Research paper

Spherical montmorillonite-supported nano-silver as a self-sedimentary catalyst for methylene blue removal

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Abstract

Supported metal nanoparticles using various substrates have been proven as a highly efficient approach for solving the problems of aggregation and recyclability of metal nanoparticles. However, the reuseability procedure often involved the abundant devices, which obviously increased the cost and heavily limited the large-scale application of metal nanoparticles. In this work, spherical montmorillonite was used firstly as the substrate for supporting silver nanoparticles on its surface through polydopamine chemistry method. The loading of silver nanoparticles with 40 nm in diameter was 15.2 wt% and the specific surface area of this prepared spherical montmorillonite supported silver nanoparticles catalyst was 45.3 m²/g, giving the catalyst an optimized apparent reduction rate constant k of 1.22 min⁻¹ for the reduction of methylene blue. Furthermore, the prepared catalyst with quickly self-sedimentary property in aqueous solution were conveniently recovered and reused without any devices involves. The spherical morphology and catalytic performance of prepared catalyst were almost unaltered after 5 cycles. Our research aims at opening a new avenue to easily realize the reusability of silver nanoparticles through using the substrate with the self-sedimentary property.

1. Introduction

Silver nanoparticles (Ag NPs) have been extensively investigated and proven as an efficient catalyst for reduction of organic dyes (Guo et al., 2017; Liu et al., 2017; Shen, Duan, Guo, Zhao, and Xu, 2015). The key issues of recovery and reuse of these Ag NPs have been significantly considered with respect to the their costs. The substrates including polymer materials (Cao et al., 2017; Xu et al., 2017; Zhu, Wang, Meng, and Diao, 2013), carbon materials (Ji et al., 2016), silica materials (Dong et al., 2014; Khriazianov et al., 2018), cellulose materials (Torkamani and Azizian, 2016), clay materials (Arora, Mehta, Mishra, and Basu, 2018; Zhang et al., 2018), and metal oxides (Xiao et al., 2018; Sedghi, Asadi, Heidari, and Heravi, 2017) have been fully functionalized through various methods and employed to support Ag NPs. This supported approach is now emerging as the most effective strategy to address this issue (Nasrollahzadeh, Atarod, Jaleh, and Gandomirouzbahani, 2016). Among these substrates, recently, magnetic nanoparticles have received greatly attention due to their easy-recovered property from wastewater containing organic dyes through an external magnetic field (Kurtan, Baykal, and Sözeri, 2015; Xie et al., 2014; Yao et al., 2014). Besides, the stable substrates with large surface area for highly loading Ag NPs and strong organic dyes adsorbed capacity were enormously expected due to the increased opportunity for the contacting between Ag NPs and organic dyes to enhance the catalytic performance of the fabricated catalyst (Jeon et al., 2013). These kinds of substrates such as graphene oxide and clay can be recovered through filtration or centrifugation. Until now, despite solving this issue in the laboratory-scale, approaches through the added extra separation and recovery process have yet to be promoted to industrial scale. A major shortcoming of these supported-Ag NPs catalyst is abundant devices involved during the reuse process, which heavily increases the cost of the catalytic procedure and is not well suitable for the industrial applications.

As we know, these substrates with higher density than the wastewater can slowly sink, which can deposit and be recovered from the bottom for some time. Based on the self-sedimentation phenomenon, the substrates with high density and quick sinking ability in wastewater are highly desirable and it is very meaningful to solve an engineering
matter in a physical way. Montmorillonite (Mt) with a mean density of 2–3 g/cm³ is a natural nanomaterial, can strongly adsorb organic dyes and is considered as a promising candidate as a self-sedimentary substrate (Rafatullah, Sulaiman, Hashim, and Ahmad, 2010). In our previous work, Mt was used as the substrate to in situ supported Ag NPs through polydopamine chemistry method (Wang, Hu, and Zhang, 2017). Although the prepared Mt supported Ag NPs exhibited excellent catalytic performance and good recyclability, their large-scale applications are still restricted by the problem of recyclability during the catalytic procedure since the deposited rate of catalyst is limited by the hydration and swelling of Mt and thus abundant expensive recovered devices are involved. Therefore, the fabricated catalyst with quick sedimentary property is highly desirable.

