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Synthesis of Nitrogen Functional Derivatives of 5-substituted-6-azauracil as one of the Four Nucleobases in the Nucleic Acid of RNA

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Abstract

3-Arylhydrazono-2,4-dioxo-4-phenylbutanoateshave been prepared by the coupling of benzoylpyruvate with aryldiazonium chlorides.Reactions of the 3-arylhydrazono-2,4-dioxo-4-phenylbutanoateswith 1-aminoguanidine, semicarbazide, and thiosemicarbazidegave5-substituted 2-imino-6-azauracil (**3a**),6-azauracil (**3b**), and 2-thio-6-azauracil (**3c**), respectively. The analytical data of these compounds - IR, 1H, and ¹³C NMR spectral data - are reported.

Key words: Coupling; Benzoylpyruvate; Arylhydrazone; 1, 2, 4-Triazin-5-one, 6-Azauracil.

Introduction

4-Aryl-2,4-dioxobutyrates are the formal derivatives of pyruvic acid and are as such trivially referred to as benzoylpyruvates. Being endowed with multiple functionalities, they are important synthetic precursors, capable of interacting with both electrophilic as well as nucleophilic reagents.In particular, in the latter case they offer a versatile scaffold on which to mould annulated rings carrying distinct structural features. This versatility draws its impetus from the conspicuous qualitative differences between the three carbonyl functionalities, which makes regioand chemoselective discrimination possible. Through a judicious matching of the applied nucleophilic species, the mode of annulation may be predicted, rendering a powerful tool for the construction of a variety ofheterocyclic compounds [1].Reactions of benzoylpyruvates with *N*,*N*- and *N*,*O*-dinucleophiles carrying a two-unit linker lead to formation of heterocyclic compounds [1-10]

It is know that benzoylpyruvate react with aryldiazonium salts to form the corresponding 3-arylhydrazono-2,4-dioxo-4phenylbutanoates [11].

3-arylhydrazono-2,4-dioxo-4-

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