊*}b	18.13±2.90^{◊*}a	$22.02\pm2.10^{\circ*}a$
Bacillus subtilis	19.00±2.30 ^{°*} b	20.013±3.11 ^{◊*} a	27.05±1.09 ^{◊*} a
Streptococcus pyogenes	19.44\pm1.82^{°*}b	21.01±2.26 ^{°*} a	27.13±2.89 ^{◊*} a

Table.6: Statistical Calculations of Biological Activity (Inhibition Zone in ((mm)) in (Concentrations (1, 3, 5 μ gm) for Compound [13]

Bactria	(1) μ gm	(3) µ gm	(5) μ gm
Staphylococcus aureus	12.31±1.01b	15.3±1.73a	17±2.1a
K. Pneuomona	14.21±1.03b	13.11±1.83a	16.4±1.17a
E- Coli	14.13±2.31b	12.11±2.01a	16.9±1.34a

Table.7: Statistical Calculations of Biological Activity (Inhibition Zone in ((mm)) in (Concentrations (1, 3, 5 μ gm) for Compound [14]

Bactria	(1) μ gm	(3) µ gm	(5) μ gm
Staphylococcus aureus	8.16±1.10b	11.13±1.00a	12.00±2.00a
K. Pneuomona	9.12±1.09b	14.21±1.18 a	12.41±1.39a
E- Coli	9.14±2.06b	11.14±1.05a	11.69±1.17a

The results of statistical tables appeared data of inhibition for the three types of bacteria on imine derivatives that the Activity of imine compounds which bearing active groups as (nitro group, carboxyl

group) have good activity toward microbes.



Photo. 1: The inhibition of the compounds (9, 10, 11) on Staphylococcus Aureu



Photo. 2: The inhibition of compounds(8, 12, 13, 14) on Staphylococcus Aureu

REFERENCES

- Nagham Mahmood Aljamali , Sajida Hadi Ridha and Noorhan Ali Hamza ., " Synthesis & Studying of Liquid Crystalline 1. Applications of New Oxadiazole Compounds Via (Polarized Optical and Differential Scanning Calorimetry)"., Pak. J. Biotechnol., Vol. 15, 1, 135-144, (2018).
- Alshalchi, S.A.; Alnaib, A.; Kandala, N. (1999).," Antibiotic sensitivities of urinary pathogenic E.coli isolated from pregnant women. 2. AL-Mustansiriya . Journal of science, vol, 10.no,2.
- Alwan, A.S; Abou, S.Z. (1990).,"IRAQI Drug Guide. First edition national, board for the selection of drugs. IRAQ. 3.
- 4. Clark, M.S. (2000)., "The bactericidal activity of gemifloxacin. Journal Medical microbial, Vol. 49, and p: 841-844.
- Clark, W.G.; Brater, D.C.; Johnson, A.R. (1992).,"Medical pharmacology Goths. Introduction to chemotherapy mechanisms of 5. antibacterial. International edition.

