



A NOVEL SYNTHESIS OF 4-(5-SUBSTITUTEDTHIOCARBAMIDO)-1,2,4-THIADIAZOLOPYRIDINES

S.G. Khobragade^a, D.T. Tayade^{b*}, S.S. Padhen^c, N.S. Dixit^d

a: Brijlal Biyani Science College, Amravati 444 605.

b: Govt. Vidarbha Institute of Science and Humanities, Amravati 444 604.

c: Rajarshree Shahu Science College, Chandur Railway

d: G.S. Tompe Arts, Commerce & Science college, Chandur Bazar, Amravati (M.S.) India

*Email: skdtayade@gmail.com, sgkhobragade29@gmail.com

Abstract:

A novel series of 4-(5-substitutedthiocarbamido)-1,2,4-thiadiazolopyridines (**IIIa-h**) was synthesized by the interactions of 4-(5-amino)-1,2,4-thiadiazolopyridine (**I**) with various alkyl/aryl isothiocyanate (**IIa-h**) in acetone medium. The synthesized compounds were recrystallised and their structures were justified and established on the basis of elemental analysis, chemical characteristics and through spectral studies.

Key words:

4-(5-amino)-1,2,4-thiadiazolopyridine, various alkyl/aryl isothiocyanate and acetone

Introduction:

The literature survey reveals that thiocarbamido, thiadiazolo nucleus containing drugs possess an important applications and significances in medicinal, pharmaceutical, drug, industrial, agricultural and biochemical sciences[1-9]. The thiadiazolo derivatives of cyanopyridine showed remarkable anti-cancer[10], anti-tumor[11], anti-malarial[12], anti-tubercular[13], anti-fungal[14], anti-bacterial[15], cytotoxic[16], anti-inflammatory and anti-pyretic properties[17-18]. Thiocarbamido, thiadiazolo nucleus containing drugs have been shown to possess a diverse range of physiological activities[19], plant growth promoting activity, herbicidal^[20], amoebicidal[21], anti-diabetic[22] properties. Some thiadiazoles were found to be active against *S. aureus*, *E. coli*, and *C. albicans*[23]. The present work described somewhat suitable and direct