

Graphical abstract





Heterocyclic Letters 14: iss.-2 (2024), 249-254 Paper-3 A one pot method for the synthesis of 1-(2-trifluoromethanphenyl)-5-phenyl-1h-tetrazole (1, 5-disubstituted tetrazole) catalyzed by zirconium oxychloride and their antibiotic activity *G.Nageswara Rao, M.Rajeshwari Department of Chemistry, Telangana University, Nizamabad, Telangana State-503322, India. 1-(2-trifluoromethan phenyl)-5-phenyl-1H-tetrazole from amide a mixture of 2-Trifluoromethyl aniline, sodium azide and zirconium oxychloride in dry acetonitrile with Reflux 43hr. 43 hr CH₃CN NaN₃ + ZrOCL Reflux R=CF₃ scheme 1





Heterocyclic Letters 14: iss.-2 (2024), 259-265 Paper-5 Studies in designing of substituted 4h-pyrano [3,2-h]quinoline-3-carbonitrile Derivatives as potential neuraminidase inhibitors of swine flu in silico approach. Prasanna B. Ranade¹*, Dinesh N. Navale¹, Pramod P. Gaikwad¹, Usama A. Anware¹, Santosh W. Zote², Dnyaneshwar K. Kulal³, Swapnil J. Wagh⁴, Vaijayanti Ghase⁵ ¹Department of Chemistry, Vivekanand Education Society's College of Arts, Science and Commerce, (Autonomous), Chembur, Mumbai 400 071 INDIA. ²Department of Chemistry, PTVA's Sathaye College (Autonomous), Dixit Road, Vile Parle (East), Mumbai-400 057, Maharashtra, INDIA. ³Department of Chemistry, Ramnarain Ruia Autonomous College, L. N. Road, Matunga, Mumbai-400 019, Maharashtra, INDIA. ⁴Department of Chemistry, R.S.S. Prasarak Mandal's Nanasaheb Yashvantrao Narayanrao Chavan, Arts, Science & Commerce College Chalisgaon, Jalgaon-424101Maharashtra, INDIA. ⁵Department of Chemistry, S.K. Somaiya College Vidyavihar, Ghatkopar Mumbai, INDIA Corresponding author: prasannabranade@gmail.com N294 S246 R292 E276 E PDB ID 3TI6 Docking R371 Y406 E227 E119 R156 L184

 Paper-6
 Heterocyclic Letters 14: iss.-2 (2024), 267-272

 Indium-mediated reduction of aromatic nitro groups in β-lactams to oxazines

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An indium-induced reduction of nitro-substituted β -lactams has been used for facile synthesis of oxazines in aqueous ethanol using ultrasound at 40°C through molecular rearrangement.



Paper-7 [Heterocyclic Letters 14: iss2 (2024), 273-286 Physicochemical parameters involved in the interaction of some phenanthroline derivatives with janus kinase-3 protein using a theoretical model Figueroa-Valverde Lauro ¹⁹ , Rosas-Nexticapa Marcela ² , Alvarez-Ramirez Magdalena ² , López-Ramos Maria ¹ , Mijangos Sánchez Juliette ¹ . ¹ Laboratory of Pharmaco-Chemistry, Faculty of Chemical Biological Sciences, University Autonomous of Campeche, Av. Agusti Melgar s/n, Col Buenavista C.P. 24039 Campeche, Camp., México.			
Physicochemical parameters involved in the interaction of some phenanthroline derivatives with janus kinase-3 protein using a theoretical model Figueroa-Valverde Lauro ^{1*} , Rosas-Nexticapa Marcela ² , Alvarez-Ramirez Magdalena ² , López-Ramos Maria ¹ , Mijangos Sánchez Juliette ¹ . ¹ Laboratory of Pharmaco-Chemistry, Faculty of Chemical Biological Sciences, University Autonomous of Campeche, Av. Agusti Melgar s/n, Col Buenavista C.P. 24039 Campeche, Camp., México.	Paper-7	Heterocyclic Letters 14: iss2 (2024), 273-286	
Figueroa-Valverde Lauro ^{1*} , Rosas-Nexticapa Marcela ² , Alvarez-Ramirez Magdalena ² , López-Ramos Maria ¹ , Mijangos Sánchez Juliette ¹ . ¹ Laboratory of Pharmaco-Chemistry, Faculty of Chemical Biological Sciences, University Autonomous of Campeche, Av. Agustí. Melgar s/n, Col Buenavista C.P. 24039 Campeche, Camp., México.	Physicochemical parameters involved in the interaction of some phenanthroline derivatives with janus kinase-3 protein using a theoretical model		
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$H_{AR} H_{BA}$	¹ Laboratory of Pharmaco-Chemistry, Faculty of Chemical Biological Sciences, University Autonomous of Campeche, Av. Agustín Melgar s/n, Col Buenavista C.P. 24039 Campeche, Camp., México.		
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Paper-8