- 6. Cruickshank, R.; Duguid, J.P.; Matmion B.P. (1973).,"A Guide to the laboratory diagnosis and control of infections. 12Edition . London Vol:1
- 7. Finkelstein, R; Kassis, E.; Reinhertis, G. (2005)., "Community a quaired urinary tract infection in adult. Journal hospital infection, 38:193-202
- 8. Forrest, A.; Weir.M.; Plaisance, I.K. (1988). ,"Relation between renal function and Disposition oral ciprofloxacin. Journal Antimicrobial agent and chemotherapy, p:1537-1540.
- 9. Gebe, S.S.(2006)., "Asymptomatic bacteriuria. Ethiopia-medical Journal. 36(3):158-192.
- 10. Green W.D; Pearson, N; Eley, A.(1980).," Comparative in vitro-activities of cefotaxime and ceftazoxime. Journal Antimicrobial agents and chemotherapy.p:397-401.
- Guold, J.C. (1973).,"The comparative bacteriology of acute and chronic urinary tract infection. Urinary tract infection . 8th Edition . 11. Nephrology series. England.
- 12. Joseph, L.H. (1998). Chemotherapeutic drugs. Clinical pharmacy and thetapeutics. 3th Edition.
- 13. Kafaf P.A. (2000).," Gentic study on antibiotic resistance of some gram- negative bacteria isolated from urinary tract infection. Thesis, M.Sc. College of science. ALMustansiriya university.
- 14. Kafaf P.A. (2000).," Gentic study on antibiotic resistance of some gram- negative bacteria isolated from urinary tract infection . Thesis, M.Sc. College of science. ALMustansiriya university.
- 15. Knight J.A. (1999).," Encyclopedia of genetics. Second edition. New York.
- 16. Mims H.M. (1995). Antimicrobial agents and chemo therapy .Medical microbiology Third edition.
- 17. Sachadev, K.N. (1989). Examination of urine clinical pathology and bacteriology seventh edition. India .
- 18. Yana, Y; David, M.L. (1998), "Chromosomal B-lactamase expression and resistance to B-lactam antibiotics in proteus vulgaris and morganella morganii. Journal Antimicrobial agents and chemotherapy, p:1385-1391.
- 19. Bramhananda, N. R.; Venkataramudu, B.; Ravindranath, L. K.; Aleem, S. A. and Narendra, N. S., Der Pharma Chemica, 2016, 8,4,101-112.
- Subbiah, R .; Tanmoy, G.; , Tanushree, S .; Puspita, R.; Benu, P.S.; Jayatri, N. ; Avijit, D. and Tapan, K .M., Der Pharma Chemica, 20. 2016,8,4,446-452.
- 21. Sahar B. A. and Ammar, A. R., Der Pharma Chemica, 2016, 8,4, 63-66.
- 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, 2016, 8,4,159-169.
- 23. Kiran M. K.; Sagar, A. J.; Pramod, B. P.; Vikas, R. D. and Shitalkumar, S. P., , Der Pharma Chemica, 2016, 8, 4,1-5.
- 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, 2005, 52, 539-544 539.
- Srinivas, K.; Srinivas, U.; Bhanuprakash, K.; Harakishore, K. "Synthesis and antibacterial activity of various substituted s-triazines". 25. Eur J Med Chem ;2006, 41, 1240-1246.
- Woese, C.R.; Kandler, O.; Wheelis, M.L. "Towards a natural system of organisms: proposal for the domains Archaea, Bacteria, and 26. Eucarya "., Proceedings of the National Academy of Sciences of the United States of America., 1990, 87, 12, 4576–9.
- 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 ,1998, 53, 541-544.
- 28. Montalbetti, C. A.; Falque, V. "Amide bond formation and peptide coupling". Tetrahedron., 2005, 61,46, 10827–10852. doi:10.1016/j.tet. 2005.08.031
- 29. Valeur, E.; Bradley, M. "Amide bond formation: beyond the myth of coupling reagents". Chem. Soc. Rev., 2009, 38,606-631. doi:10.1039/B701677H.
- 30. Nanjunda S.; Swamy, S.; Basppa, P.B.; Prabhuswamy, B.; Doreswamy, B. H. "Crystal Structure of Novel2-butyl-4-chloro-1HImidazolyl-5-Carboxaldehyde". European Journal. of Medicinal Chemistry, 2006, 41, 531-538.3.
- 31. Jin, J. C.; Baoan, S.; Zhuo, C.; Song, Y. "Synthesis, structure, and bioactivity of N0-substituted benzylidene-3,4,5-Trimethoxybenzo hydrazide and - acetyl-2-substituted phenyl-5-(3,4,5-trimethoxyphenyl)-2,3-dihydro-1,3,4-oxadiazole derivatives.", Bioorganic & Medicinal Chemistry Letters, 2006, 16,5036–504.
- 32. Aboraia, S. A.; Rahman, A. M.; Mahouz, M. N.; "Novel 5-(2 hydroxyphenyl)-3-substituted-2,3-dihydro-1,3,4-oxadiazole-2-thione derivatives: Promising anticancer agents.", Bioorganic & Medicinal Chemistry ,2006, 14, 1236-1246.
- 33. Mieaad Mohammed, Nagham Mahmood Aljamali, Sabreen Ali Abdalrahman, Wassan Ala Shubber., "Formation of Oxadiazole Derivatives Ligands from Condensation and Imination Reaction with References To Spectral Investigation, Thermal and Microbial Assay" ., Biochem. Cell. Arch. , 2018 , 18, 1, pp. 847-853.
- 34. Nagham Mahmood Aljamali. "Synthesis and Biological Study of Hetero (Atoms and Cycles) Compounds", Der Pharma Chemica, 2016, 8, 6, 40-48.
- 35. Nagham Mahmood Aljamali.; Intisar, O. "Synthesis of Sulfur Heterocyclic Compounds and Study of Expected Biological Activity" ,Research J. Pharm. and Tech., 2015, 8,9 ,1225-1242 , DOI: 10.5958/0974-360X.2015.00224.3.
- Nagham Mahmood Aljamali .; Saher Mahmood J .; Zainab M J .; Seena K. "Microbial Studying of (Thiazole, Oxadiazole, 36. Thiadiazole)-Derivatives on Mouth and Teeth Bacteria", International Journal of Medical Research and Pharmaceutical Sciences, 2016, 3, 8, 30-39, DOI:10.5281/zenodo.61357.
- 37. Rappé, M. S.; Giovannoni, S. J. "The uncultured microbial majority". Annual Review of Microbiology., 2003, 57, 369-94.
- 38. DeLong, E.F.; Pace, N.R. "Environmental diversity of bacteria and archaea". Syst Biol., 2001, 50, 4, 470-8.
- 39. Rosenberg, S.M.; Slack, A. "Antibiotic-induced lateral transfer of antibiotic resistance". Trends Microbiol., 2004, 12, 9, 401-4.
- 40. Gitai, Z. "The new bacterial cell biology: moving parts and sub cellular architecture". Cell, 2005, 120, 5, 577-86.
- 41. Vanita S .; Supriya, M. "Development of furfuraldehyde formazans as potential antitubercular agents " ., Der Pharma Chemica, 2016,8,18, 144-148.
- 42. Park, S.; Park, B.; Yun, S.; Kang, H. and Yun, S. "Antimicrobial activities of honey bee venom against pathogens isolated from clinical bovine mastitis in Korea", Planta Med., 2013, 79, , PL16.
- 43. Habibe, T .; Mehmet, L. A . "Electrochemical properties of 1-(o-, m-, pnitrophenyl)-3-(m-nitrophenyl)-5-phenylformazans and their nickel complexes" ., Turk J Chem , 2010, 34 , 465 - 479.

