SYNTHESIS OF NEW 2-OXO-1,2-DIHYDROPYRIDINE-3-CARBOXYLIC ACID DERIVATIVES

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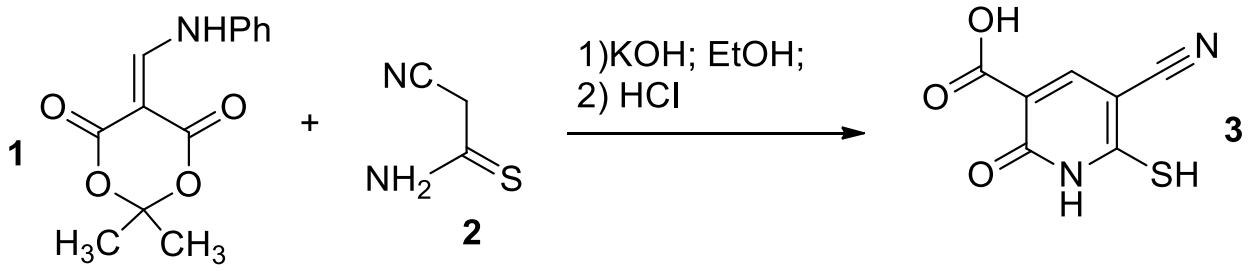
Abstract

2,2-Dimethyl-5-((phenylamino)methylene)-1,3-dioxane-4,6-dione prepared by reaction of Meldrum's acid with triethyl orthoformate and aniline, reacts with active methylene nitriles to afford 2-oxo-1,2-dihydropyridine-3-carboxylic acid derivatives useful as drug precursors or perspective ligands.

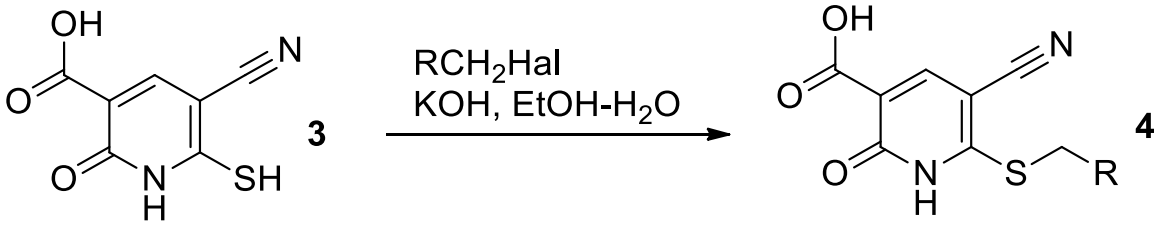
Keywords: nicotinic acids, Meldrum's acid, 2-oxo-1,2-dihydropyridine-3-carboxylic acid, cyanothioacetamide, cyanoacetamide.

It is known that nicotinic acid (niacin, vitamin PP) and its derivatives have a wide spectrum of biological activity. Thus, nicotinic acid and nicotinates showed hypolipidemic, hypocholesterolemic, neuroprotective and other effects. 2-Oxo-1,2-dihydropyridine-3-carboxylic acid are less studied. However, they are of interest as complexating agents [1,2] and as pharmaceuticals [3].

Earlier we have developed [4] the method for synthesis of 6-mercapto-2-oxonicotinic acid **3** based on the heterocyclization of aminomethylidene derivative of Meldrum's acid **1** with cyanothioacetamide **3**.

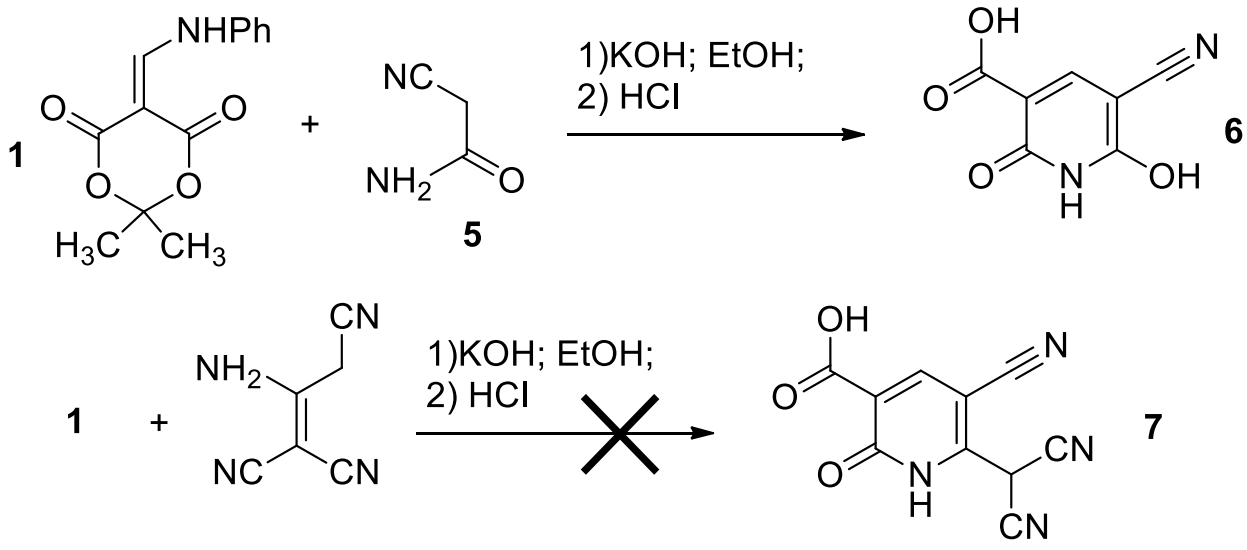


We decided to study the reactions of 5-cyano-2-oxo-1,2-dihydropyridine-3-carboxylic acid **3**. Compound **3** easily reacts with alkyl halides regioselectively at S atom to give sulfides **4**.



