STABLE DEEP EUTECTIC SOLVENT DOPED WITH METAL NANOPARTICLES

1,2Anastasiia TULUPOVA, 1Vasilii BURTSEV, 1Vaclav ŠVORČÍK, 1,2Oleksiy LYUTAKOV

1VŠCHT - University of Chemistry and Technology, Prague, Czech Republic, EU, lyutakoo@vscht.cz
2Tomsk Polytechnic University, Research School of Chemistry and Applied Biomedical Sciences, Tomsk, Russian Federation

https://doi.org/10.37904/nanocon.2021.4340

Abstract

In this work, we prepared MeNPs doped DES systems. Gold nanoparticles were prepared on aluminum oxide microparticles surface. DES was prepared from choline chloride and urea, and embedded with the nano/micro particles. Created system was characterized using UV-Vis and FTIR spectroscopy, as well as TEM technique. As the result, we demonstrate that stable deep eutectic solvents that can be used for a range of applications.

Keywords: Deep eutectic solvents, doped, metal nanoparticles

1. INTRODUCTION

Metal nanoparticles (MeNPs) are of high interest since their unique optical, magnetic, electrical and physiological properties [1]. MeNPs can be used in different application areas, such as biomedicine [2], electrochemical catalysis [3], chemical sensing [4] and many others. MeNPs are commonly used as a colloidal suspension in water or traditional organic solvents. In turn, Deep Eutectic Solvents (DES) are recently proposed new alternative media for applications as a “green” reaction medium in organic synthesis. In contrast to traditional organic solvents, DES is non-toxic and non-flammable. Finally, deeply eutectic solvents do not produce toxic metabolites and are biodegradable [5]. The main uses of DES technology include the synthesis of nanostructures with a controlled shape [6], metal electrodeposition [7], CO2 adsorption [8] and others. But recently, attention has been paid to the study of DES as alternative solvents for organic synthesis and photochemistry. In this work we propose a creation of MeNPs doped DES for a range of potential applications.

2. EXPERIMENTAL

2.1. Materials

Gold(III) chloride trihydrate (HAuCl4, ≥99.9 %), sodium citrate tribasic dihydrate (≥99.5 %), choline chloride (≥98,0 %), urea (≥99,0 %), aluminum oxide, 4-aminothiophenol (≥97,0 %), p-Toluenesulfonic acid monohydrate (≥98,5 %), tert-Butyl nitrite (≥90,0 %), methanol, acetonitrile and deionized water were purchased from Sigma-Aldrich and used without further purification.

2.2. Samples preparation

Preparation of Deep eutectic solvent

At first stage, choline chloride is dried under vacuum at 90 °C and then mixed with powdered urea in a ratio of 1:2. The two components were heated with constant stirring until a homogenous mixture was obtained (around 5 hours).
Preparation of AuNPs

Gold nanoparticles on aluminum oxide microparticles surface was prepared according to following methodology: Al₂O₃ powder first was modified with functional group -SH using a diazonium salt. Modified 0.2 g Al₂O₃ was added to deionized water and boil. After the water boiled was added 10 mM aqueous HAuCl₄ solution to it, then 40 mM sodium citrate tribasic dihydrate water solution was added dropwise within 30-40 seconds. The resulting solution was boiled for 10 min, then was transferred to a cold hotplate and continue stirring for 12 hours. The resulting particles were washed by water and methanol by centrifugation at 7000 rpm for 5 min and dried in oven at 60 °C. Figure 1 schematically shows the preparation route Al₂O₃-AuNPs.

![Figure 1 Schematic representation of the preparation of gold nanoparticles on aluminum oxide microparticles surface](image)

2.3. Measurement Techniques

UV-Vis absorbance spectra of the samples were measured using a HR2000 (Ocean Optics) spectrometer in 200-800 nm wavelengths range using the AvaLight-DHS light source (Avantes).

Fourier transform infrared (FTIR) attenuated total reflection (ATR) spectra were recorded using a Nicolet 6700 spectrometer (Thermo Scientific, France) with a Smart ATR accessory device (1000 scans and 4 cm⁻¹ resolution).

