

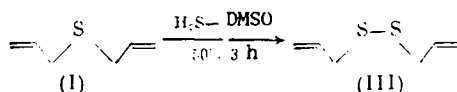
FORMATION OF DIALLYL DISULFIDE FROM DIALLYL SULFIDE IN THE H₂S-DMSO SYSTEM

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UDC 542.97:547.379.3

Diallyl sulfide (I) reacts with the H₂S DMSO LiOH system to form 3,7-dimethyl-1,2,5-trithiacycloheptane and 4-thia-1-heptene-6-thiol [1]. Di(propen-1-yl) sulfide (II) reacts with this system to give dipropyl polysulfides [2].

We have found that sulfide (I) reacts with the H₂S-DMSO system to give diallyl sulfide (III) in 76.3% yield. The conversion of sulfide (I) was 58.8%.



Sulfide (II) does not undergo this reaction. The yield of disulfide (III) in the reaction of sulfide (I) with the S₈-DMSO system at 50°C over 3 h was 8.2% and the conversion of sulfide (I) was 16.3%.

A sample of 5.1 g sulfide (I) and 200 ml DMSO was heated at 50°C and hydrogen sulfide was introduced for 3 h. The mixture was cooled to room temperature, diluted with water, and extracted with ether. The ethereal extracts were washed with water and dried over CaCl₂. Ether was distilled off. The residue was fractionated in vacuum to give 2.9 g (76.3%) disulfide (III), 2.1 g (I), and ~1 g tarry residue.

PMR spectrum of disulfide (III) in CDCl₃ with HMDS as the internal standard (δ, ppm): 3.32 d (2H, CH₂), 5.07 m (2H, =CH₂), 5.83 m (1H, =CH-).

The physical constants of disulfide (III) were in accord with reported data [3]. The elemental analysis corresponded to the proposed formula. Mass spectrum: M⁺ 146 m/z.

LITERATURE CITED

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