



## Graphical Abstract

Paper-1	Heterocyclic Letters 12: iss.-1 (2022), 13-18
<b>Preparation of a resveratrol-ascorbic acid mem conjugate</b>	
<b>Rasidul Amin* and David A. Hunt**</b>	
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** <i>Department of Chemistry, The College of New Jersey, 2000 Pennington Road Ewing, NJ 08628</i>	
il: <a href="mailto:hunt@tcnj.edu">hunt@tcnj.edu</a>	

Paper-2	Heterocyclic Letters 12: iss.-1 (2022), 19-23
<b>Banik's Cycloaddition Reaction Towards Beta Lactams: Mechanistic Insights</b>	
<b>Bimal Krishna Banik</b>	
<i>Department of Mathematics and Natural Sciences, College of Sciences and Human Studies, Deanship of Research, Prince Mohammad Bin Fahd University, Al Khobar 31952, KSA; Email: <a href="mailto:bimalbanik10@gmail.com">bimalbanik10@gmail.com</a>; <a href="mailto:bbanik@pmu.edu.sa">bbanik@pmu.edu.sa</a></i>	
Banik's cycloaddition describes the stereospecific synthesis of $\beta$ -lactams starting from polyaromatic imines and acid chlorides in the presence of a tertiary base. The N-polyaromatic substituent has a crucial role that controls the synthesis of thermodynamically stable <i>trans</i> -N-azetidin-2-one. A probable mechanism of this process is advanced based upon the available literature	

**In vitro antioxidant activity of different polarity extracts and metabolites isolated from *Moltkia ciliata* growing in algeria**

Soumaia CHIHI <sup>a,b</sup>, Oumelkheir RAHIM <sup>a,□</sup>, Rabab ZATOUT <sup>c</sup> and Ali DOUADI <sup>a</sup>

<sup>a</sup> Pollution & Waste Treatment Laboratory, Department of Chemistry, Faculty of Mathematics and Matter sciences, Kasdi Merbah University, Ouargla 30000, Algeria.

<sup>b</sup> VTRS Laboratory, El-Oued University, B.P. 789, El-Oued 39000, Algeria <sup>c</sup> Department of pharmacy, Faculty of medicine, Benyoucef Benkhedda Algeirs IUniversity, Algeria. \* E-mail Corresponding author: [rahoumelkheir@gmail.com](mailto:rahoumelkheir@gmail.com)

The plant *Moltkia ciliata* belong to a family *Boraginaceae*, although this plant has been traditionally used for the treatment of various ailments still no systematic pharmacognostical, phytochemical and pharmacological work has ever been carried out on this potential plant.

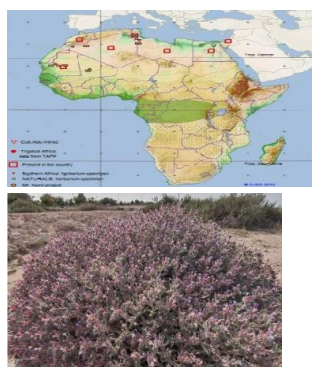


Fig.1: Geographical distribution of the plant in Africa in 2007 and image showing the outside of the plant *Moltkia ciliata*

Evaluation of antioxidant properties using electrochemical and spectrophotometrical assays and determination of *total bioactive compounds* content of the ethyl acetate (EtOAc) and *n*-butanol (*n*-BuOH) extracts of *Moltkia ciliata*. Chromatographic and HPLC analysis of extract shows many compounds with different quantities. The study concludes this plant is a rich source of phenols and flavonoids, and also showed good in-vitro antioxidant activity by all methods.

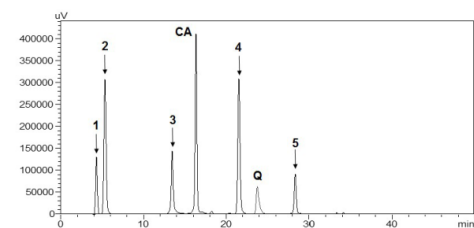


Fig. 2: Chromatograms HPLC

The results showed that EtOAc extract which had the highest level of *bioactive compounds* content, exhibited the most potent antioxidant capacity in each assays compared to the *n*-BuOH extract.