Heterocyclic Letters 14: iss.-2 (2024), 287-291

Dipole Moment Studies on Anticancer Polyaromatic Compounds

Aparna Das¹, Ram Naresh Yadav,² and Bimal Krishna Banik¹*

¹Department of Mathematics and Natural Sciences, College of Sciences and Human Studies, Prince Mohammad Bin Fahd University, Al Khobar, Kingdom of Saudi Arabia;²Veer Bahadur Singh Purvanchal University, Jaunpur-222003 (U.P), India; *Email: bimalbanik10@gmail.com; bbanik@pmu.edu.sa* Dipole moment values of a few anticancer polyaromatic molecules is investigated.





Paper-9 Heterocyclic Letters 14: iss.-2 (2024), 293-299 Novel Synthesis of Heterocyclic 1,3,4 Thiadiazole based bioactive Metal complexes Ajay M Patil^{1*}, Chandrashekhar G. Devkate², Uddhav Chaudhar³, Shyam Takle⁴ ^{1*}Department of Chemistry, Pratishthan College Paithan, Chh.Sambhajinagar-431107, India ²Dept. of Chemistry, IndrarajArts, Com. and Sci. College, Sillod, Chh. Sambhajinagar-431112, India ^{3*}Department of Chemistry, Kalikadevi Art's, Science & Commerce College, Shirur(Ka.), Beed, India ⁴Department of Chemistry, S.D.College Soegaon, Chh.Sambhajinagar-431120, India E-mail authors: patilam4@gmail.com Novel Heterocyclic 1,3,4 ThiadiazoleFe (III) Complexes were prepared. Metal Complexes were prepared from ligand 4-bromo-2-(((5-mercapto-1,3,4-thiadiazol-2-yl)imino)methyl)-6-methoxyphenol with Fe(NO₃)₃.9H₂O (Iron Nitrate). Structures of complexes were confirmed based on different spectroscopic techniques like elemental analysis, FT-IR, UV-Vis, magnetic and molar conductivity measurements. All complexes were non-electrolytes and had octahedral geometry. Investigation of bioactivity for complexes and ligands exposed their moderate antibacterial activities tested in vitro against bacterial StrainsS.aureusandB. subtilis and fungal strains of F. OxysporumandA. Niger using Kirby-Bauer disc diffusion method. Fe(NO₃)₃.9H₂O насо о́сн₃ осн в

Paper-10 Heterocyclic Letters 14: iss.-2 (2024), 301-308
Molecular Iodine Catalysed Synthesis Of Tetrahydroazepines Frameworks *Via* Silyl Aza-Prins Cyclization

N. Prudhvi Raju^{a*}, R. L. Satyanarayana^b, D. Ravikumar^c, K. Pavan Krishna^d

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Based on the powerful Aza-Prins cyclization in combination with the Peterson-type elimination reaction, a C–N, C– C bond and an endocyclic double bond are formed. Under mild reaction conditions, using Molecular Iodine as sustainable catalysts, tetrahydroazepines with different degrees of substitution are obtained directly and efficiently





Paper-11 Heterocyclic Letters 14: iss.-2 (2024), 309-317 Maleimide Based Donor-Acceptor Fluorophore: Microwave Assisted Green Synthesis, Photophysical and DFT Study Kirankumar S. Gosavi Department of Chemistry, KVPS's Kisan Arts, Commerce and Science College, Parola, Dist: Jalgaon, Maharashtra, India. Pin: 425111 Email-id: kirangosavi08@gmail.com ORCID ID: 0000-0003-2143-06 This paper deals with green synthesis of 3-aryl 4-methoxy maleimide derivatives, using microwave irradiation method to attribute its photophysical properties. Donor R insity (a.u.) Me 500 550 Acceptor Wavelength (nm)

