Bulletin of Pharmaceutical Research 2014;4(2):72-80

An Official Publication of Association of Pharmacy Professionals

ISSN: 2249-6041 (Print); ISSN: 2249-9245 (Online)

RESEARCH ARTICLE



SYNTHESIS AND BIOLOGICAL EVALUATION OF CLUBBED TRIAZOLE-THIAZOLIDINONE DERIVATIVES

Prabodh Chander Sharma^{*1}, Deepak Kumar¹, Ritu Gorsi¹, Archana Sharma¹ and Harish Rajak²

¹Institute of Pharmaceutical Sciences, Kurukshetra University, Kurukshetra-136 119, Haryana, India ²SLT Institute of Pharmaceutical Sciences, Guru Ghasidas University, Bilaspur-495 009, Chhattisgarh, India

**E-mails*: sharma_prabodh@rediffmail.com, deepakkash4340@gmail.com *Tel*.: +91 9416025460.

Received: November 19, 2013 / Revised: June 28, 2014 / Accepted: June 29, 2014

In the present work, sixteen novel 3-(2-(4-(4-substituted benzylideneamino)-5-phenyl-4*H*-1,2,4triazol-3-ylthio)acetyl)-2-(arylimino)thiazolidin-4-one (10a-p) derivatives were synthesized by clubbing 3-(2-chloroacetyl)-2-(arylimino)thiazolidin-4-one derivatives (4a-h) with SH group of triazole Schiff bases (9a-b). The structures of the newly synthesized compounds were confirmed by analytical and spectral methods (¹H NMR and IR). The biological potential of newly synthesized compounds was investigated through hydrogen peroxide scavenging assay, anthelmintic activity and *in vitro* α -amylase inhibition activity. Among synthesized compounds, 10l showed the most potent hydrogen peroxide scavenging activity 71.44%, 72.32% and 73.99% at different concentrations 10 μ g/ml, 30 μ g/ml, and 50 μ g/ml respectively. Compound 10f showed the most potent anthelmintic activity with mean paralysis time 5.20±0.05 min and mean death time 7.64±0.16 min. Highest α -amylase inhibition activity 92.17% was exhibited by compound 10j.

Key words: Thiazolidin-4-one, 1,2,4-triazoles, Schiff's base, Hydrogen peroxide scavenging assay.

INTRODUCTION

Heterocyclic systems have been enriched with pharmacological activities since ancient times (Kumar, 2011; Mehta and Pathak, 2011; Dahiya and Mourva, 2013; Pareta et al 2013; Nusrat et al 2014). Among the heterocyclics, thiazolidin-4one is a biologically important scaffold known to be associated with several biological activities such as antimicrobial (Deep et al 2014), anticancer (Wang et al 2012), antidiabetic (Ottana et al 2012) analgesic, anti-inflammatory (Deep et al 2012), hydrogen peroxide scavenging activity (Sharma et al 2011) and anticonvulsant activity (Kaur et al 2012) etc. Another five membered heterocycles like 1,2,4-triazole and their derivatives constitute an important class of compounds which act as a core nucleus of therapeutically various important drugs. Triazole derivatives also exhibit a broad array of agricultural, industrial and biological activities. The synthesis of these heterocycles has received considerable attention in recent years and several 1,2,4-triazole derivatives have been reported for diverse biological activities. Schiff bases of 1,2,4-triazoles have also been found to possess extensive biological activities (Kharb *et al* 2011; Singhal *et al* 2011; Chandermauli *et al* 2012).

Design of new bioactive agents with the development of hybrid molecules through the combination of different pharmacophores in the same structure may lead to compounds having more efficiency in biological response. The interesting biological properties of these two classes of compounds inspired us to synthesize new compounds with both moieties clubbed together with a hope of improved biological potential. Hence in the present study, we report