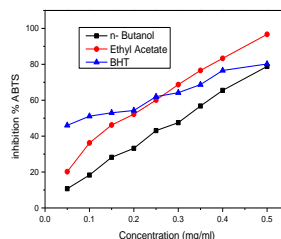


Fig. 4: ABTS radical scavenging activities of extracts of *M.Ciliata*

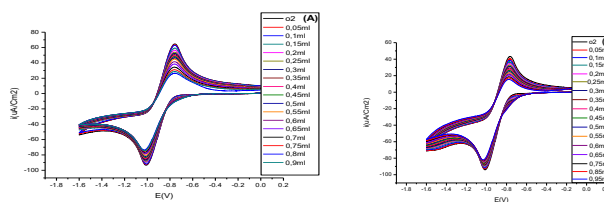


Fig. 5: Cyclic voltammograms of  $O_2$  in presence of (A) AcOEt extract, (B) *n*-BuOH extract at different concentrations in DMF +0.1M TNBHFP on GC as working electrode vs. Hg/Hg<sub>2</sub>Cl as reference at 28°C with scan rate of 100 mV/s

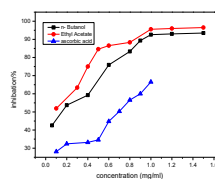


Fig. 6: The DPPH radical scavenging activity of *M. Ciliata* extracts and the ascorbic acid.

Thus, the plant *M. ciliata* can be explored as a potential source of natural antioxidant.

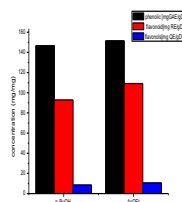


Fig. 3: Total phenolic content, flavonoid and flavonols of extracts from *M. ciliata*



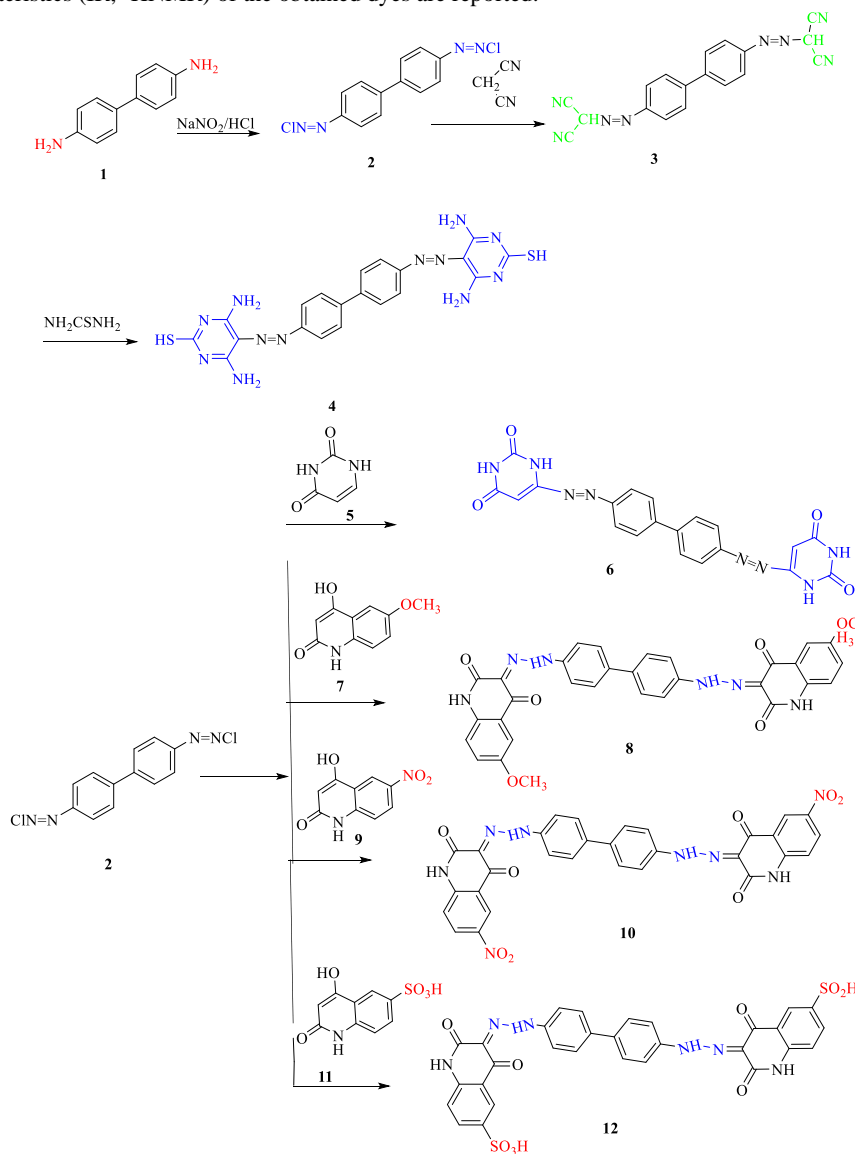
### Synthesis and molecular docking of some new azo dyes derived from benzidine and study some of their antimicrobial potential