Spray-drying process has been proposed to realize the consolidating nanoparticles into macroscopic compacts (Stunda-Zujeva, Irbe, and Berzina-Cimdina, 2017; Nandiyanoto and Okuyama, 2011). After dissolving the nanoparticles, the solution is passed through atomization, solvent evaporation, drying, and particle collection process to form the final product. Spherical montmorillonite (SMt) (Yawe et al., 2011), silica (Yu et al., 2008), TiO₂ (Suh, Jang, Suh, and Suslick, 2010), and ZrO₂ (Widijayastuti, Wang, Lenggoro, Iskandar, and Okuyama, 2007) with sub-micrometer size to micrometer size were successfully fabricated through controlling the spray-drying process condition. Here, we fabricated a supported-Ag NPs catalyst using the SMt as the substrate. Polydopamine (PDA) was used here as a coating to modify the surface of SMt and played as a reducing agent role to deposit Ag NPs on the surface of SMt. The model organic dye methylene blue (MB) was selected to estimate the catalytic performance. In addition, the recovery and reusability of the prepared catalyst were evaluated. To the best of our knowledge, this is the first report of fabricated SMt-supported Ag NPs using as a catalyst for reduction of MB, paving a new avenue to greatly economically solve an engineering problem in a physical way and exhibiting the great prospect in practical treatment of wastewater containing organic dyes.

2. Materials and methods

2.1. Instruments and reagents

Chemicals and reagents were all obtained from Sinopharm Chemical Reagent Co. Ltd., China and directly used as received. The general formula of montmorillonite is (Na, K, Ca)₀.₃₃(Al₁.₆₇Mg₀.₃₃)Si₂O₅(OH)₂·nH₂O. Na-montmorillonite (Mt) used in our research is kindly provided by Zhejiang Fenghong Co. Ltd. and used without further purification. The cation exchange capacity (CEC) of Na-montmorillonite is about 80 cmol(+) /kg. Samples were characterized with scanning electron microscopy (SEM, S-4800, HITACHI, Japan), X-ray diffraction (XRD, Bruker D8-Advantage powder diffractometer, Germany), and N₂ adsorption (ASAP2020, US). The Ag content was measured through an inductively coupled plasma mass spectrometry (ICP-MS, ELAN DRC II, Perkin Elmer, Hong Kong) and then reused. The key factors such as the concentrations of catalyst, organic dye, and reduce agent were fully investigated.

2.3. Fabrication of SMt-supported Ag NPs

Spherical Mt-supported Ag NPs (Ag@SMt) was synthesized through modifying the Mt with polydopamine (PDA), followed by depositing Ag NPs on the PDA modified Mt. Generally, 1.0 g of dopamine (DA) was added into 200 mL DI water containing 10.0 g of Mt and 20 mM Tris-HCl buffer. The pH was adjusted to 8.5. After shake at room temperature for 6 h, the PDA modified Mt was filtered from the suspension and then washed with alcohol for at least three times. Then PDA modified Mt (PDA@SMt) was further dispersed in 50 mL 100 mM AgNO₃ aqueous solution under dark with 12 h rotator (MX-RL-Pro, SCILLOGEX, USA). After filtration, the brown microspheres were dried at 70 °C for at least 12 h through vacuum oven to obtain the final product Ag@SMt.

2.4. Adsorption and catalytic experiment

About 20 mg of sample and 100 mL of 100 mg L⁻¹ MB aqueous solution were mixed under constant stirring. The adsorption process of MB was monitored by UV–vis absorption spectra. The removal percentage of MB was calculated following the Eq. (1) (Vimonses, Lei, Jin, Chow, and Saint, 2009):

\[
R_{\text{MB}} = \left( \frac{C_0 - C_t}{C_0} \right) \times 100\%
\]

where \( R_{\text{MB}} \) is the reduction efficiency of MB, and \( C_t \) and \( C_0 \) are the concentration of the dyes at time t and 0, respectively.

