

**SHORT  
COMMUNICATIONS**

## Reaction of Divinyl Sulfide with Selenourea

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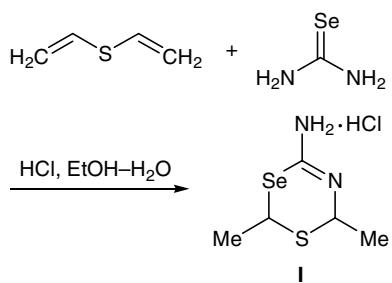
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Divinyl sulfide is a versatile intermediate product in modern organic synthesis [1–4], which is used to obtain various heterocyclic compounds [2–4]. Divinyl sulfide is known to react with thiourea and its derivatives in the presence of strong acids to produce 4-amino-2,6-dimethyl-2*H,6H*-1,3,5-dithiazinium salts [5–9]. There are no published data on reaction of divinyl sulfide with selenourea.

We have found that divinyl sulfide reacts with selenourea in the presence of hydrochloric acid, yielding 90% of previously unknown heterocyclic compound, 2,6-dimethyl-2*H,6H*-1,3,5-thiaselenazin-4-amine hydrochloride (**I**). The reaction occurs in aqueous ethanol at 50°C. The structure of compound **I** was proved by the <sup>1</sup>H, <sup>13</sup>C, and <sup>77</sup>Se NMR and IR spectra and elemental analysis. The product is an approximately equimolar mixture of *cis* and *trans* isomers.



**2,6-Dimethyl-2*H,6H*-1,3,5-thiaselenazin-4-amine hydrochloride (**I**).** mp 113–114°C. IR spectrum (KBr),  $\nu$ , cm<sup>-1</sup>: 670, 770, 960, 995, 1150, 1300, 1345, 1375, 1430, 1470, 1590, 1620, 3050, 3190, 3400. <sup>1</sup>H NMR spectrum (CDCl<sub>3</sub>),  $\delta$ , ppm: *cis* isomer: 1.53 d (6-CH<sub>3</sub>), 1.56 d (2-CH<sub>3</sub>), 4.95 q (6-H), 5.20 q (2-H),

11.40 s (NH<sub>2</sub>); *trans* isomer: 1.70 d (6-CH<sub>3</sub>), 1.89 d (2-CH<sub>3</sub>), 4.97 q (2-H), 5.27 q (6-H), 9.44 s (NH<sub>2</sub>). <sup>13</sup>C NMR spectrum (CDCl<sub>3</sub>),  $\delta$ <sub>C</sub>, ppm: *cis* isomer: 19.58, 20.06, 36.48, 51.40, 169.80; *trans* isomer: 20.63, 26.41, 39.90, 57.08, 169.72. <sup>77</sup>Se NMR spectrum (CDCl<sub>3</sub>),  $\delta$ <sub>Se</sub>, ppm: 470. Found, %: C 24.10; H 4.52; Cl 14.61; N 11.11; S 13.39; Se 32.42. C<sub>5</sub>H<sub>11</sub>ClN<sub>2</sub>SSe. Calculated, %: C 24.45; H 4.51; Cl 14.45; N 11.40; S 13.03; Se 32.16.

### REFERENCES

1. Trofimov, B.A., Amosova, S.V., Gusarova, N.K., and Musorin, G.K., *Tetrahedron*, 1982, vol. 38, p. 713.
2. Trofimov, B.A. and Amosova, S.V., *Sulfur Rep.*, 1984, vol. 3, p. 323.
3. Trofimov, B.A. and Amosova, S.V., *Divinilsulfid i ego proizvodnye* (Divinyl Sulfide and Its Derivatives), Novosibirsk: Nauka, 1983, p. 135.
4. Amosova, S.V. and Gavrilova, G.M., *Heteroatom Chem.*, 2006, vol. 17, p. 491.
5. Trofimov, B.A., Gavrilova, G.M., Kalabin, G.A., Bairov, V.V., and Amosova, S.V., *Khim. Geterotsikl. Soedin.*, 1979, p. 1466.
6. Gavrilova, G.M., Amosova, S.V., Trofimov, B.A., Kositsina, E.I., Pertsikov, B.Z., Gostevskaya, V.I., Musorin, G.K., Al'pert, M.L., and Borodina, N.M., *Khim. Geterotsikl. Soedin.*, 1986, p. 697.
7. Trofimov, B.A., Gavrilova, G.M., Voronov, V.K., and Amosova, S.V., *Khim. Geterotsikl. Soedin.*, 1980, p. 1136.
8. Gavrilova, G.M., Kositsina, E.I., Trofimov, B.A., Gostevskaya, V.I., and Amosova, S.V., *Khim. Geterotsikl. Soedin.*, 1982, p. 622.
9. Gavrilova, G.M., Trofimov, B.A., Gostevskaya, V.I., and Amosova, S.V., *Khim. Geterotsikl. Soedin.*, 1982, p. 652.