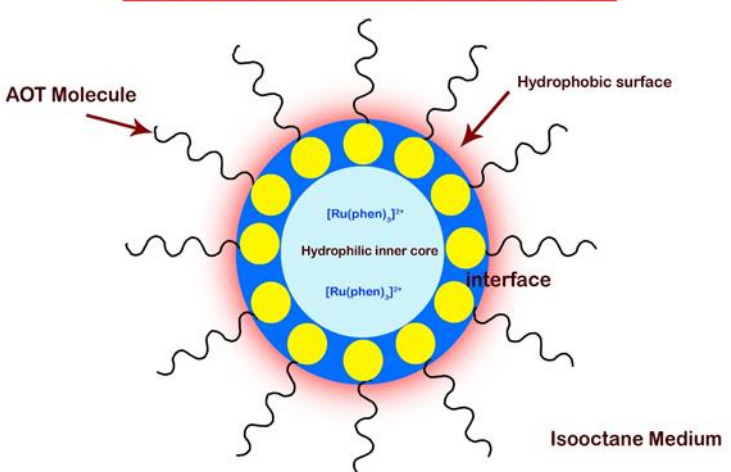
A. A. Ghoneim

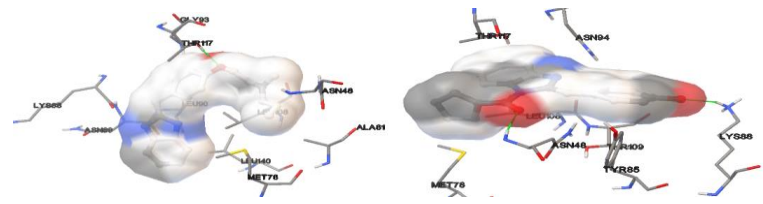
Chemistry Department, College of Science, Jouf University, P.O. Box: 2014, Sakaka, Saudi Arabia.

