

Synthesis of Some New Scaffolds of Isoxazolidine & Isoxazoline Derivatives using Novel Class of Nitrones via 1,3-Dipolar Cycloaddition Reaction using Greener Chemistry Protocol and their Further applications including Potential Biological activities of the Cycloadducts

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Abstract

Some environment friendly greener methodologies have been described for the synthesis of new isoxazolidine and isoxazoline derivatives using new nitrones via 1,3-dipolar cycloaddition reactions. These also include synthesis of bisisoxazolidine and bisisoxazoline derivatives synthesized from glyoxal and terephthalaldehyde respectively. Few new spiro isoxazolidine derivatives have been also reported using new dipolarophiles. Furthermore, these new isoxazolidine and isoxazoline derivatives are found to have vast synthetic potential as they could be used as precursors for the synthesis of a variety of new organic molecules including peptides, 1,3-amino alcohols with potential biological activities. For the synthesis of peptides, it has been observed that CDMT (chloro dimethyl triazine) has found to be better coupling reagent than conventional DCC (dicyclohexyl carbodiimide) due to the formation of insoluble by-product (*N,N*-dicyclohexylurea) and purification becomes tedious. The new nitrones reported are synthesized from furfural, dihydropyran, chlorohydrin, glyoxal, terephthalaldehyde and formamide respectively. Significant increase in the reaction rates, excellent yields, and high selectivity (diastereo and regioselectivity) are the important features observed in these cycloaddition reactions following greener methodologies. It has been observed that high diastereoselectivity in these cycloaddition reactions have been observed when the reactions are performed in water. Synthesis of aldehydes and ketones with new nitrones in atom efficient reactions are the most attractive features as they have future scopes in these reactions. The side products (enamines) obtained during the synthesis of aldehydes and ketones has been successfully utilized as new dipolarophiles in these cycloaddition reactions for the synthesis of spiro cycloadducts. Potential biological activities including cytotoxicity of the new molecules have made these new syntheses much more attractive and useful as well.

Biography:

Professor of Chemistry at Sikkim Government College (NBBDC), Gangtok, Sikkim, India with 25 years of teaching experience in teaching Undergraduate and Post graduate Chemistry students. Has active research experience of 24 years in synthetic organic chemistry in the field of "Nitrone cycloaddition reactions and their further applications following green chemistry methodologies". Has established few environment friendly greener methodologies along with atom efficient reactions in the synthesis of few novel isoxazolidine and isoxazoline derivatives using new nitrones via 1,3-dipolar cycloaddition reactions.

Speaker Publications:

1. Chakraborty, B. J. Heterocyclic. Chem. 2019, DOI: 10.1002/jhet.3736 (Online published).
2. Chakraborty, B. J. Heterocyclic. Chem. 2019, DOI: 10.1002/jhet.3804 (Online published).
3. Chakraborty, B. ; Rai, N. J. Heterocyclic. Chem. 2018, 55, 1053-1063.
4. Chakraborty, B. ; Chettri, E. J. Heterocyclic. Chem. 2018, 55, 1157-1165.
5. Chakraborty, B.; Chettri M.S; Chettri E. J. Heterocyclic Chem. 2017, 54, 110-120.

10th European Chemistry Congress:

Vienna, Austria- July 15-16, 2020.

Abstract Citation:

Bhaskar Chakraborty, Synthesis of Some New Scaffolds of Isoxazolidine & Isoxazoline Derivatives using Novel Class of Nitrones via 1,3-Dipolar Cycloaddition Reaction using Greener Chemistry Protocol and their Further applications including Potential Biological activities of the Cycloadducts, Euro Chemistry 2020, 10th European Chemistry Congress; Vienna, Austria- July 15-16, 2020. (<https://europe.chemistryconferences.org/abstract/2020/synthesis-of-some-new-scaffolds-of-isoxazolidine-isoxazoline-derivatives-using-novel-class-of-nitrones-via-1-3-dipolar-cycloaddition-reaction-using-greener-chemistry-protocol-and-their-further-applications-including-potential-biological-activities-of-the->)

