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PYRIDINETHIONES PREPARATION AND CHEMISTRY OF 1-SUBSTITUTED-2-FORMYL-2(1H)-PYRIDINETHIONES AND -SELONES

Jan Becher^a

^a Department of Chemistry, Odense University, Odense M, Denmark

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PYRIDINETHIONES

PREPARATION AND CHEMISTRY OF 1-SUBSTITUTED-2-FORMYL-2(1H)-PYRI-DINETHIONES AND -SELONES.

Jan Becher

Department of Chemistry, Odense University, DK-5230 Odense M, Denmark.

Ring closure reactions of glutacondialdehyde derivatives have been reported. We have reported the synthesis of 1-substituted-3-formy1-2(1H)-pyridinethiones $\underline{2}$ from the glutacondialdehyde anion 1 and isothiocyanates:

R	yield	m.p.	R	yield	m.p.
2a methy1			2d pheny1		180-2 ⁰
<pre>2b cyclohexy1</pre>	98%	152-4°	2e 2,6-dimethylphenyl	79%	190-2°
2c tert-buty1	38%	86-7 ⁰	2f 2-naphthy1	94%	195-6 ⁰

Scheme 1

Due to the low solubility of the glutacondial dehyde anion (sodium or potassium salt) in organic solvents of low polarity, the reactions were run in DMF or DMSO. In the aromatic series the exothermic reactions took place at room temperature, whereas an elevated temperature (ca. 80° C) was necessary to complete the reactions in the aliphatic series.

Attempts to prepare 3-formy1-2(1H)-pyridinethione, $\underline{3}$, from $\underline{1}$ and reagents such as (SCN)₂ and HSCN were unsuccessful. However, $\underline{3}$ was isolated in quantitative yield by thermolyses (ca. 190°C/

latm.) of $\underline{2}c$. Thermal syn-elimination of alkenes from corresponding structural arrangements are well known ($\underline{e}.\underline{g}$. Chugaev reaction).

The poster will discuss the chemistry of $\underline{3}$ and the preparation of a number of related systems.

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