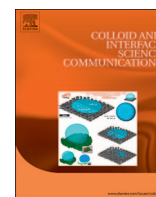


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journal homepage: www.elsevier.com/locate/colcomSilica Nanospheres Coated by Ultrasmall Ag⁰ Nanoparticles for Oxidative Catalytic Application

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ABSTRACT

The present work introduces optimal modification of core-shell composite nanomaterial, where small (2–8 nm) Ag⁰ nanoparticles are deposited onto large (about 140 nm) silica spheres for application in oxidative catalysis. The size of Ag⁰ and density of its deposition onto silica spheres was modified by the post treatment of initially deposited Ag⁰ (about 30 nm) by hydrogen peroxide in specific conditions. The comparison of catalytic effect of the post-treated and initial SN-Ag⁰ in electrochemical phosphonation of benzo(thia)oxazoles by diethyl phosphite in oxidative conditions revealed the difference between the composite nanoparticles. In particular, the post-treated SNs-Ag⁰ nanoparticles exhibit efficient catalytic effect in oxidative conditions resulting in facile and green method for synthesis of phosphonated benzooxa(thia)zoles, while no catalytic effect is observed under the use of larger Ag⁰ nanoparticles deposited onto silica spheres. The use of Ag⁰-based nanomaterial in oxidative catalysis had been never demonstrated before.

Core-shell composite materials, where silica spheres are coated by a noble metal (mainly, gold and silver) shell, are of great interest in nanoscience due to their potential applications in medicine [1], biosensing [2], photonics [3] and catalysis [4]. The deposition of silver (Ag⁰) nanoparticles onto larger-sized silica supports is a convenient tool to prevent their uncontrollable agglomeration [5]. Thus, the choice of the core-shell morphology for colloid stabilization of Ag⁰ nanoparticles is good alternative to their stabilization by another type of core-shell morphology, where Ag⁰ cores are coated by polymeric or dendrimeric shell [6]. The choice of optimal core-shell morphology is of great impact in the catalytic application of silver nanoparticles.

An easy preparation, low cost, high monodispersity, and large particle size range of spherically shaped silica nanoparticles (SNs) are the reasons for their application as the cores for the deposition of Ag⁰. Moreover, both chemical reactivity of Si–OH groups and adsorption capacity of silica surface are the factors facilitating a surface decoration of SNs [7]. The optimal surface decoration facilitates the deposition of Ag⁰ nanoparticles through electrostatic attraction [8] or coordination bonds [9].

Silver nanoparticles have been of great interest as catalysts in organic synthesis among the metal-based nanoparticles, and their catalytic application has expanded rapidly in the past decade [10]. The applicability of the core-shell SN-Ag⁰ nanomaterial in catalysis in great extent is exemplified in the literature by reductive capacity of Ag⁰ [10], while, to the best of our knowledge, the applicability of Ag⁰-based nanomaterial in oxidative catalysis has not been reported. Oxidative catalysis of C-H phosphonation reactions of aromatic compounds is of great impact in organic synthesis, although all documented examples including benzo(thia)oxazoles [11,12] represent homogeneous catalysis. Nevertheless, our previous reports are worth noting as successful application of nanoheterogeneous catalytic electrochemical transformations in organic synthesis [13].

Catalytic application of the core-shell composites based on Ag⁰-deposition onto SNs (SN-Ag⁰) requires optimal Ag⁰ particle size and silica surface decoration density [10]. Nevertheless, a controllable optimization of both the parameters is still open for an improvement, although some fine literature examples are worth noting. The work [14]

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