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This research includes preparing some new heterocyclic compounds derived from benzidine used as an azo dye such as thiopyrimidine, from the reaction between 2, 2'-((*1E, 1'E*)-[1, 1'-biphenyl]-4, 4'-diylbis (diazene-2, 1-diyl)) dimalononitrile **3** and thiourea in absolute ethanol to 5, 5'-((*1E, 1'E*)-[1, 1'-biphenyl]-4, 4'-diylbis (diazene-2, 1-diyl)) bis (4, 6-diaminopyrimidine-2(*5H*)-thione **4** and another way four compounds were synthesized from diazonium salt of benzidine with 4-hydroxy-6-methoxyquinolin-2(*1H*)-one (**7**), 4-hydroxy-6-nitroquinolin-2(*1H*)-one (**9**) and 1, 2-dihydro-4-hydroxy-2-oxoquinoline-6-sulfonic acid (**11**) to give (*3E*)-3, 3'-([1, 1'-biphenyl]-4, 4'-diylbis (hydrazin-2-yl-1-ylidene)) bis (6-methoxyquinoline-2,4 (*1H,3H*)-dione) **8**, (*3E*)-3,3'-([1, 1'-biphenyl]-4,4'-diylbis (hydrazin-2-yl-1-ylidene)) bis (6-nitroquinoline-2,4 (*1H,3H*)-dione) **10** and (*3E*)-3,3'-([1, 1'-biphenyl]-4,4'-diylbis (hydrazin-2-yl-1-ylidene)) bis (2,4-dioxo-1,2,3,4-tetrahydroquinoline-6-sulfonic acid) **12** respectively. The spectral characteristics (IR, <sup>1</sup>HNMR) of the obtained dyes are reported.



Paper-5	Heterocyclic Letters 12: iss.-1 (2022), 51-59
<p><b>Photoemission characteristics of ruthenium(ii) phenanthroline complexes in aot reverse micelle</b></p> <p><b>Helen Ratna Monica Jeyarajan*</b></p> <p><i>Post Graduate &amp; Research Department of Chemistry, The American College, Madurai - 625 002, Tamilnadu, India.</i></p> <p>*Email: <a href="mailto:helenmonica@americancollege.edu.in">helenmonica@americancollege.edu.in</a></p> <p style="text-align: center;"><u>[Ru(phen)<sub>3</sub>]<sup>2+</sup> in AOT/isooctane/water Reverse Micelle</u></p> 	

Paper-6	Heterocyclic Letters 12: iss.-1 (2022), 61-67
<p><b>Synthesis and Antimicrobial activity of Substituted 4-[(1<i>H</i>-benzo[<i>D</i>]imidazol-2-yl)methyl]phenol derivatives</b></p> <p><b>Srinivasulu Chittimalla, Jalapathi Pochampally*, Sreedhar Maroju, Ganapathi Velupula, Veera Bh dram, Vishnu Thumma and Umapathi Nalla</b></p> <p><i>Department of Chemistry Rayalaseema University, Kurnool, India, 500004.</i>              Email: <a href="mailto:srinivas6meena@gmail.com">srinivas6meena@gmail.com</a></p> <p>A new set of <i>N</i>-substituted benzimidazole analogues were prepared by the reaction of respective benzimidazoles with different acids, acid chlorides and alky halides under appropriate conditions. The synthesised derivative were screened for antimicrobial activity and studied their molecular docking</p> 	

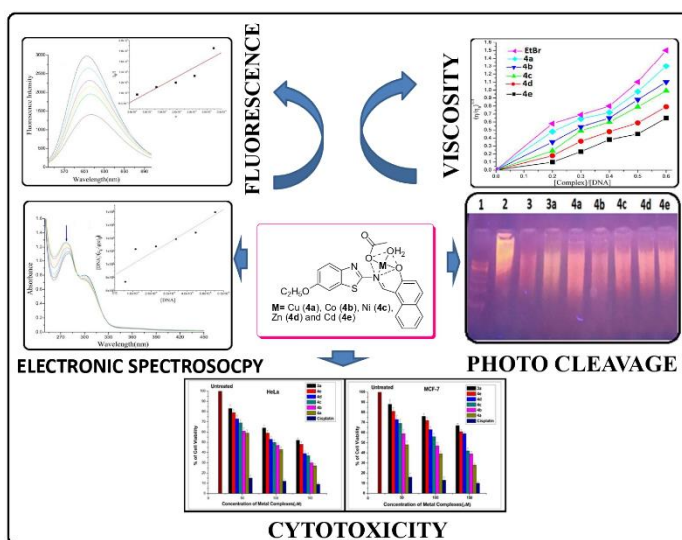
**DNA binding, Photo cleavage, Antibacterial and Cytotoxicity activities of Mono nuclear metal complexes of an N, O donor Benzothiazole derivative Schiff base ligand**

