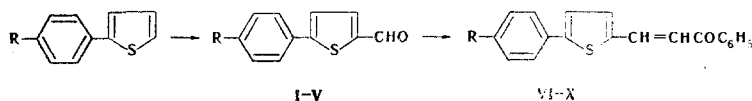


CARBONYL DERIVATIVES OF 2-ARYLTHIOPHENES

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The reaction of 4-R-benzenediazonium chlorides (R = H, CH₃, CH₃O, Cl, Br) with thiophene (Gomberg reaction) has given 2-arylthiophenes, which have been converted by Vilsmeier formylation into 5-arylthiophene-2-carbaldehydes (Table 1), as described by Demerseman et al. [1]. The latter, on condensation (methanolic solution of caustic soda, 45-55°C) with acetophenone has given chalcone analogs (Table 2):



Previously, 3-(5-phenyl-2-thienyl)-1-phenylprop-2-en-1-one (VI) has been described [2] with a yield of 51% and mp 110-112°C. The structure of the compounds obtained has been confirmed by UV and IR spectroscopy. The electronic spectra showed that the thiophene ring is a better transmitter of electronic influences than a 1,4-phenylene system.

TABLE 1. 5-Arylthiophene-2-carbaldehydes (I-V)

Compound	R	Mp, °C	Empirical formula	S, %			Yield, %	λ _{max} , nm (log ε) (in ethanol)	
				found	calculated	found		calculated	
I	H	90	C ₁₁ H ₈ OS	—	—	—	87	231 (3,96)	332 (4,33)
II	CH ₃	93	C ₁₂ H ₁₀ OS	15,9	16,3	15,8	83	235 (4,00)	335 (4,37)
III	CH ₃ O	116	C ₁₂ H ₁₀ O ₂ S	14,9	15,0	14,7	85	241 (4,00)	350 (4,30)
IV	Cl	87	C ₁₂ H ₁₀ ClOS	14,1	14,2	14,4	81	235 (4,01)	331 (4,34)
V	Br	114	C ₁₂ H ₁₀ BrOS	11,8	12,1	12,0	80	237 (4,00)	331 (4,38)

TABLE 2. 3-(5-Aryl-2-thienyl)-1-phenylprop-2-en-1-ones (VI-X)

Compound	R	Mp, °C	Empirical formula	S, %			Yield, %	λ _{max} , nm (log ε) (in ethanol)	
				found	calculated	found		calculated	
VI	H	122	C ₁₉ H ₁₄ OS	—	—	—	98	270 (4,12)	378 (4,48)
VII	CH ₃	132	C ₂₀ H ₁₆ OS	10,7	10,6	10,5	97	273 (4,14)	387 (4,47)
VIII	CH ₃ O	149	C ₂₀ H ₁₆ O ₂ S	10,2	10,3	10,0	98	282 (4,16)	397 (4,43)
IX	Cl	153	C ₁₉ H ₁₄ ClOS	10,0	10,1	9,9	99	272 (4,13)	382 (4,50)
X	Br	166	C ₁₉ H ₁₄ BrOS	8,7	8,8	8,7	98	268 (4,52)	381 (4,65)

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