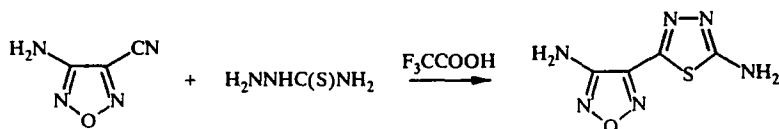


SYNTHESIS OF 4-(5-AMINO-1,3,4-THIADIAZOL-2-YL)-1,2,5-OXADIAZOLE-3-AMINE

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To expand the group of 1,3,4-thiadiazole derivatives which could be of interest as efficient stabilizers of silver halide photographic emulsions and the components of light-sensitive and photoconductor materials and drugs with antimicrobial and antiviral activity, we synthesized the first member of a series of 1,3,4-thiadiazoles containing an amino-1,2,5-oxadiazole fragment — 4-(5-amino-1,3,4-thiadiazol-2-yl)-1,2,5-oxadiazole-3-amine. It was formed with a yield of more than 75% in condensation of 3-amino-4-cyano-1,2,5-oxadiazole with thiosemicarbazide in trifluoroacetic acid.



4-(5-Amino-1,3,4-thiadiazol-2-yl)-1,2,5-oxadiazole-3-amine. A mixture of 6.1 g (5.5 mmole) of 3-amino-4-cyano-1,2,5-oxadiazole, 5.5 g (6.0 mmole) of thiosemicarbazide, and 20 ml of trifluoroacetic acid was heated for 4 h in a boiling water bath while stirring, cooled, and poured into 50 ml of 25% ammonia solution. The lightly colored sediment was filtered off, washed with water (to neutral reaction of the washing water), and dried in air. Yield of 7.6 g (75%); mp = 236-237°C. IR spectrum: 3456, 3368 (NH₂), 1688, 1648 (NH), 1512 (C-N, thiadiazole), 1468 (C-N, furazan), 1088 cm⁻¹ (N-O, furazan). PMR spectrum (DMSO): 7.95 (2H, s, NH₂, thiadiazole) and 6.54 ppm (2H, s, NH₂, oxadiazole). ¹³C NMR spectrum (DMSO): 169.978 (C₅, thiadiazole) and 143.56 (C₂, thiadiazole); 154.389 (C₃, oxadiazole), and 139.818 ppm (C₄, oxadiazole). Found, %: C 25.9; H 2.4; N 45.5; S 17.8. C₄H₄N₆OS. Calculated, %: C 26.0; H 2.2; N 45.5; S 17.4.

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