**N. Nageswara Rao, K. Gopichand, E. Kishan, R. Nagaraju, Abdul Majeed Ganai and P. Venkateswar Rao\***

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Imine base ligand and its mono nuclear metal complexes have been synthesized and were characterized by various spectroscopic and analytical techniques. The DNA binding, Photo cleavage studies have been studied. The synthesized compounds were screened for two gram negative bacterial species and three gram positive bacterial species. The In-vitro cytotoxicity of the synthesized compounds was also evaluated by MMT assay

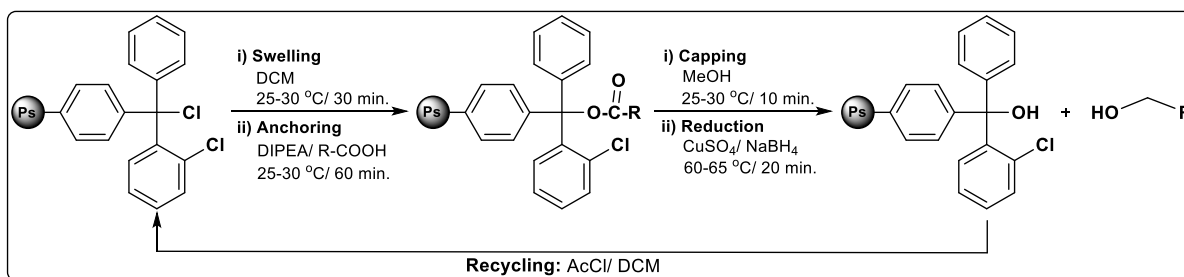


**Reduction of aliphatic, aromatic and heteroaromatic carboxylic acid derivatives to alcohol promoted by trityl resin under presence of copper sulphate and sodium borohydride catalytic system**

**Anirudhdha G. Kalola <sup>a</sup>, Pratibha Prasad <sup>a</sup> and Manish P. Patel <sup>a\*</sup>**

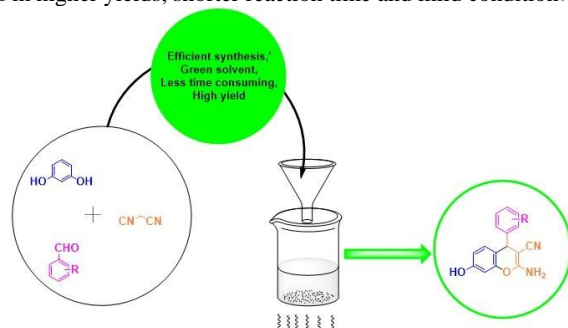
<sup>a</sup> Department of Chemistry, Sardar Patel University, Vallabh Vidyanagar-388120, Gujarat, India

Corresponding Author. Tel.:02692-226856; e-mail: [patelmanish1069@yahoo.com](mailto:patelmanish1069@yahoo.com)

