Molecules 1999, 4, M111

## 4,5-Dimethoxy-2-nitroacetophenone

## Dmitrij S. Zavgorodniy, Alexander V. Butin and Tat'yana A. Stroganova

Research Laboratory of Furan Chemistry, Kuban State Technological University, Moskovskaya 2, Krasnodar, Russian Federation. Phone: +7 8612 55 95 56; E-mail: <a href="mailto:nemol@kubstu.ru">nemol@kubstu.ru</a>, E-mail: <a href="mailto:strog@kuban.net">strog@kuban.net</a>

Received: 13 September 1999 / Accepted: 20 September 1999 / Published: 8 October 1999

The general part of the experimental section [1] has been presented elsewhere. To solution of 4,5-dimethoxyacetophenone (20.0 g, 110 mmol) in 45 ml of glacial acetic acid, red fuming nitric acid (22.0 ml) was added dropwise with cooling in an ice-water bath. After 20 minutes the reaction mixture was poured into water. Increasing the reaction time caused by-product formation. The precipitate was filtered off, washed with water and recrystallized from ethanol to yield 14.3 g (58 %) 4,5-dimethoxy-2-nitroacetophenone.

M.p. 134°C (ethanol).

<sup>1</sup>H NMR (CDCl<sub>3</sub>, 80 MHz): 7.60 (s, 1H, 3-H<sub>Ar</sub>); 6.75 (s, 1H, 6-H<sub>Ar</sub>); 3.93 (s, 6H, OCH<sub>3</sub>); 2.42 (s, 3H, CH<sub>3</sub>).

IR (cm<sup>-1</sup>): 1680 (C=O).

Anal. calc. for C<sub>10</sub>H<sub>11</sub>NO<sub>5</sub> (225.21): C 53.33, H 4.92; Found: C 53.17, H 5.09.

## Reference

1. Gutnov, A.V.; Butin, A.V.; Abaev, V.T.; Krapivin, G.D.; Zavodnik, V.E. Furyl(aryl)alkanes and Their Derivatives.19. Synthesis of Benzofuran Derivatives via 2-Hydroxyaryl-R-(5-methylfur-2-yl)methanes. Reaction of Furan Ring Opening - Benzofuran Ring. *Molecules* **1999**, *4*, 204-218.

Sample availability: available from the authors and from MDPI.

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