

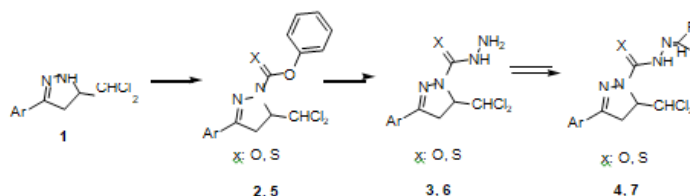
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Synthesis and biological evaluation of novel 2-pyrazoline derivatives

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Owing to the wide variety of biological activities of heterocyclic compounds comprising a 2-pyrazoline moiety, the development of new preparative methods for this class of compounds has become a subject of great interest. We previously described the first synthesis of 3-aryl-5-dichloromethyl-2-pyrazolines **1** involving chloral and several chloral derivatives. Pyrazolines **1** were able to react with phenyl chloroformate to give the corresponding phenyl pyrazolinecarboxylates **2**. These compounds were used as intermediates to obtain carbohydrazides **3**, which were used in preparing carbohydrazones **4**. A similar reaction sequence but using O-phenyl chlorothioformate instead of phenyl chloroformate provided good approaches to the respective intermediates **5**, carbothiohydrazides **6** and carbothiohydrazones **7**. The novel substances **3,4,6,7** were assayed to test their antitumoral properties against two leukemia cell lines, resulting IC₅₀ values between: 4.38-20.01 μ M (HL-60) and 2.19-17.05 μ M (K562).



Biography

María Vera Tárraga has completed her BSc in Chemistry and MSc in Organic Chemistry from Murcia University (Spain). She is currently a PhD student working on the synthesis of bioactive heterocyclic compounds.

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