International Journal Of Advanced Research In Medical & Pharmaceutical Sciences(IJARMPS) Volume.3, Issue.8, August. 2018

- **44.** Nagham Mahmood Aljamali .; Saher Mahmood J .; Zainab Mahmood J .; Intisar O. "Inhibition Activity of (Azo Acetyl acetone) on Bacteria of Mouth", Research J. Pharm. and Tech. ,2017, 10, 6,1683-1686 .,DOI: 10.5958/0974-360X.2017.00297.9
- **45.** Angela, K.; Lucica, V.; Nicoleta, C. " Synthesis and structural studies of complexes of Cu, Co, Ni and Zn with isonicotinic acid hydrazide and isonicotinic acid (1-naphthylmethylene)hydrazide", J. Serb. Chem. Soc, 2010, 75, 2, 229-242.
- **46.** Wright, G.D. " The antibiotic resistome: the nexus of chemical and genetic diversity"., Nature Reviews Microbiology, 2007, 5, 175-186.
- **47.** Nagham Mahmood Aljamali . "Synthesis and Chemical Identification of Macro Compounds of (Thiazol and Imidazol)".,Research J. Pharm. and Tech, 2015, 8,1, 78-84., DOI : 10.5958/0974-360X.2015.00016.5 .
- **48.** Hegazi, A.G.; Abdou, A.M. and Abd, A. F. "Evaluation of the antibacterial activity of bee venom from different sources", World Applied Sciences Journal, 2014, 30, 3, 266-270.
- **49.** Intisar Obaid Alfatlawi , Nuha Salman S, Zainab Mahmood J , Nagham Mahmood Aljamali , " Synthesis of New Organic Compounds Via Three Components Reaction with Studying of (Identification ,Thermal Behavior, Bioactivity on Bacteria of Teeth) "., Journal of Global Pharma Technology. 2017; 11, 9, 157-164.
- Eman H. S., Nagham Mahmood Aljamali ., "New Azo-Thiadiazole Ligands (Preparation, Spectral, Thermal, Biochemical, Physical properties) Studying " ., Journal of Global Pharma Technology. 2017; 11, 9, 165.
 Mieaad Mohammed, Nagham Mahmood Aljamali, Nadheema Abed Abbas ., " Preparation, Spectral Investigation, Thermal
- **51.** Mieaad Mohammed , Nagham Mahmood Aljamali , Nadheema Abed Abbas ., " Preparation, Spectral Investigation, Thermal Analysis, Biochemical Studying of New (Oxadiazole Five Membered Ring)-Ligands".,Journal of Global Pharmacy Technology, 2018; 10, 1, 20-29.