Paper-12Heterocyclic Letters 14: iss.-2 (2024), 319-327Synthesis and characterization of chromium (iii) complexes of
p-vanillin semicarbazone and thiosemicarbazone

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Chromium (III) complexes of p-vanillin semicarbazone (L^1) and thiosemicarbazone(L^2) have been synthesized. These complexes were characterized by elemental analysis, molar conductance, magnetic moment, IR, electronic and epr spectral studies. Complexes were found to have Cr (L_1 and L_2)₂ X₃ Composition. Molar conductance indicates that chloro complexes are 1:1 whereas nitrato complexes are 1:2 electrolyte in nature. Both the ligands act as bidentate. On the basis of spectral studies an octahedral geometry has been assigned for all the complexes.



Paper-13

Heterocyclic Letters 14: iss.-2 (2024), 329-352

Cu(II) complexes of derivatives of (thiophen-2-yl)methanamine: Design, Synthesis, Spectral characterization, DNA binding investigations and antimicrobial activity.

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The synthesis and characterization of six new Schiff base ligands, including the imine ligands TMMCFP, TMMFMOP, TMMFP, TMMDMOP, TMMBD, and TMMDEP, as well as their complexes with 3D series copper metal, included elemental analyses, ¹H, ¹³C NMR, IR, ESI mass, UV-Visible, ESR, magnetic moment, and TGA investigations. A square planar shape is ascribed to the Cu (II) compounds based on the findings from the studies to surround the metal ion. Using UV-visible electronic, fluorescence, and viscosity studies, the interaction of synthesized metal complexes with calf thymus Deoxyribonucleic acid (CT-DNA) was investigated.



 Paper-14
 Heterocyclic Letters 14: iss.-2 (2024), 353-361

 Synthesis, spectral and thermal studies of Schiff bases derived from N-(4-((2-hydroxybenzylidene) amino) phenyl)

 pyridine-2-sulfonamide

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Paper-15 Heterocyclic Letters 14: iss.-2 (2024), 363-372 Study of Novel Metal Complexes of Thiosemicarbazones Sandip D. Patel, Mohit R. Joshi, Dr. Harshad Kumar P. Patel Sankalchand Patel University, Visnagar-384315 sandippatel77790@gmail.com The novel metal complexes of heterocyclic ligand, i.e., 2-(1-(5-chloro -1H-benzo[d] [1,2,3] triazol-1-yl)propan-2-ylidene) hydrazine carbothioamide (CBTTS) were synthesized from reaction between novel ligand with selected Transition Metal salts. The novel metal complexes characterized for their elemental contents, Spectral studies, metal: ligand ratio and magnetic properties of novel ligand and their metal complexes, which are confirmed the predicted structure and best antibacterial and antifungal activities of ligand CBTTS and its all metal complexes. H=N + \(\mathcal{F}\) + \(\mathcal\) + \(\mathcal{F}\) + \(\mathcal{F}\) + \(\ma

 Paper-16
 Heterocyclic Letters 14: iss.-2 (2024), 373-383

 One-pot synthesis of 2-substituted benzoxazole/benzimidazole/Benzothiazoles
 using (diacetoxyiodo)benzene (DIB)as an efficient catalyst

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Using DIB as a catalyst in CH_3CN , we reported the synthesis of 2-substituted benzoxazole/benzimidazole/benzothiazoles. This approach greatly increases its utility of as a special and potent tool in chemical synthesis. This technique offers a novel approach to the synthesis of diverse benzoxazole, benzimidazole, and benzothiazole derivatives using various aldehydes and o-substituted amino aromatics. This technique gives good to exceptional yields of products and is relatively simple to build up. Following the reaction, the products were examined by mass spectroscopy, infrared spectroscopy, and NMR.







