

Prebiotic Formation of Sugars and Amino Acids Using Aminomethylene

Project Manager
Dr. Artur Mardyukov

Researchers
Felix Keul

Principal Investigator
Prof. Dr. Peter R. Schreiner

Project Term
2020 - 2020

Clusters
Goethe-HLR Frankfurt

Software
GAUSSIAN

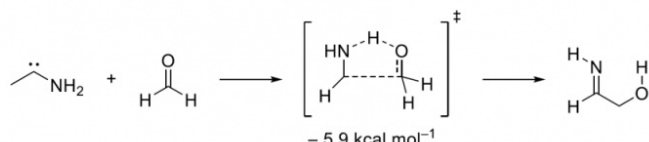
Additional Software
CFOUR

Institute
Institute of Organic Chemistry

University
Justus Liebig University Giessen

Funded by

Volkswagenstiftung



Scheme 1: Reaction between aminomethylene and formaldehyde towards glycolonitrile.

Introduction

After the matrix isolation of aminomethylene (H-C-NH_2)^[1] we study further the formation of amino acids and sugars in quantitative flow pyrolysis experiments with cyclopropylamine as starting material. After pyrolysis at 900 °C aminomethylene is generated by ethylene extrusion and undergoes [1,2] H-shift to methanimine. During the pyrolysis a huge amount of HCN is released by the decomposition of aminomethylene or methanimine which reacts with methanimine to aminoacetonitrile during warm-up of the cold trap. Hydrolysis with *conc.* HCl should lead to the amino acid glycine according to Strecker's synthesis. By rinsing the cold trap with water imine can also be hydrolyzed to formaldehyde which immediately forms glycolonitrile with HCN in solution. Computations are used as a helping tool to investigate these reactions.

Methods

For computational studies, the program gaussian16 is used. For geometry preoptimizations we use hybrid functionals as B3LYP with high basis sets. For accurate computations we used coupled cluster methods.

Results

We were able to identify aminoacetonitrile and glycolonitrile via NMR spectroscopy and GC-MS in the condensed residue of flow pyrolysis experiments which confirm our working hypothesis. We were not able to identify formaldehyde in the spectra suggesting that the subsequent reaction with HCN is very fast. The high level ab initio computations indicate that the reaction is almost barrierless as for hydroxymethylene and formaldehyde. Formaldehyde and aminomethylene form a pre-reactive complex of $-6.3 \text{ kcal mol}^{-1}$. Over a small barrier of $0.4 \text{ kcal mol}^{-1}$ that corresponds to a rotation of formaldehyde in the plane of aminomethylene it undergoes to iminoethanol. The C-H-insertion

goes over an unfavored barrier of $21.3 \text{ kcal mol}^{-1}$ to aminoacetaldehyde.

Outlook

Based on these results, we continue the experiments that we can show the simultaneous formation of sugars and amino acids. Hereby, aminomethylene can further react in a barrierless carbonyl-ene reaction with formaldehyde to give iminoethanol, which then can be further converted to serinonitrile or be hydrolysed to glycolaldehyde.

Reference

[1] Eckhardt, A. K.; Schreiner, P. R.: *Angew. Chem. Int. Ed.*, 2018, 57, 5248-5252. <https://doi.org/10.1002/anie.201800679>

Last Update: 2022-04-22 12:48