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THE PREPARATION OF 1, 5-Se₂S₂N₄

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The reaction of [(Me₃Si)₂N]₂S with an equimolar amount of SeCl₄ or the reaction of [(Me₃Si)₂N]₂Se with 1:1 mixture of SCl₂ and SO₂Cl₂ produces 1,5-Se₂S₂N₄ in good yields. The product is identified and characterized using X-Ray diffraction, vibrational analysis, mass spectroscopy, and NMR-spectroscopy.

<u>Keywords:</u> Chalcogen-nitrogen-compounds; Raman spectroscopy; NMR spectroscopy; crystal structure

INTRODUCTION

The chemistry of selenium-nitrogen compounds is a relatively limited but a rapidly growing field. [1,2] We report here a convenient synthesis of 1,5-Se₂S₂N₄ from [(Me₃Si)₂N]₂S and SeCl₄, or from [(Me₃Si)₂N]₂Se and SCl₂ and SO₂Cl₂. Both reactions afford the same product that is identified using X-ray diffraction, vibrational analysis, as well as NMR and mass spectroscopy.

EXPERIMENTAL SECTION

All reactions were carried under an argon atmosphere. The solvents were dried by freshly distilling under a nitrogen atmosphere.

Caution! The eight-membered Se₂S₂N₄ ring molecule is explosive when heated or subjected to mechanical stress.

Reaction of [(Me₃Si)₂N]₂S with SeCl₄

A solution of [(Me₃Si)₂N]₂S in CS₂ was added dropwise to a slurry of SeCl₄ in CS₂ at -78° C. The reaction mixture was allowed to warm slowly to room temperature with stirring for 12 hours. The yield was 70 mol % based on the initial amount of [(Me₃Si)₂N]₂S.

Reaction of [(Me₃Si)₂N]₂Se with SCl₂ and SO₂Cl₂

[(Me₃Si)₂N]₂Se was dissolved in CS₂, and a mixture of SCl₂ and SO₂Cl₂ in CS₂ was added dropwise at -78° C. The yield was 73 mol % based on the initial amount of [(Me₃Si)₂N]₂Se.

RESULTS AND DISCUSSION

Se₂S₂N₄ can be prepared in good yields according to the following reactions.

$$[(Me_3Si)_2N]_2S + SeCl_4 \rightarrow \frac{1}{2} Se_2S_2N_4 + 4 Me_3SiCl$$

$$[(Me_3Si)_2N]_2Se + SCl_2 + SO_2Cl_2 \rightarrow \frac{1}{2} Se_2S_2N_4 + 4 Me_3SiCl$$

The resulting dark brown-red material was almost insoluble in organic solvents and precipitated during the reaction. The elemental analysis of

the solid product can be inferred in terms of a mixture containing 91 mol % of Se₂S₂N₄ and 9 mol % of Se₈ (Anal. calcd for Se₂S₂N₄: N, 20.1; S, 23.0; Se 56.9. Found: N, 16.5. The ⁷⁷Se NMR spectrum of the product indicates a mixture of 90-95 % Se₂S₂N₄ and 5-10 % Se₈).

Se₂S₂N₄ crystallizes in a monoclinic space group $P2_1/c$, Z=4, with the unit cell dimensions a=8.818(2), b=7.387(1), c=8.981(2) Å, $\beta=93.14(3)$ ° (T = 298 K); R=0.0545. The compound is isostructural with S₄N₄ ^[3] and β -Se₄N₄. ^[4] The structure is disordered with sulfur and selenium statistically distributed over chalcogen atom sites (site occupation factors of selenium in every position ca. 50 %).

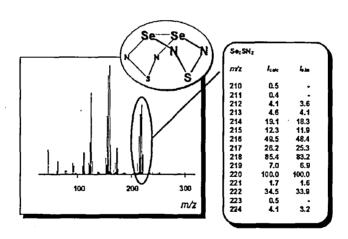


FIGURE 1 12 eV EI mass spectrum of 1,5-Se₂S₂N₄.

The 12 eV mass spectrum of the product showed Se_2SN_2 as the largest fragment. It can be inferred as a part of the $1,5-Se_2S_2N_4$ ring (see Fig. 1). The ¹⁴N NMR spectrum of the crude reaction mixture showed two resonances at -46 and -238 ppm. The resonance at -238 ppm is assigned to $Se_2S_2N_4$ (c.f. -256 ppm for S_4N_4 ^[5]). The resonance

at –46 ppm is due to an unknown species that is removed during the purification of the product by distillation. The ⁷⁷Se NMR spectrum showed two resonances at 1418 ppm and 620 ppm that are assigned to 1,5-Se₂S₂N₄ and Se₈, ^[6] respectively. The single ¹⁴N resonance also indicates that the product is the 1,5-isomer.

The observed Raman lines are in a good agreement with the fundamental vibrations calculated for Se₂S₂N₄ by using the general valence force field approach. The calculations also yield reasonable force constants.

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