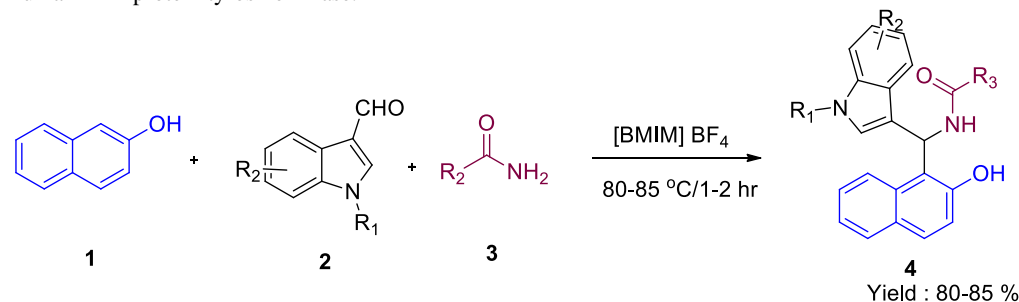



**Ultrasonic assisted synthesis and optimization of 2-amino-4h-chromene's derivatives**
**Sarika Patel, Shweta Patel, Hasit Vaghani\*, Jasmin Kumbhani\*, Ravibhai Bhola**
*Faculty of Science, Department of Chemistry, Ganpat University, Kherva, Mehsana-384012, Gujarat, India*
*\*Correspondence: [jhk01@ganpatuniversity.ac.in](mailto:jhk01@ganpatuniversity.ac.in), [hvv01@ganpatuniversity.ac.in](mailto:hvv01@ganpatuniversity.ac.in)*

A one pot three component reaction of 2-amino-4-aryl-4H-chromene and its derivatives were synthesized under ultrasonic irradiation using morpholine as a catalyst in aqueous condition. The synthesized titled derivatives were confirmed by IR, <sup>1</sup>H NMR, <sup>13</sup>C NMR, mass spectra and elemental analysis. When compared to traditional method, ultrasonic irradiation is a modern technology for improving prepared in higher yields, shorter reaction time and mild condition.


**An efficient ionic liquid medium for the synthesis of new naphthol-indole hybrids and their anti cancer evaluation**
**Kurapati Ravi<sup>a</sup>, Bala Narsimha Dhoddi<sup>a</sup> and Jalapathi Pochampally<sup>a\*</sup>**
*<sup>a</sup> Department of Chemistry, Osmania University, Hyderabad, Telangana, India - 500 007.*
*Corresponding author E-mail ID: [pochampalli.ou.chemi@gmail.com](mailto:pochampalli.ou.chemi@gmail.com)*

A one-pot three-component condensation of 2-naphthol **1** with indol aldehydes **2** and amides **3** in the presence of ionic liquid medium [BMIM] BF<sub>4</sub> has been described as an environmentally friendly and efficient process for the synthesis of N-((2-hydroxynaphthalen-1-yl)(1-methyl-1H-indol-3-yl)methyl)acetamide derivatives **4**. This innovative process has various advantages, including high yields, quick reaction times, operational simplicity, and no time-consuming set-up. Further, the synthesized compounds were examined for their anticancer potential towards PC-3 and SKOV-3 cells. Among the screened, compound 4b and 4f are found to be the promising compounds with IC<sub>50</sub> values ranges from 7.9 to 9.1 μM against PC-3 and SKOV-3 cells. In additional, the docking studies revealed that compound 4b and 4f showed good binding affinities in relation to human RET protein tyrosine kinase.





Synthesis of Novel Products by Using Aniline and Ethyl Benzoate

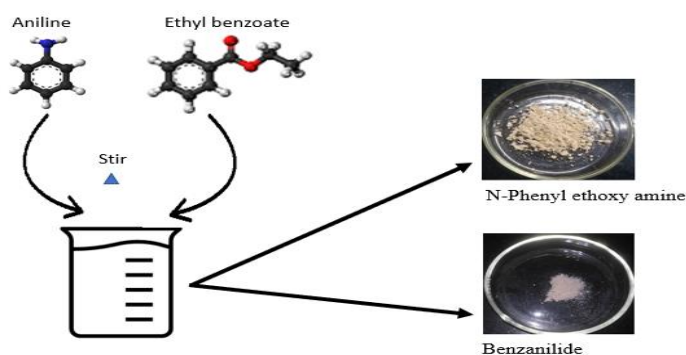
Megha Shankar Gharge<sup>a,\*</sup>, V. M. Kamble<sup>b</sup>

<sup>a</sup>Research Scholar, <sup>b</sup>Associate Professor

<sup>a</sup>Department of Chemistry, <sup>a</sup>The Institute of Science, Mumbai University, 15, Madam Cama Road, Mumbai-32.

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Simple method of Synthesis of novel products by using aniline and ethyl benzoate. This preparation was performed by using aniline and ethyl benzoate without any catalyst.

