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CONTENTS

Papers

1291 selenoureas by iodine-mediated deselenization

Synthesis of carbodiimides from 1,3-disubstituted A rapid and facile one-pot synthesis of symmetrical and unsymmetrical carbodiimides from 1,3-disubstitued selenoureas by iodine-mediated oxidative deselenization in high yields is reported. Selenophilicity of iodine is explored to obtained carbodiimides having alkyl/aryl substitution, which were not easily accessible from the earlier reported protocols.

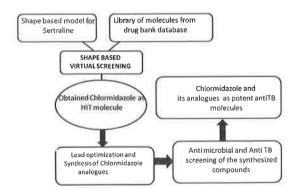
Chlormidazole and its analogues have been synthesized and

screened for anti microbial and anti-TB activities.

L Santhosh, Shekharappa, L Roopesh Kumar & Vommina V Sureshbabu*

109, Peptide Research Laboratory, Department of Studies in Chemistry, Central College Campus, Dr. B. R. Ambedkar Veedhi, Bangalore University, Bangalore 560 001, India

Computer assisted drug repurposing: Anti TB An approach towards shape based virtual screening method has 1295 been conceptualized for drug repurposing. It has helped in finding activity in non antibiotics an obsolete drug chlormidazole with potent TB activity.



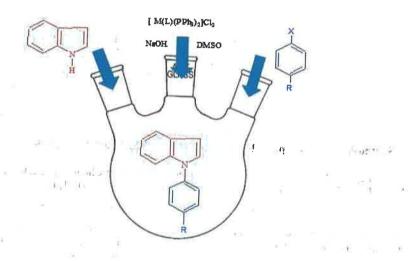
G P V Sangeeta, K Purna Nagasree, Risy Namratha Jamullamudi & Murali Krishna Kumar Muthyala*

Medicinal Chemistry Research Labs, AU College of Pharmaceutical Sciences, Andhra University, Visakhapatnam 530 003, India

N-Arylation of heterocycles by mixed ligand Mixed ligand transition metal complexes [M(L)(PPh₃)₂]X₂ 1-4 where L-2-transition metal complexes under mild conditions phenyl-3-benzovlamino-12-dihydro-quinazolin-4(34), one (PRADO)

Mixed ligand transition metal complexes [M(L)(PPh₃)₂]X₂ 1-4 where L-2-phenyl-3-benzoylamino-1,2-dihydro-quinazolin-4(3H)-one, (PBADQ) where M is Co, Ni, Cu, Zn and X is Cl, have been synthesized and characterized by elemental analysis, IR, NMR spectral studies and their role as catalyst studied in N-arylation of indole and isatin with aryl halides. It has been observed that all the complexes worked as efficient catalysts, especially [Ni(L)(PPh₃)₂]Cl₂ complex 2 which is found to be most suitable for N-arylation of indole and isatin.

Nu: Indole, Isatin X: Cl, Br, I



S S Hegade*, G A Gaikwad & G N Mulik

Department of Chemistry, Shrimant Babasaheb Deshmukh Mahavidyalaya, Atpadi 415 301, Dist Sangli, India (Affiliated to Shivaji University Kolhapur)

Antibacterial and antifungal activities of some The investigation of antibacterial and antifungal activity of five hydroxyacetophenone derivatives

and mentally by

The investigation of antibacterial and antifungal activity of five synthesized compounds 1-5 have been reported. The synthesized compounds have demonstrated poor antifungal, but good antibacterial activity.

I G Mamedov*, A E Farzaliyeva, Y V Mamedova, N N Hasanova, M R Bayramov & A M Maharramov

Baku State University, Chemical Faculty, Z. Khalilov 23, Baku, Azerbaijan

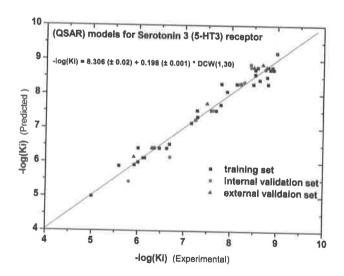
Synthesis, characterization, anti inflammatory Novel pyrazoline derivatives incorporated chalcone backbone 1315 pyrazoline derivatives bearing chalcone backbone

activity and molecular docking study of novel have been synthesized, characterized and evaluated for their antiinflammatory activity.

Upendra Bhadoriya* & Dinesh Kumar Jain

IPS Academy College of Pharmacy, A. B. Road, Rajendra Nagar, Indore 452 012, India

1322 Simplified molecular-input line-entry system based quantitative structure-activity relationship (QSAR) models for serotonin 3 (5-HT₃) receptor



Sanija Begum, P Ganga Raju Achary*, Andrey A Toropov & Alla P Toropova

Department of Chemistry, Institute of Technical Education and Research (ITER), Siksha 'O' Anusandhan University, Bhubaneswar 751 030, India