2.5. Catalytic reduction of MB

The catalytic performance of the Ag@SMt was evaluated by its catalytic reduction capacity towards MB. Typically, 2 mg catalyst, dye solution (1 mL of 250 mg L⁻¹), and NaBH₄ (2 mL of 0.1 M) aqueous solutions were mixed with 7 mL DI water under stirring. The reduced process of MB was monitored at 665 nm using a UV–vis spectrometer. After the whole reduction process was completed, the self-sedimentary Ag@SMt catalyst was easy recovered from the bottom of the solution and then reused. The key factors such as the concentrations of catalyst, organic dye, and reduce agent were fully investigated.

3. Results and discussion

3.1. Synthesis and characterization of Ag@SMt

The spray-drying technology is widely used in practical uses, which has been fully proved by its application in the manufacturing of inorganic materials (Park, Kang, Kwon, and Park, 2018). The main principle of this technology is to deliver and rapidly heat the slurry via the direct injection of very small droplets. Spherically shaped particles are typically generated using the spray-drying method because the spherical form is the most stable structural morphology (Vehring, 2008). Therefore, SMt in our experiment was prepared through this technology and used as the substrate for supporting Ag NPs. In addition, the strong adhesive property of polydopamine (PDA) and excellent reduced capacity towards Ag precursor have extensively investigated (Fu et al., 2015; Liu, Ai, and Lu, 2014). Hence spherical Mt-supported Ag NPs (Ag@SMt) was facile fabricated through PDA chemistry method, as shown in Scheme 1.

To prove the successful preparation of Ag@SMt, SEM was employed to observe the morphologies of spherical Mt (SMT), PDA coated SMt (PDA@SMt), and Ag@SMt, respectively. As shown in Fig. 1a, SMt prepared by the spray-drying technique presents a typical spherical morphology with an average diameter of 25 μm and its spherical morphology does not change under the PDA coated process (Fig. 1b). After the Ag-supported process, small bright nanoparticles with a mean diameter of 40 nm are well observed on the surface of PDA@SMt.
and are further examined as Ag NPs by XPS exhibited in Fig. 1d. These results demonstrate that Ag NPs can be uniformly dispersed on the PDA modified SMt surface and the prepared Ag@SMt can maintain a stable spherical morphology even through long time coating and supported process. The curves of N₂ adsorption-desorption isotherm of samples are shown in the inset of Fig. 1. SMt with a typical mesoporous structure is observed. The specific surface area and total pore volume are calculated as 60.2 m²/g and 0.37 cm⁻³/g, which is much higher than those of pristine Mt (31.8 m²/g and 0.26 cm⁻³/g, Fig. S1 and Table 1), indicating the effective spherical-made process. After the coated with PDA, the specific surface area and total pore volume slightly decrease 54.2 m²/g and 0.36 cm⁻³/g. Compared with the SMt, after the Ag-supported process, the specific surface area of Ag@SMt further decreases to 45.3 m²/g, while the total pore volume obviously decreased to 0.30 cm⁻³/g. Combined with Ag NPs content on Ag@SMt which is calculated to be 15.2 wt% from the quantified by ICP/MS, the dramatic decreases of specific surface area and the total pore volume are mainly due to the occupation of large amount of Ag NPs on SMt surface (Wang et al., 2017).

To investigate the effect of granulation process, PDA coated process, and Ag-supported process on Mt, XRD was performed to measure the d-spacing of all the samples. Fig. 2 exhibits that the d-space of SMt is increased to 1.54 nm after acid treatment, indicating the acid dissolves the octahedral layer (Myriam, 1998). Upon the coated process, the d-space further increases to 1.95 nm, suggesting the thickness of PDA coated on SMt is about 0.21 nm. However, d-space is almost unaltered while the (001) reflection has low intensity after the Ag-supported process, illustrating that delamination happened and most Ag NPs grew on the external surface of SMt, which is well consistent with the reported pristine Mt supported Ag NPs in our previous work (Wang et al., 2017). In addition, four peaks at 2θ of 38.1°, 44.4°, 64.4°, and 77.8° appear, which should be assigned to the (111), (200), (220), and (311) crystal faces of Ag NPs (fcc, JCPDS card no. 04–0783), confirming the successfully supported Ag NPs on the SMt surface (Guo et al., 2017). Furthermore, we used the Scherrer equation \( Dhkl = \frac{k \lambda}{\beta \cos \theta} \) and calculated the crystallite of the Ag NPs as 0.4 nm. Considering the size of Ag NPs observed from SEM as 40 nm, the Ag NPs are composed by a large amount of crystallite.