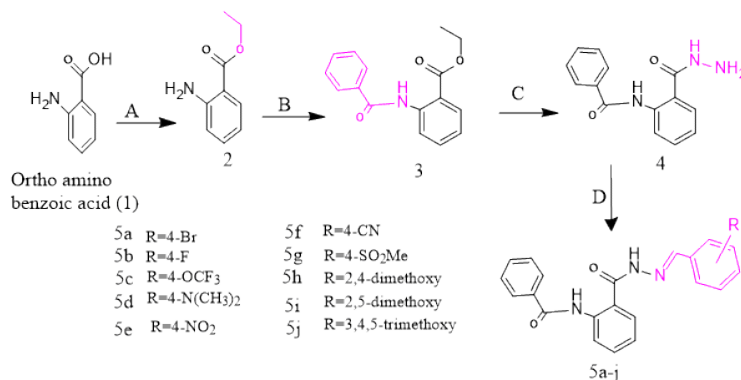


Synthesis and biological evaluation of hydrozone derivatives of ortho-aminobenzoic acid

Dachepally Raju<sup>a,\*</sup>, J. Sreeramulu<sup>a</sup>, P. Malleswarareddy<sup>b</sup> and Nagaraju<sup>c</sup>

Department of Chemistry, Sri Krishnadevaraya University, Anantapuramu, Andhra Pradesh, India-515003.

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A. Conc. H<sub>2</sub>SO<sub>4</sub>, Ethanol, reflux, 10hr. B. Benzyl chloride, TEA, DCM, 10°C, reflux 3hr, C. Hydrazine-hydrate, ethanol, reflux, 6hr. D. Aldehydes (a-j), Ethanol reflux, 1hr.



**Synthesis, spectral and biological properties of 4-aminoantipyrine-based schiff base transition metal complexes.**

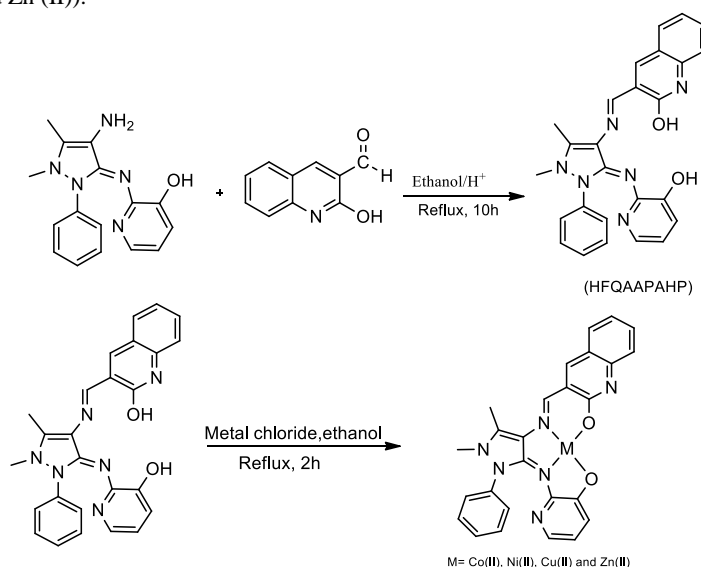
Shyam A. Takle<sup>1\*</sup>, Sunil R. Mirgane<sup>2</sup>

<sup>1</sup>Department of Chemistry, Sant Dnyaneshwar Mahavidyalaya Soegaon, Aurangabad-431120

<sup>2</sup>Department of Chemistry, J.E.S. College, Jalna-431203

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4-Aminoantipyrine moiety containing Schiff bases mono nuclear complexes has been synthesized by reaction of transition metals (Co (II), Ni (II), Cu (II) and Zn (II)).