Paper-17 Heterocyclic Letters 14: iss.-2 (2024), 385-406 Synthesis of Warfarin

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 Paper-18
 Heterocyclic Letters 14: iss.-2 (2024), 407-422

 Biologically potent novel heterocyclic azomethines and their co (ii) metal complexes: green synthesis, characterization and molecular docking studies with sars-cov-2 main protease 6lu7

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Paper-19

Heterocyclic Letters 14: iss.-2 (2024), 423-430

Green synthesis and characterization of copper nanoparticles using allamanda blanchetii flower extract

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The Allamanda Blanchetii flower extract is used for the biosynthesis of Coppernanoparticles and the successful formation of the resultant product was confirmed through several physicochemical techniques. The crystalline structure and crystallite size were investigated through an X-ray diffractometer (XRD). The antioxidant test is also carried out against 1,1-diphenyl -2- picrylhydrazyl freeradicals and the antioxidant potential of CuNPs were found to be higher than ascorbic acid



Paper-20 Heterocyclic Letters 14: iss.-2 (2024), 431-442 . CF3SO3H.SiO2 Catalyzed, Solvent Free, 'N' and 'O' Formylation of Amines, Anilines Alcohols and Phenols'

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Ethyl Methanoate has been successfully employed as an 'N' and 'O' formylating agent in a straightforward way in presence of catalytic amount of $CF_3SO_3H.SiO_2$. This methodology is solvent free and economical for the synthesis of formamides and formats. The methodology tolerates different aromatic and aliphatic amines, anilines and phenols offering well to excellent yield of the intended products within short reaction time. Late stage synthesis of formates and formates also feasible using this methodology.





Paper-21

Heterocyclic Letters 14: iss.-2 (2024), 443-448

Synthesis, characterization, & antimicrobial screening of N-thiadiazolyl thiazolidinone derivatives

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A new series of 2-(5-(substitutedbenzyl)-1,3,4-thiadiazol-2-yl)-(substitutedphenyl)- thiazolidin-4-one (5) have been synthesized by the reaction of Schiff base of 2-amino-5-(substituted)-benzyl-1,3,4-thiadiazole (3) with α -mercaptoalkanoic acid (4) in glacial acetic acid. The structures of the compounds have been confirmed by IR, NMR and Mass Spectroscopy. Representative compounds were screened for their anti-microbial activity against gram-negative bacteria, E coli and P.aeruginosa and grampositive bacteria, S aureus, and C diphtheriae using disc diffusion method. Some of these compounds have been found to exhibit excellent antibacterial activity.



PERSPECTIVES

Perspectine No.1 Microwave in Research-More Miracles Heterocyclic Letters 14: iss.-2 (2024), 449-456

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This perspective describes the future possible trends in microwave research. As an example, future trends in organic and inorganic chemistry, material science, biochemistry, flow chemistry, communication, earth science, the food industry, and medicinal chemistry are considered.



Perspectine No.2

Heterocyclic Letters 14: iss.-2 (2024), 457-467

Expeditious Synthesis of Oxygen and Sulfur Heterocycles by Microwave

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The oxygen- and sulfur-containing heterocyclic compounds are important compounds due to their diverse medicinal activities. Microwave-induced reactions were performed for the preparation of O-heterocycles and S-heterocycles as environmentally benign, fast, high yielding and economical methods.

REVIEWS





 Review No.2
 Heterocyclic Letters 14: iss.-2 (2024), 479-492

 2-[Bis(methylthio)methylene]malononitrile: A Versatile Reagent in the Diversity-Oriented Synthesis of Complex

 Heterocyclic Scaffolds

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2-[Bis(methylthio)methylene]malononitrile extensively applied as a synthon in the diversity-oriented synthesis of diverse heterocyclic compounds. These include mono/polycyclic-heterocyclic compounds, fused heterocyclic compounds, bicyclic bridged nitrogen, sulphur, oxygen-containing heterocycles, substituted heterocycles, and pyrimido-heterocycles. These compounds are synthesized via various reactions, such as cyclo-condensation, cyclo-addition, cascade reactions, and multi-component reactions. The compound's versatility and high reactivity as a multi-functional reagent stem from its easily available structure, featuring electron-withdrawing two cyano functional groups, two methylthio as the best leaving groups, and an α , β -unsaturated alkene part. Its unparalleled potential for crafting complex and functional heterocyclic frameworks has prompted us to compile a comprehensive review on the synthetic utility of 2-[Bis(methylthio)methylene]malononitrile, highlighting its role as a potent reagent in the synthesis of heterocyclic scaffolds.





Review No.3

Heterocyclic Letters 14: iss.-2 (2024), 493-506

Piperidine scaffold drugs and their medicinal applications

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Piperidine containing compounds represents one of the most important synthetic medicinal block for drug synthesis as, it has many biological applications. This review article discusses the various naturally occurring and synthetic piperidine derivatives with applications.