Other active methylene nitriles were also introduced in the reaction. Thus, cyanoacetamide **5** in the presence of KOH reacts with

enamino-1,3-diketone **1** following by acidification to give pyridine **6** in a good yield. However, we failed to prepare the compound **7** starting from maljnonitrile dimer.



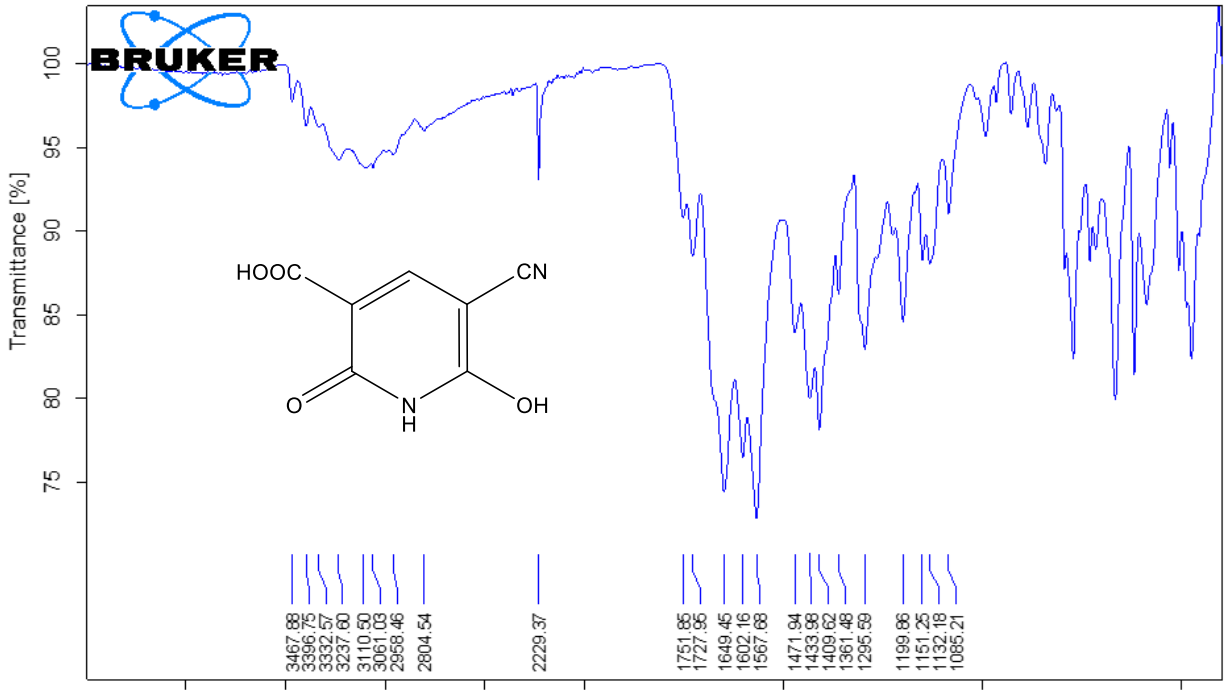


Figure 1. IR spectrum of compound **6**

Experimental

Anilinomethylidene derivative of Meldrum's acid. A mixture of the powdered Meldrum's acid (0.1 mol), triethyl orthoformate (21.6 mL, 0.13 mol), and freshly distilled aniline (9.1 mL, 0.1 mol) was refluxed with vigorous stirring for 5 min to afford a syrupy reaction mass. It was diluted with 30 mL of EtOH and refluxed for an additional 3 min. Then it was cooled with stirring to ~20 °C and diluted with water to 100 mL. After 2 h, the product was filtered off and washed with water, twice with 60% EtOH, and with hexane.

2,2-Dimethyl-5-(phenylamino)methylene-1,3-dioxane-4,6-dione (1). Yield 92%, m.p. 156—157 °C. Found (%): C, 63.19; H, 5.32; N, 5.66. C13H13NO4. Calculated (%): C, 63.15; H, 5.30; N, 5.67. ^{1}H NMR, δ: 1.70 (s, 6 H, 2 Me); 7.19-7.51 (m, 5 H, Ph); 8.58 (d, 2 H, CH=, ^{3}J = 14.7 Hz); 11.27 (d, 1 H, NH, ^{3}J = 14.7 Hz).

Compounds 3 and 6 (general procedure). Potassium hydroxide (1.12 g, 0.02 mol) was added to a vigorously stirred suspension of compound **1** (0.01 mol) and cyano(thio)acetamide (0.01 mol) in 10 mL of EtOH. After 24 h, the reaction mixture was acidified with conc. HCl to pH 5 and kept for 3 h. The precipitate that formed was filtered off and washed successively with water and EtOH. The yield of pyridine **3** was 68% and pyridine **6** – 74%.

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