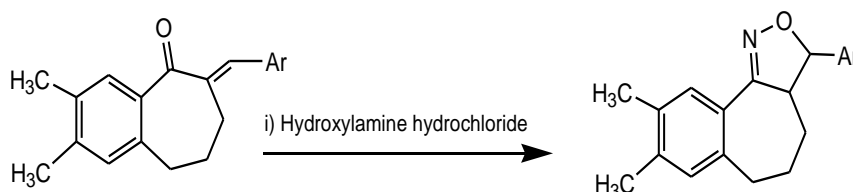
**Synthesis and Biological Studies of Isoxazoles from Benzosuberones**

Srinivas Bathini\* and Sadvik. B

Srizanta Bio Lab, Research and development, Dammiguda, Hyderabad, Telangana, 500048, India.

Corresponding author, E-mail: [drbathinis@gmail.com](mailto:drbathinis@gmail.com)

6-Arylidene-3,4-dimethyl-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-5-ones (**6a-g**) were obtained by the condensation of 3,4-dimethyl-6,7,8,9-tetrahydro-5H-benzo[a]cyclohepten-5-ones (**5**) with appropriate aromatic aldehydes. Cycloaddition of **6a-g** with hydroxylamine hydrochloride in alkaline medium yielded 8,9-dimethyl-3-phenyl-3a,4,5,6-tetrahydro-3H-benzo[6,7]cyclohepta[c]isoxazole derivatives **7a-g**.



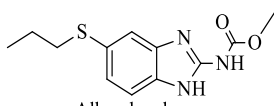
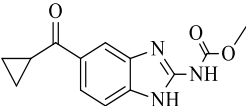
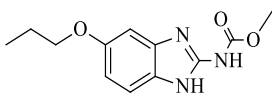
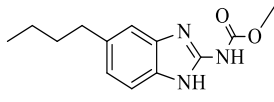
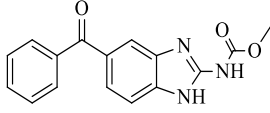
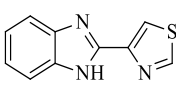




## PERSPECTIVE

Perspective No.1	Heterocyclic Letters 12: iss.-1 (2022), 149-154
<b>Clay-Supported Reactions for the Synthesis of Biologically Active Molecules</b>	
<b>Bimal Krishna Banik</b>	
<i>Department of Mathematics and Natural Sciences, College of Sciences and Human Studies, Deanship of Research, Prince Mohammad Bin Fahd University, Al Khobar 31952, KSA; Email: <a href="mailto:bimalbanik10@gmail.com">bimalbanik10@gmail.com</a>; <a href="mailto:bbanik@pmu.edu.sa">bbanik@pmu.edu.sa</a></i>	
Clay has been used successfully to perform diverse reactions to prepare a variety of organic compounds. In some instances, the acidity of the clay is responsible for the success of these reactions. Some reactions require clay in combination with other acidic catalysts.	

## REVIEWS

Review No.1	Heterocyclic Letters 12: iss.-1 (2022), 155-168	
<b>Mini review on biological activity of imidazole and their derivatives</b>		
<b>F.M. Zahou<sup>a</sup>, Ruba A. Alolayan<sup>b</sup>, Nadia A. A. Elkanzi<sup>b,c</sup></b>		
<sup>a</sup> Biology Department, college of Science, Jouf University, sakaka, 2014, Saudi Arabia		
<sup>b</sup> Chemistry Department, college of Science, Jouf University, sakaka, 2014, Saudi Arabia		
<sup>c</sup> Chemistry Department, Faculty of Science, Aswan University, Aswan, 81528, Egypt		
Corresponding author (N.A.A.Elkanzi)		
*e-mail: <a href="mailto:nahasan@ju.edu.sa">nahasan@ju.edu.sa</a> & <a href="mailto:kanzi20@yahoo.com">kanzi20@yahoo.com</a>		
In our review paper we discuss the imidazole derivatives as drug		
 Albendazole	 Ciclobendazole	 Oxibendazole
 Parbendazole	 Mebendazole	 Thiabendazole



**Advances and perspectives of fe metal nanoparticles synthesized in ionic liquid and their applications**

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