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Research Article

SYNTHESIS OF 3-[4-AMINO-N-SUBSTITUTED-o-ANISAMIDO-5-YL]AMINO-5-SUBSTITUTED-1,2,4-DITHIAZOLE.D.T.Tayade^{1*} R. D. Thombare¹, S.A. Waghmare²¹ Department of Chemistry, Government Vidarbha Institute of Science and Humanities, Amravati 444606.² Department of Chemistry, Ghulam Nabi Azad Arts, Comm. & Science College, Barshitakli, Dist. Akola 444401.**Abstract:**

A novel series of 3-[4-Amino-N-[2-(dithethylamino)ethyl-o-anisamido-5-yl]amino-5-substitutedimino-1,2,4-dithiazole was synthesized by the oxidative cyclization of 4-amino-5-substituteddithiobiureto-N-[2-(dithethylamino)ethyl-o-anisamide in chloroform medium by making the use of liquid bromine as oxidizing agent. The products were characterized and justified on the basis of elemental analysis, chemical characteristics and spectral studies.

Keywords:3-[4-Amino-N-[2-(dithethylamino)ethyl-o-anisamido-5-yl]amino-5-substituted -imino-1,2,4-dithiazole, 4-amino-5-substituteddithiobiureto-N-[2-(dithethylamino)ethyl-o-anisamide, Bromine in Chloroform.

Corresponding author:**D.T.Tayade,**

Department of Chemistry,
Government Vidarbha Institute of Science and Humanities,
Amravati 444606.

Email:- skdtayade@gmail.com,
rupalidhombare30@gmail.com

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INTRODUCTION:

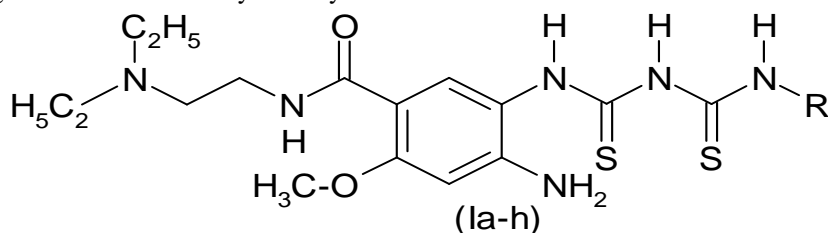
The literature survey shows that the dithiazole, dithiazino and thiadiazino nucleus containing compounds are widely used in medicinal, industrial, agricultural, biochemical sciences [1-7]. It was considered that these nucleus containing drugs possesses, antibacterial [8], antidiabetic [9], amoebicidal [10] herbicidal [11] properties. Iminosubstitutedthiadiazolo nucleus also possesses noticeable pharmaceutical activities.

The heterocyclic compounds containing nitrogen, nitrogen and sulphur have gained huge important in our life. It was shows that thiadiazoles are also effective against copper corrosion [12] and additive in lubricating oil [13].

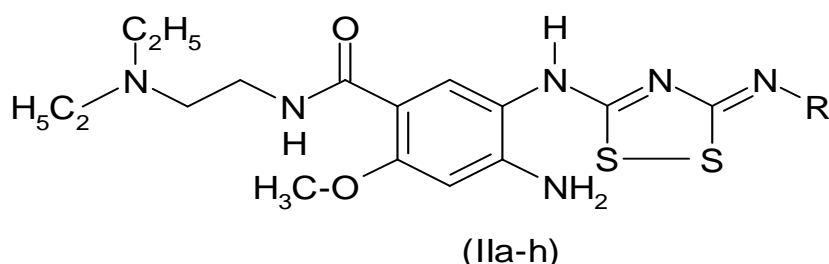
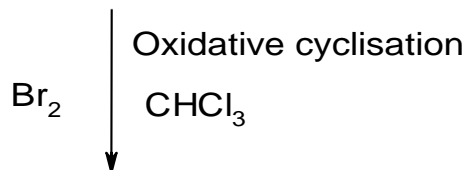
In recent years several 1,2,4-dithiazole and their derivatives were found to have prominent pharmacological activities such as anticonvulsant, analgesic anti-inflammatory activity. Dobolkar and

Ansari [14] had successfully investigated oxidative cyclisation of cyanoamidinosubstitutedthiocarbamide and N-substitutedformidinothiocarbamides. Various researchers [15-19] studied oxidative cyclisation of 1,3,4-thiadiazoles, 1,3,4-thiadiazolines and 1,2,4-triazoles.

