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(Organic including Medicinal)



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# **Indian Journal of Chemistry**

## Sect. B: Organic Chemistry including Medicinal Chemistry

**VOL. 58B June 2019** NUMBER 6 **CONTENTS Papers** 669 Stereoselective synthesis of subincanadine alkaloids framework MeMgBr/CH<sub>3</sub>CHO 8 steps (-)-Indolizinoindolone N-Tosyl-(S)-Acetoxysuccinic (subincanadine framework) tryptamine anhydride Manojkumar G Kalshetti & Narshinha P Argade\* Division of Organic Chemistry, CSIR-National Chemical Laboratory, Pune 411 008, India 674 An efficient one-pot four-component synthesis of 1H-pyrazolo[1,2-b] phthalazine-5,10-dione derivatives catalyzed by proline Proline. **RCHO** EtOH:H2O (2:1), 3 CN NH CN 80°C, Ar atm CN NH<sub>2</sub> Ö 1 Δ NH<sub>2</sub>NH<sub>2</sub>.H<sub>2</sub>O 5a-r (62-96 %) 2 R = Aromatic or aliphatic group Dipanwita Banerjee & Gourhari Maiti\* Department of Chemistry, Jadavpur University, Kolkata 700 032, India

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Synthesis, free radical scavenging and α-glucosidase inhibitory activities of 2*H*-chromenylphenyloxazolones A series of 2*H*-chromenylphenyloxazolones have been prepared starting from 2*H*-chromene-3-carbaldehydes. The compounds have been evaluated for their DPPH,  $ABTS^+$  free radical scavenging and  $\alpha$ -glucosidase inhibitory activities. Compound 4g has been identified as the most potent ABTS.+ free radical scavenger when compared to the standard drug. The compounds 4k, 5a-c and 5g are identified as potent  $ABTS^+$  free radical scavengers in the present series of the compounds. Compound 5g has been identified as a promising  $\alpha$ -glucosidase inhibitor. Molecular modeling studies have been carried out for compound 5g to explain the molecular basis of  $\alpha$ -glucosidase inhibitor.



E Varsha Reddy, K S Hariprasad, A Zehra, P Vijaykumar, A K Tiwari, A Addlagatta & B China Raju\*

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691 Biocatalyst mediated synthesis of tryptanthrins performed under ultrasonication



#### Ananda Mane, Siddharth Kamat, Audumbar Patil & Rajashri Salunkhe\*

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A simple synthesis of 1,3-di-aryl-quinolone derivatives by palladium catalyzed cross-coupling reaction

An approach for the preparation of N-aryl-quinolone via enaminone synthesis and derivatization at 3 position has been accomplished by the palladium catalyzed Suzuki cross coupling reaction.



## P Ravi Kumar, A Jaya Shree, K Raghavulu, K Basavaiah, Venu Kandula, Anindita Chatterjee, Satyanarayana Yennam & Manoranjan Behera\*

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WEB: A green and an efficient catalyst for Knoevenagel condensation under grindstone method A greener, economic and efficient approach for Knoevenagel condensation of substituted heteroaryl/aryl aldehydes with malononitrile catalyzed by WEB under grindstone method at room temperature is described. The total process is advantageous over conventional methods that involve reflux, heating, expensive approaches. The present approach also reduced reaction time.



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Solid-phase synthesis and solution conformation of two overlapping fragments of SPFE peptide using 2D-NMR



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