

Notice for the PhD Viva Voce Examination

Mr V P Rama Kishore Putta (Reg. No. 1650082), PhD scholar at CHRIST (Deemed to be University), will defend his PhD thesis at the public viva-voce examination on Wednesday, 16 December 2020 at 3.00 pm on the Webex Meeting platform.

Title of the Thesis : Synthesis of Benzothiazinones,

Benzothiazines and their Selenium

Analogues through Novel Synthetic Routes

Discipline : Chemistry

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The faculty members of the Department and the School, interested experts and research scholars of all the branches of research are cordially invited to attend this open viva.

Place: Bengaluru

Date: 10 December 2020

Registrar

ABSTRACT

Benzo fused N-heterocyclic scaffolds containing oxygen, sulphur or selenium have found wide interest in the field of drug-discovery. Among these N-heterocycles, benzothiazine, benzoxazine, benzoselenazine and benzothiazinone derivatives are a unique class of compounds and have a larger scope towards the development of efficient and simple synthetic methodologies for their synthesis with readily available substrates. During the course of the present thesis a convenient and simple synthetic procedures were developed for the synthesis of benzothiazines, benzoxazines, benzoselenazines and benzothiazinones in an onepot methodology. 2-aryl/alkyl substituted 1,3-benzothiazines and selenazines were synthesized by reacting 2-amino benzyl alcohols and thio or seleno benzamides in the presence of T3P. A reagent controlled methodology was developed for the synthesis of 2amino substituted 1,3-benzothiazines and oxazines. Initially, various 2-amino benzylalcohols are reacted with isothiocyanates to form the corresponding thioureas. The formed thioureas undergo cyclodehydration in the presence of T3P to yield 2-amino substituted 1,3benzothiazines and on the other hand molecular iodine facilitates desulfurization of the thiourea to yield 2-amino substituted 1,3-benzoxazines. 2-amino substituted 1,3benzothiazinones were synthesized by reacting anthranilic acids and isothiocyanates in the presence of EDC.HCl. 2-aryl substituted 1,3-benzothiazinones were synthesized by employing thiobenzamides in the presence of T3P. All the compounds synthesized were characterized by 1HNMR, 13C, Mass spectroscopic (LCMS, HRMS) analysis. Docking studies against TANKYRASE-1 enzyme for colorectal cancer (CRC) and antibacterial studies were also discussed.

Keywords: Benzothiazines, Benzoxazines, Benzoselenazines, Benzothiazinone, Cyclodehyration, Desulfurization, T3P, EDC.HCl, Tankyrase-1, CRC, 1HNMR, 13C, LCMS, HRMS.