



**Synthesis of ethoxyphthalimido derivatized thiadizole assembled imidazolidinone and chloroazetidione systems from common intermediate Schiff's bases and evaluation of their antibacterial activity**

**Monika Kumawat and Ganpat L.Talesara\***

\*Synthetic Organic Chemistry Research Laboratory, Department of Chemistry,  
Mohan Lal Sukhadia University, Udaipur-313001, Rajasthan, **INDIA**

Email: [glntalesara@yahoo.com](mailto:glntalesara@yahoo.com), [monika.kumawat84@gmail.com](mailto:monika.kumawat84@gmail.com)

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**ABSTRACT**

*In present investigation synthesis of 2-(2-(3-(5-((2-methyl-1H-enzo[d]imidazol-1-yl)methyl)-1,3,4-thiadiazol-2-yl)-4-oxo-2-phenylimidazolidin-1-yl)ethoxy)isoindoline-1,3-dione **9a-d** and 3-chloro-4-(4-phenyl)-1-(5-((2-methyl-1H-benzo[d]imidazol-1-yl)methyl)-1,3,4-thiadiazol-2-yl)azetidion-2-one **6a-d** are described. 2-Methyl benzimidazole **1** is converted to carbothioamide **3** by the reaction with ethylchloroacetate followed by thiosemicarbazide. It is cyclised to thiadizole **4** by treatment with conc. H<sub>2</sub>SO<sub>4</sub> and NH<sub>3</sub>. Condensation of **4** with different araldehyde yielded corresponding Schiff's bases **5a-d**. Compounds **5a-d** has acted as key intermediate for both series of the final compounds. In one pathway, **5a-d** are converted to azetidione plugged compounds **6a-d** by treatment with chloroacetylchloride in presence of triethylamine. In an another route, reaction of **5a-d** with glycine has afforded the formation of 3-(5-((2-methyl-1H-benzoimidazol-1-yl)methyl)-1,3,4-thiadiazol-2-yl)-2-phenylimidazolidin-4-one **7a-d**. Finally the targeted molecules **9a-d** were obtained by the base induced condensation of **7a-d** with bromoethoxyphthalimide. Structure confirmation was accomplished by spectral studies (IR, <sup>1</sup>HNMR, Mass) and elemental analysis of all the synthesized compounds.*

**Keywords:** Bromoethoxyphthalimide; Azetidione; Benzimidazole; Thiadizole; Chloroacetylchloride.

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