. UDC 547.779.07:542.953.1

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It has been shown that mixtures of stereoisomeric dimethyl- β -thiodiacrylic esters are formed in the reaction of 1-substituted thiosemicarbazides with methyl propiolate [1]. The reaction of terminal α -acetylenic ketones with 1-phenylthiosemicarbazide in alcohol at 20°C leads to 2-acylmethyl-5-imino-3-phenyl-4H-1,3,4-thiadiazoles [2].

We have found that 1,3,5-triphenylpyrazole, with mp 137-138°C (from alcohol) (mp 137-138°C [3]), is formed in 68% yield when equimolar amounts of 1-benzoyl-2-phenylacetylene and 1-phenylthiosemicarbazide are refluxed in methanol for 5 h.

The intermediately formed 1-benzoylvinylthiosemicarbazide is unstable and upon heating splits out a molecule of HNCS to give 1-benzoylvinyl-1-phenylhydrazine. Under the reaction conditions

the latter readily undergoes dehydrocyclization to 1,3,5-triphenylpyrazole. IR spectrum: 700, 770, 980, 1070 (=CH deformation vibrations); 1420, 1442, 1465, 1505 (pyrazole ring C=N and C=C); 1600, 1620 (aromatic ring C=C); 3070 cm⁻¹ (=CH stretching vibrations). The results of elementary analysis were in agreement with the calculated values.

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Irkutsk Institute of Organic Chemistry, Siberian Branch, Academy of Sciences of the USSR, Irkutsk 664033. Translated from Khimiya Geterotsiklicheskikh Soedinenii, No. 8, pp. 1136-1137, August, 1985. Original article submitted February 18, 1985.