As a part of research work presently undertaken in this laboratory in the synthesis of heteroacycles and heterocycles, it was thought interesting to investigate the cyclisation of 4-amino-5-substituteddithiobiureto-N-[2-(diethylamino)ethyl]-o-anisamides with liquid bromine in chloroform medium to obtain a novel series of 3-{4-amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-substitutedimino-1,2,4-dithiazoles which are heither to unknown. The present work describes suitable, convenient and some what direct method for this synthesis and depicted below



4-Amino-5-substituteddithiabiurato-N-[(2-diethylamino)ethyl]-o-anisamide



3-{4-Amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-substitutedimino-1,2,4-dithiazole.

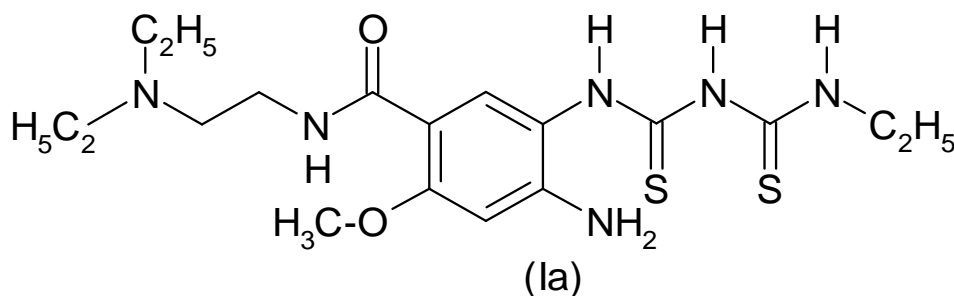
R = , methyl, t-butyl, phenyl, p-chlorophenyl, ethyl, o-tolyl, m-tolyl, p-tolyl.

EXPERIMENTAL:

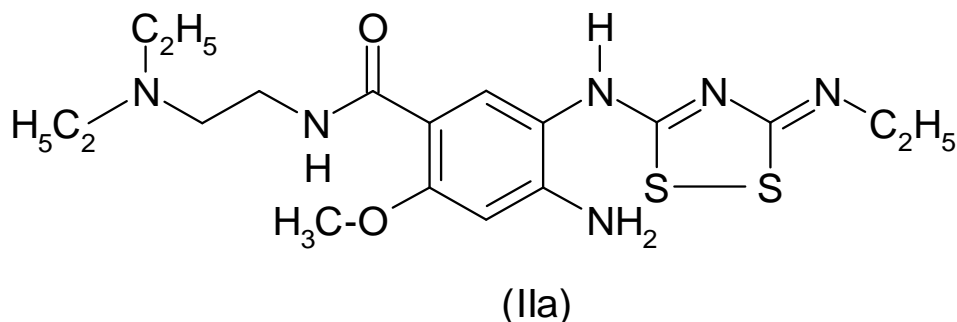
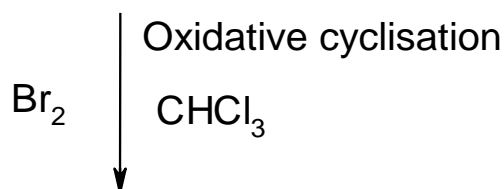
The melting point of the all synthesized compounds was recorded using hot paraffin bath. The carbon and hydrogen analysis were carried out on Carlo-Ebra 1106 analyzer. Nitrogen estimation was carried out on Colman-N-analyzer-29. IR spectra were recorded on Perkin Elmer Spectrometer in range 4000-400 cm^{-1} in KBr pellets. PMR spectra were recorded on Bruker 400F spectrometer with TMS as internal standard using CDCl_3 and DMSO-d_6 as solvent. The purity of compound was checked on silica Gel-G Pellets by TLC with layer thickness of 0.3 mm. All chemicals used were AR-grade.