### 3.2. Adsorption of Ag@SMt

Recently, most of the researchers have paid attention to organic dyes absorbed capacity of substrates since it is possible to increase the opportunity for the contacting between supported Ag NPs and organic dyes to enhance the catalytic performance of catalysts (Xie et al., 2014).
and PDA@SMt show the large adsorption of MB with 48.0 mg g\(^{-1}\) due to the di
stages. In the primary stage, the MB quickly adsorbs on the surface of PDA@SMt, and Ag@SMt present the adsorption behavior with three
increased concentration of NaBH\(_4\) to 60 mM. It can be understood that k decreases with MB while increases with NaBH\(_4\) concentration under limited Ag NPs surface because the coverage degree of MB and NaBH\(_4\) tends to a constant
increased valid surface area of Ag NPs (Wunder, Lu, Albrecht, and Ballau, 2011). Further increasing the concentration of Ag@SMt would not enhance the catalytic performance due to enough surface area providing for electron transfer.

The catalytic mechanism of Ag NPs is based on the electron transfer from the electron donor NaBH\(_4\) to electron acceptor MB. Therefore, the catalytic reduction process mainly occurs on the Ag NPs surface. Therefore, the competition between the electron donor and the electron acceptor on the available Ag NPs surface would greatly influence the k.

As shown in Fig. 4c, when fixing the concentrations of NaBH\(_4\) and Ag@SMt as 0.02 M and 0.2 g L\(^{-1}\), the k first decreases to 0.20 min\(^{-1}\) as MB concentration increases to 400 mg L\(^{-1}\), and then unaltered with the further increased MB concentration. On the contrary, as shown in Fig. 4d, when fixing the concentrations of MB and Ag@SMt as 250 mg L\(^{-1}\) and 0.2 g L\(^{-1}\), the k increases to 0.52 min\(^{-1}\) as the NaBH\(_4\) concentration increases to 40 mM, and then unchanged with the further increased concentration of NaBH\(_4\) to 60 mM. It can be understood that k decreases with MB while increases with NaBH\(_4\) concentration under limited Ag NPs surface because the coverage degree of MB and NaBH\(_4\) on the Ag NPs surface exist an equilibrium. The increasing of MB results in the increasing of the coverage degree of MB while decreasing the coverage degree of NaBH\(_4\), k keeps a constant at higher concentration of MB since the coverage degree of MB and NaBH\(_4\) tends to a constant with a large coverage degree of MB and small coverage degree of NaBH\(_4\), vice versa. All these results demonstrate that the k of our synthesized Ag@SMt can be well optimized by tailoring the concentrations of catalyst, dyes, and NaBH\(_4\). It is worth to note that the optimized k (1.22 min\(^{-1}\)) is much higher than those (0.34 min\(^{-1}\) (Zhu et al., 2013), 0.43 min\(^{-1}\) (Xie et al., 2014), 0.058 min\(^{-1}\) (Tang et al., 2015), and 0.25 min\(^{-1}\) (Sahoo, Kumar, Thiyagarajan, Thakur, and Panda, 2018)) reported in the literature, which can be contributed to the uniform dispersion and high loading of Ag NPs on the SMt with strong adsorption capacity towards organic dyes.

