**The synthesis of derivatives of 1,3-thiazolan-4-one and exhibit the desired biological and pharmacological activities: A Review**

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**ABSTRACT**

The researchers have extensively investigated the approaches for the develop- ment of the synthesis of derivatives of 1,3-thiazolan-4-one, and those that are practical for the synthesis of titile compounds 1,3-thiazolan-4-one and its derivatives, which are significant and exhibit the desired biological and pharmacological activities

**Keywords:** 1,3-thiazolan-4-one, pharmacological activity, development, biological

**1. INTRODUCTION**

Because many biologically active molecules, including thiamine, penicillin, and other antibiotics1,2, include these heterocyclic rings, the nitrogen and sulfur-containing heterocyclic compounds are important from a biological perspective. The same goes for 1,3-thiazolan-4-one and its derivatives, which are significant and exhibit the desired biological and pharmacological activities, such as hypnotic3, antiinflammatory4, antibacterial5, antifungal6, antitubercular7, anticancer8, antitumor9, analgesic10, anesthetic11, antiproliferative12, anti-HIV13, and nematicidal14. 1,3-Thiazolan-4-one have also been utilised to treat schizophrenia15, diabetic problems such cataract, nephropathy, and neuropathy16, as well as selective antiplatelet activating factor (PAF)17. Additionally, cyanine dyes, which are utilised in the photographic film industry18, are made using 1,3-thiazolan-4-one derivatives. Some of the important biologically active heterocyclic compounds incorporating the 1,3-thiazolan-4-one nucleus are briefly discussed in the following few pages.

The researchers have extensively investigated the approaches for the develop- ment of the synthesis of derivatives of 1,3-thiazolan-4-one, and those that are practical for the synthesis of titile compounds were briefly reviewed in the following sections.

**Scheme 1.** Desai *et al*.19 reported the heterocyclization reaction of 2-(benzothiazol-2-ylthio)-*N*'-benzylideneacetohydrazide with thioglycolic acid in DMF in the presence of a catalytic amount of anhydrous ZnCl2 under microwave irradiation is described and compared with conventional synthesis methods. This reaction produced 4-thiazolidinones in good yields. Elemental analyses and spectrum data allowed for the complete elucidation of all freshly synthesised compounds' structures. Some of the new compounds were tested against bacteria (Gram− ve and Gramt+ ve) and fungi. (**Scheme 1**).



**Scheme 1**

**Scheme 2.** Tetrahydro-6-pyridazinone-3-carboxylic acid hydrazides of 1,4,5,6-tetrahydro-6-pyridazinone-3-carboxylic acid react with aromatic aldehydes to produce hydrazones of 1,4,5,6-tetrahydro-6-pyridazinone-3-carbonyl aromatic aldehydes. The 1,3-thiazolidinone derivatives were produced by cyclocondensing the intermediate 1,3,4-oxadiazole with acetic anhydride in absolute ethanol and then cyclizing it with mercaptoacetic acid in DMF in the presence of anhydrous ZnCl2. (**Scheme 2**)20.

**Scheme 2**

**Scheme 3.** Parekh *et al*.21 reported the thiazolidinones derivatives by the cyclocondensa- tion of the 2-amino-4-(-methoxyiminocarbomethoxymethyl)-thiazole with mercapto- acetic acid and mercaptopropionicacid in dry benzene. Spectral data and elemental analysis were used to determine the compounds' structures. The products' ability to prevent the growth of several bacteria *in vitro* was tested. Some of the substances had notable antifungal and antitubercular action (**Scheme 3**).



**Scheme 3**

**Scheme 4.** Based on the starting material 2-[(2,6-dichlorophenyl)amino] phenylacetic acid, Patel *et al*.22 reported synthesis of 1,3-thiazolan-4-one derivatives, screening them for antibacterial and antifungal activity at two concentrations, and comparing them to penicillin-G, ampicillin, and amoxicillin. When compared to the usual medications, the compounds containing 4-OCH3 & 5-(OCH3)3 displayed good action (**Scheme 4**).



**Scheme 4**

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**Scheme 5.** Two novel series of 4-thiazolidinone derivatives, including 2-(5-nitro-2-furyl/ substituted phenyl)-4-thiazolidinone-3-yl]amides of 2',4'-difluoro-4-hydroxybiphenyl-3-carboxylic acid and 2-(2',4'-difluoro-4-hydroxybiphenyl-3-carbonylhydrazono) 1-(2,4-difluoro-4-hydroxybiphenyl-3-carbonyl)4-arylthiosemicarbazides. The antimyco- bacterial, antiviral, and antimicrobial properties of all produced compounds were tested against Mycobacterium tuberculosis H37Rv as well as against other virus, bacteria, and fungal strains. (**Scheme 5**)23.



**Scheme 5**

**Scheme 6.** Panwar *et al*.24 synthesised a number of 1,3-thiazolan-4-one derivatives and tested them for antimicrobial susceptibility against *S. aureus, E. coli, K. pneumonia, P. vulgaris*, and *A. fumigates*, *C. albicans, A. albicans*, and *C. krusei*, respectively. Some of the synthetic compounds displayed the strongest antibacterial and antifungal effects while still being completely toxin-free. (**Scheme 6**).

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**Scheme 6**

**2. CONCLUSION**

This review provided a resource for basic and application study on the topic by outlining 1,3-thiazolan-4-one and its derivatives.

**3. ACKNOWLEDGEMENTS**

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**4. CONFLICTS OF INTERESTS**

There is no conflict of interest with the publication of this essay, according to the author.

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