Synthesis of 3-{4-Amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl} amino-5-ethylimino-1, 2, 4-dithiazole

3-{4-Amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl} amino-5-ethylimino-1,2,4-dithiazoles was synthesized by the oxidative cyclization of 4-amino-5-ethylthio-biureto-N-[2-(diethylamino)ethyl]-o-anisamide with liquid bromine in presence of chloroform. A paste of 4-amino-5-ethylthio-biureto-N-[2-(diethylamino)ethyl]-o-anisamide was prepared in chloroform to it 10% liquid bromine in chloroform was added with constant stirring. Initially the colors of bromine disappear in the reaction mixture. The reaction mixture was allowed to stand for 8 hours, it afforded brown colored product. It was recrystallised from ethanol, yield 90%, m.p. 251°C.



4-Amino-5-ethylthio-biureto-N-[2-(diethylamino)ethyl]-o-anisamide



3-{4-Amino-N-[2-(diethylamino)-ethyl]-o-anisamido-5-yl}-amino-5-ethylimino-1,2,4-dithiazole.

Table No. 1

Sr. No.	3-{4-Amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-substitutedimino-1,2,4-dithiazoles	Yield (%)	m.pt. (°C)
1.	3-{4-Amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-methylimino-1,2,4-dithiazole	88	245
2.	3-{4-Amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-t-butylimino-1,2,4-dithiazole	81	249
3.	3-{4-Amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-p-chlorophenyl imino-1,2,4-dithiazole	85	265
4.	3-{4-Amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-otolylimino-1,2,4-dithiazole	90	261
5.	3-{4-Amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-m-tolylimino-1,2,4-dithiazole	80	255
6.	3-{4-Amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-p-tolylimino-1,2,4-dithiazole	90	270

PROPERTISE:

It is brown colour crystalline solid having melting point 251°C. It gave positive test gave for nitrogen and sulphur. It was desulphurized by alkaline plumbite solution. It formed picrate having melting point 179°C. **Elemental Analysis** :C[(found 50.03%) calculated 50.94], H[(found 06.20%) calculated 06.60], N[(found 19.00%) calculated 19.81], S[(found 14.50%) calculated 15.09]. **IR Spectrum**: The IR spectrum was carried out in KBr pellets : 3429.00 (N-H stretching), 2923.24 (Ar-C=C stretching), 1639.05 (N-C=O stretching), 1506.08 (C=N stretching), 1092.21 (C-N stretching), 774.31 (C-S stretching). **NMR Spectrum** : The NMR spectrum of compound was carried out in CDCl₃ and DMSO-d₆. This spectrum distinctly displayed the signals due to Ar-H protons at δ 7.3000-7.152 ppm, -NH proton at δ 4.9072-4.2568 ppm, NH₂ protons at δ 3.9058-3.2062 ppm, -OCH₃ protons at δ 3.2009-3.0183 ppm, CH₂ protons at δ 2.9955-2.0929 ppm and N-CH₃ protons at δ 1.2269-1.1268 ppm.

From the above properties and spectral analysis of the compound (**IIa**) was assigned the structure as 3-{4-amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-ethylimino-1,2,4-dithiazoles (**IIa**)

Similarly, 4-amino-5-methylthiobiureto-N-[2-(diethylamino)ethyl]-o-anisamide (**Ib**), 4-amino-5-t-butylthiobiureto-N-[2-(diethylamino)ethyl]-o-anisamide (**Ic**),