Transmission electron microscopy (TEM) images were obtained with a JEOL JEM-1010 instrument (JEOL Ltd., Japan), with a SIS MegaView III digital camera (Soft Imaging Systems, acceleration voltage 80 kV) and analysis was performed by using AnalySIS Software 2.0.

3. RESULTS AND DISCUSSION

Deep eutectic solvent including choline chloride (ChCl) as the hydrogen bond acceptor urea as a hydrogen bond donor were prepared. The synthesized DES had the following physicochemical properties: a homogenous and transparent liquid of high viscosity, without any evident solid particles and stable at room temperature. To confirm the formation of hydrogen bonding, infrared spectra of choline chloride, urea and deep eutectic solvents were obtained. The results are presented in Figure 2. Absorption bands of urea at 3425 cm⁻¹ and 3333 cm⁻¹ correspond to the stretching mode of –NH₂. On the contrary, these absorption bands moved towards the lower values of wavenumber, and specifically 3402 cm⁻¹ and 3315 cm⁻¹. This shift could be affected by the formation of hydrogen bonds between urea and ChCl. Also, the bands associated to ChCl and urea
appeared in the spectrum of DES. Their corresponding wave number of the vibrational modes and associated terminal groups are listed in Table 1.

Figure 2 FTIR spectra of choline chloride, urea and synthesized DES

Table 1 Wave numbers and their assignments obtained from FTIR spectra.

| Observed frequencies (cm\(^{-1}\)) | Assignments       |
|------------------------------------|--------------------|
| Urea                              | Choline Chloride   | DES         |
| 3425                               | 3402               | \(v_{as} \text{ NH}_2\) |
| 3333                               | 3315               | \(v_s \text{ NH}_2\) |
| 3256                               |                    | \(v_{as} \text{ NH}_2\) |
| 3219                               | 3182               | \(v_{as} \text{ OH}\) |
| 3016                               | 3026               | \(\delta_{as} \text{ OH}\) |
| 1674                               | 1665               | \(\delta_{s} \text{ NH}_2\) |
| 1592                               | 1601               | \(\delta_{as} \text{ NH}_2\) |
| 1482                               | 1472               | \(\rho \text{ CH}_3\) |
| 1454                               | 1430               | \(\rho_s \text{ CH}_3\) |
| 1151                               | 1169               | \(v_{as} \text{ CN}\) |
| 1086                               | 1082               | \(\rho \text{ CH}_2\) |
| 1008                               | 1008               | \(v \text{ C-O}\) |
| 953                                | 953                | \(v_{as} \text{ CCO}\) |
| 889                                | 870                | \(v_{as} \text{ N-CH}_3\) |
| 788                                | 783                | \(\omega \text{ C=O}\) |

In the next step, prepared material was characterized using TEM technique. Figure 3A shows the TEM image of microparticle of Al\(_2\)O\(_3\) before modification. Figure 3B indicates the apparent presence of gold nanoparticles located over the surface of aluminum oxide. In the results, we confirmed the success of synthesis of gold nanoparticles on Al\(_2\)O\(_3\) microparticles surface.
Figure 3 TEM images of Al₂O₃ (A), Al₂O₃-AuNPs (B).

The optical characterization of the created system was performed by UV-Vis measurement (Figure 4). The characteristic for pristine Al₂O₃ absorption band is located at 260 nm. Synthesis of AuNPs on Al₂O₃ surface leads to the appearance of peak with the maximum at 540 nm, which is typical absorption band, characteristic for gold nanoparticles.

Figure 4 UV-VIS spectrum of Al₂O₃ and gold nanoparticles on Al₂O₃.

4. CONCLUSION

In this work, deep eutectic solvent was prepared and doped with metal nanoparticles, previously immobilized on Al₂O₃ powder surface. Created system was characterized by FTIR, TEM and UV-Vis method. Such system can be further used for a range of attractive application, with especial attention on green chemistry field.

ACKNOWLEDGEMENTS

This work was supported by under the project IGA 126 88 2109.
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