

# Quantification of urea electrolysis products in alkaline medium

Christine Cachet-vivier, R. Benyahia, Lamia Rebiai, Laure Latapie, Guillaume Hopsort, Karine Groenen Serrano, Karine Loubière, Theodore Tzedakis, Sam Azimi, Vincent Rocher, et al.

### ▶ To cite this version:

Christine Cachet-vivier, R. Benyahia, Lamia Rebiai, Laure Latapie, Guillaume Hopsort, et al.. Quantification of urea electrolysis products in alkaline medium. 72nd Annual Meeting of the International Society of Electrochemistry, Aug 2021, Jeju Island, South Korea. hal-04796324

## HAL Id: hal-04796324 https://hal.science/hal-04796324v1

Submitted on 21 Nov 2024

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

#### Quantification of urea electrolysis products in alkaline medium

<u>C. Cachet-Vivier</u><sup>(a)</sup>, R. Benyahia<sup>(a)</sup>, L. Rebiai<sup>(a)</sup>, L. Latapie<sup>(b)</sup>, G. Hopsort<sup>(b)</sup>, K. Serrano<sup>(b)</sup>, K. Loubière<sup>(b)</sup>, T. Tzedakis<sup>(b)</sup>, S. Azimi<sup>(c)</sup>, V. Rocher<sup>(c)</sup>, E. Torralba-Penalver<sup>(a)</sup>, S. Bastide<sup>(a)</sup>

<sup>(a)</sup> Université Paris Est, ICMPE (UMR 7182), CNRS, UPEC, F-94320 Thiais, Fance

<sup>(b)</sup> Laboratoire de Génie Chimique (UMR 5503), INP, CNRS, Université Paul Sabatier, 118, route de Narbonne, 31062 Toulouse Cedex 9, France

<sup>(c)</sup> Département Innovation du Syndicat Interdépartemental pour l'Assainissement de l'Agglomération Parisienne (SIAAP), Colombes, France

#### mail: cachet@icmpe.cnrs.fr

The treatment of nitrogen compounds of sewage waters in wastewater treatment plants (WWTPs) is very costly in energy and has an impact on the environment with discharges of nitrates into rivers and N<sub>2</sub>O (greenhouse gas) in the atmosphere. WWTPs operators are therefore looking for innovative treatment concepts that enhance the value of both mater and energy. Because nitrogen derives mainly from urea, which is contained in large quantities in urine (0.33 mol L<sup>-1</sup>), an electrochemical remediation of urine can be foreseen to alleviate the problems of its treatment in WWTPs.

The objective of our works is the implementation of an electrochemical system allowing the degradation of urea (pollutant) at the anode and the production of hydrogen (green fuel) at the cathode, according to the overall process:

 $CO(NH_2)_2 + H_2O \rightarrow CO_2 + N_2 + 3 H_2$ 

In recent years, many works have concerned the evaluation of the electrocatalytic properties for urea electro-oxidation by voltammetry using Ni-based catalysts, because of the Ni ability in this matter [1-2]. Two main axes have followed: (i) the nanostructuring of the electrode material to increase the active surface area in various forms (nanowires, nanosheets, nanoribbons, nanoparticles) or (ii) the addition of a co-element to nickel that mainly modifies the oxidation potential of urea. For our part, we have studied from an analytical point of view composites made of Ni-Rh nanoparticles embedded in carbon matrix that combines these two aspects [3].

Concerning the electrolysis process, the studies generally focus on the proportions of  $N_2$  and  $O_2$  produced in the gas phase at the anode and the percentage of  $H_2$  formed at the cathode. However, to the best of our knowledge, none of these works report mass balances performed on the various soluble intermediates arising from secondary reactions that can be obtained, nor their faradic yields.

This led us to proceed with electrolysis to identify and quantify both gaseous and solution species produced. For this purpose, we carried out electrolysis on a simple nickel plate in alkaline solutions as well as in alkalinized synthetic urine and quantified the reactions products by different analytical means (Gas chromatography, UV-Vis spectrometry, ionic chromatography).

Our results with simple urea solutions show that at the cathode,  $H_2$  is collected with a yield of 100% (*i. e.* hydrogen is the only reaction). At the anode, nitrites are detected as a by-product with levels depending on the applied potential. This is the first time that this result is reported in the literature. The problem with the obtention of nitrites is that nitrogen is not removed from the solution (as a gaseous product like  $N_2$ ). We will also present results obtained with synthetic urine and report on the improvement of  $N_2$  production rates through modification of the (Ni-based) electrode compositions.

The authors wish to thank the French National Research Agency for funding this research in the frame of the HYUREA project ANR19CE04-0009.

[1] E. Urbańczyk, M. Sowa, W. Simka, Urea removal from aqueous solutions—a review, J. Appl. Electrochem. 46 (2016) 1011–1029. <u>https://doi.org/10.1007/s10800-016-0993-6</u>.

[2] K. Ye, G. Wang, D. Cao, G. Wang, Recent Advances in the Electro-Oxidation of Urea for Direct Urea Fuel Cell and Urea Electrolysis, Top. Curr. Chem. 376 (2018) 42. <u>https://doi.org/10.1007/s41061-018-0219-y</u>.

[3] P. Mirzaei, S. Bastide, A. Dassy, R. Bensimon, J. Bourgon, A. Aghajani, C. Zlotea, D. Muller-Bouvet, C. Cachet-Vivier, Electrochemical oxidation of urea on nickel-rhodium nanoparticles/carbon composites, Electrochimica Acta. 297 (2019) 715–724. <u>https://doi.org/10.1016/j.electacta.2018.11.205</u>.