4-amino-5-p-chlorophenylthiobiureto-N-[2-(diethylamino)ethyl]-o-anisamide (**Id**), 4-amino-5-otolylthiobiureto-N-[2-(diethylamino)ethyl]-o-anisamide (**Ie**), 4-amino-5-m-tolylthiobiureto-N-[2-(diethylamino)ethyl]-o-anisamide (**If**), 4-amino-5-p-tolylthiobiureto-N-[2-(diethylamino)ethyl]-o-anisamide (**Ig**), were successfully oxidative cyclised with bromine in chloroform medium to isolate 3-{4-amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-methylimino-1,2,4-dithiazole (**IIb**), 3-{4-

amino-N-[2-(diethyl-amino)ethyl]-o-anisamido-5-yl}-amino-5-t-butylimino-1,2,4-dithiazole (**IIc**), 3-{4-amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-o-chlorophenylimino-1,2,4-dithiazole (**IId**), 3-{4-amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-otolylimino-1,2,4-dithiazole (**IIe**), 3-{4-amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-m-tolylimino-1,2,4-dithiazole (**IIIf**), 3-{4-amino-N-[2-(diethylamino)ethyl]-o-anisamido-5-yl}-amino-5-p-tolylimino-1,2,4-dithiazole (**IIg**) respectively by the above mention method in **Table No. 1**

REFERENCES:

- Bhattacharyya A, Singh T, and Verma V.K., *Tribol.Int.*, 28(3), **1995**,189.
- Ghosh S.K., *Advanced Organic Chemistry*, 2nd Ed., Calcutta, **1998**, (a) P-410,(b)P-412.
- Lapman G., Pavia D. and Kriz G., *Introduction to Spectroscopy*, Asia a Pta Ltd., 3rd Ed., Singapor, **2004**, (a) P-68-69,(b) P-43.
- Tayade D.T., *Asian Jr. of Chemistry*, 7(4),**1995**, 890-891.
- Hamady N.A., Abdel-Aziz H.A., Farog A.M. and Fakhr Issa. M.A., *Monatshefte fur chemie.*,138, **2007**,1001-1010.
- Patel B.V., Patel H.S. and Patel K.C., *Ind. J. Chem.*, B, 47B (06), **2008**, 0376-4699.
- Ali T. El-sayad and Ibrahim M.A. *J.Braz. Chem. Soc.*, 446(2117) , **2010**, 1445-1468.
- Chan-Thaw C.E., Villa A., Katekonon P., Sus D., Thomas A. and Prate L.,*NanoLett.*, 10(2), **2010**, 537-541.
- Kurumurthy C., Veeraswamy B. Rao P. S., Kumar G.S., Reddy V.L., Rao J.V. and Narsaiah B., *Bioorganic and Medicinal Chemistry Lett.*, 24(3),**2014**, 746-749.
- Sztanke K., Pasternak K., Rajtar B., Sztakne M., Majek M. and Polz Dacewicz M., *J.Bioorganic and Med.Chem.*,15, **2007**, 5480-5486.

11. Simanek E.E., Abdou H., Lalwani S., Lim J.J., Mintzer M., Venditto V.J. and Vittur B., *Proc. R. Soc. A*, 466(2117), **2003**, 1445-1468.
12. Krutz L.J., Shaner D.L., Weaver M.A., Webb R.M.T Zablutowicz R.M., Reddy K.N. Huang Y. and Thomson S.J., *Pest Management Sci.*, 66(5), **2010**, 461-481.
13. Sherif E.M., Park S.M., *Electrochimica Acta*, 51, **2006**, 6556-6562.
14. Dobolkar V.V., Ansari F.Y., *Acta Polonica Pharmacologica-Drug research*, 65(5), **2008**, 521-526.
15. Lim J., Mintzer M.A., Perez L.M. and Simanek E.E., *Org. Lett.*, 12(6), **2010**, 1148-1151.
16. Mirano K., *Chem. Abstr.*, 79, **1973**, 137200.
17. Zhuo J., He C., Yao W., *United States, Patent Application Publication*, US2013/0345224 A1, **2013**.
18. Pittis W.J., Guo J., Dhar T.G., Shen Z., Gu H., Watterson S.H. and Bednarz M.S., *J. Bioorg. Med. Chem.*, 12(2), **1997**.
19. Hajiduk P.J., Dinges J., Schkeryantz J.M., Janowick D., Kaminski M., Tufano M., Augeri., *J. Med. Chem.*, 42, **1999**, 3852-3859.