SYNTHESIS OF 2-AMINO-5-(3,5,6-TRICHLORO-1,4-BENZOQUINONE-2-YL) THIAZOLES

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We present an original two-step synthesis of compounds that contain electron donor (aminothiazole) and electron acceptor (trichlorobenzoquinone) residues; these are of interest in the study of inter- and intramolecular charge transfer.

We determined for the first time that when equimolar amounts of 2-(2-dialkylaminovinyl)-3,5,6-trichloro-1,4-benzoquinones (Ia,b) [1] and thiourea or N-monosubstituted thiourea are heated in dioxane in the presence of hydrochloric acid, the aminothiazole ring closes to form 2-amino-5-(2,5-dihydroxy-3,4,6-trichlorophenyl)thiazoles (IIa-e). Oxidation of the latter with ferric chloride in aqueous DMFA at room temperature gives the quinones IIIa-e.

I a $R = CH_3$; b $R = C_2H_5$; II, III a $R^1 = H$; b $R^1 = CH_3$; c $R^1 = C_6H_5$; d $R^1 = CH_2CH = CH_2$; $e R^i = COCH_3$

Each step of the synthesis takes 2 h, and the yield of III (calculated on I) is 40-60%. The compounds are intensely colored (IIIa-d, blue; IIIe, violet), poorly soluble in orgnic solvents (except DMFA, DMSO, HMTP), and decompose when heated above 250°.

The structure of III was confirmed by the IR and UV spectra. IR spectrum (in Nujol), 1683-1653 (quinone C=0), 1645-1620 (thiazole ring), 1548-1532 (C=C), 3400-3200 cm⁻¹ (N-H). UV spectrum of IIIa (in ethanol), λ_{max} (log ϵ): 318 (4.14), 597 nm (3.67). Elemental composition data of the compounds were satisfactory.

LITERATURE CITED

1. D. Buckley, H. B. Henbest, and Pl Slade, J. Chem. Soc., No. 12, 4891 (1957).

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