3.3. Catalytic properties of Ag@SMt

Ag NPs have been proven as an efficient catalyst towards the reduction of organic dyes and the supported Ag NPs exhibited higher catalytic performance than single Ag NPs due to the enhanced adsorption capacity (Guo et al., 2017). Therefore, the catalytic capacity of Ag@SMt was evaluated through the reduction of MB in the presence of NaBH\(_4\). Typically, at the beginning of MB reduction, the peak at 665 nm decreases its intensity, hence the catalytic reduction process of MB is feasibly monitored by its time-dependent absorbance change using UV–vis spectroscopy. As shown in Fig. 4a, with the prolonged reduction time, the absorbance peak of MB at 665 nm rapidly decreases to zero in 15 min with the blue color of MB solution gradually disappears, indicating the degradation of carcinogenic MB to colorless and inactive form leucomethylene blue (Rice, 2001). The catalytic performance of Ag@SMt catalyst is much related to the valid surface area of Ag NPs for the electron relay from NaBH\(_4\) to MB. Therefore, the reaction kinetics of the reduction of MB under different catalytic conditions are calculated by monitoring their time-dependent UV–Vis absorption spectra. From the linear relation of ln(C\(_0\)/C) with the prolonged reaction time, the apparent reduction rate constant (k) is received. Fig. 4b shows that k increases to 0.98 min\(^{-1}\) with increased Ag@SMt concentration when fixing the concentrations of MB and NaBH\(_4\) as 250 mg L\(^{-1}\) and 0.02 M, respectively. The increased k is mainly due to the increased valid surface area of Ag NPs (Wunder, Lu, Albrecht, and Ballau, 2011). Further increasing the concentration of Ag@SMt would not enhance the catalytic performance due to enough surface area providing for electron transfer.

Recyclability is the key property of catalyst to maintain its sustainable and efficient performance besides lower its cost for its practical applications. The successive 5 cycles were carried out to determine its recyclability, as shown in Fig. 5a. For one cycle, we first mixed Ag@...
SMt, MB, and NaBH₄ together under a shaker, with the prolonged reaction time, the blue color vanishes. Then static settlement for 5 min, followed by pouring the reactive solution. The prepared Ag@SMt exhibits an excellent catalytic performance with unaltered reduction efficiency in the tested five cycles (Fig. 5b). Furthermore, the k slight decreases to 90% in the 5th cycle (Fig. 5c). Obviously, the self-sedimentary substrate SMt plays an important role in the recovery of Ag catalyst. The weight loss percentage of Ag@SMt after 1st, 3rd, and 5th cycle is determined to be 3.2 wt%, 9.5 wt%, and 12.1 wt%, respectively. In addition, after 5th cycle, as shown in Fig. 5d, the prepared Ag@SMt still maintains spherical morphology with Ag NPs supported on SMt, reflecting the good stability of the prepared catalyst in the aqueous solution.

Fig. 4. Time-resolved UV–Vis spectra of MB solutions (a). The apparent reduction rate constant (k) of MB changes as a function of the concentration of Ag@SMt (b), MB (c), and NaBH₄ (d), respectively.

Fig. 5. Scheme of the cycle process of Ag@SMt to catalytic reduction of MB (a), the changes of reduction efficiency (b) and apparent reduction rate constants k of Ag@SMt in the reduction of MB with the prolonged cycle number (c), and SEM images of Ag@SMt microspheres after 5th cycle (d).
4. Conclusions
By fabricating spherical montmorillonite through spray-drying technology, the engineering problem is conveniently solved in a physical way. In this work, we first selected sphericalmontmorillonite (SMt) as the substrate to in situ support Ag NPs through polylamine chemistry method. The prepared Ag@SMt with 15.2 wt% of Ag NPs loading and 45.3 m2/g of the specific surface area exhibits an optimized apparent reduction rate constant k of 1.22 min⁻¹ for catalytic reduction of methylene blue. More importantly, the prepared Ag@SMt can be easily recovered and reused without any devices involved due to the self-sedimentary property of SMt. In addition, Ag@SMt shows excellent stability and recyclability for five cycles with unaltered spherical morphology and almost unchanged catalytic performance. Our strategy to solve the engineering issue in a physical way may pave a new avenue to easily realize the application of Ag NPs to the large-scale treatment of wastewater containing organic dyes.

Author contributions
All authors have given approval to the final version of the manuscript. Ning Wang and Fengyan Xiao contributed equally to this work.

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Appendix A. Supplementary data
Supplementary data to this article can be found online at https://doi.org/10.1016/j.clay.2019.03.021.

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