

CTAB micelles assisted rGO-AgNP hybrids for SERS detection of polycyclic aromatic hydrocarbons Postprint

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Abstract

A structure of hexadecyl trimethyl ammonium bromide (CTAB) micelle-assisted reduced graphene oxide-Ag nanoparticle (rGO-AgNP) hybrids is designed and fabricated for SERS detection of nonpolar polycyclic aromatic hydrocarbons (PAHs), in which CTAB micelles

Full Text

Preamble

CTAB Micelles Assisted rGO-AgNP Hybrids for SERS Detection of Polycyclic Aromatic Hydrocarbons

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A structure of hexadecyl trimethyl ammonium bromide (CTAB) micelle-assisted reduced graphene oxide-Ag nanoparticle (rGO-AgNP) hybrids is designed and fabricated for SERS detection of nonpolar polycyclic aromatic hydrocarbons (PAHs), in which CTAB micelles act as the host material to capture PAH molecules. This method provides stable aqueous suspensions of functionalized graphene with an alkyl chain, since the rGO-AgNP hybrids do not need to be pre-modified by CTAB. The result shows that the CTAB-assisted rGO-AgNP substrate has excellent SERS performance toward PAHs and ideal stability under continuous laser radiation. With further optimization, the detection limits of pyrene and perylene were $10(\text{cid:2})6 \text{ M}$ and $10(\text{cid:2})7 \text{ M}$, respectively. Two different PAH molecules could be detected simultaneously by their characteristic peaks.

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Introduction

Surface-enhanced Raman scattering (SERS) is a very powerful tool for chemical and biological analysis, and has aroused extensive research attention.¹⁻⁴ When an analyte is adsorbed on noble metallic nanostructures, SERS offers several orders of magnitude enhancement in Raman signal intensity of the analyte. The enhancement factor is crucially dependent on the magnitude of the localized electromagnetic field originating from the excited localized surface plasmon within the metallic nanostructures, which can reach a high value of 10⁶-10⁸.^{5,6} This allows single-molecule detection using SERS.⁷⁻⁹ Considering the limited influence of water and the narrow bandwidth of the Raman peaks, the SERS technique is suited especially for the direct detection of multicomponent mixtures in water.^{10,11}

Graphene, a monolayer composed of sp²-hybridized carbon atoms,¹² is widely used in the energy, electronic and biology fields due to its unique physical and chemical properties.¹³⁻¹⁵ Owing to its two-dimensional (2D) planar structure, graphene has been utilized as a building block for preparing nanocomposites. Several investigations of organic-molecule-decorated graphene and inorganic-nanoparticle-decorated graphene composites were reported.¹⁶⁻²⁰ Among them, graphene-metal hybrids used as a new type of SERS substrate have attracted increasing interest.²¹⁻²³ Because of the large surface area and interaction with the target molecules, graphene sheets play a role in gathering the analyte molecules. Graphene sheets act as a holder which makes metal nanoparticles (NPs) aggregate, thus generating more hot-spots.²⁴ Therefore, graphene-metal hybrids exhibited a more sensitive response to organics such as folic acid, crystal violet, amaranth, and chrysoidin.²⁵⁻²⁸

Nonetheless, there are few reports on SERS detection of non-polar polycyclic aromatic hydrocarbons (PAHs) by graphene-metal nanoparticle hybrids. Although graphene enriches PAHs via p-p interaction, the poor affinity between PAHs and metal NPs prevents them approaching, which results in low SERS activity. Metal surface hydrophobization makes it possible to catch PAHs that are close to metal NPs. Du et al.²⁹ and Jones et al.³⁰ used thiol-modified Fe₃O₄@AgNPs and an Ag film as SERS substrates for PAH detection, respectively. However, when alkyl groups are grafted on the metal NPs, the resultant graphene-NP hybrids can no longer be re-dispersed in water or ethanol because they become more hydrophobic. On the contrary, the graphene-NP hybrids float on the surface of water or ethanol. Hence, the application of graphene-metal hybrids for the detection of PAHs is still a challenge.

Here, we prepared graphene-AgNP (rGO-AgNP) hybrids through one-pot reduction of silver nitrate and graphene oxide (GO) by trisodium citrate. With the aid of hexadecyl trimethyl ammonium bromide (CTAB), the resultant

rGO-AgNP hybrids were used as active SERS substrates for the ultrasensitive and label-free detection of PAHs in water. CTAB micelles anchored the PAH molecules on the rGO-AgNP sheets. On the basis of the strong SERS enhancement and enrichment toward PAHs, a low detection limit was achieved. The SERS signal possesses temporal stability under continuous laser radiation as well as a good resolution. Scheme 1 illustrates the proposed fabrication of rGO-AgNP hybrids and the detection procedure.

Experimental

Materials

Silver nitrate (AgNO₃, AR), trisodium citrate (Na₃C₆H₅O₇(cid:3)2H₂O, AR), CTAB (C₁₉H₄₂NBr, AR), and ethanol (AR) were purchased from Sinopharm Chemical Reagent Co., Ltd. (Shanghai, China). Pyrene (C₁₆H₁₀, AR) and perylene (C₂₀H₁₂, AR) were purchased from Aladdin Reagent Co., Ltd. (Shanghai, China). The GO sample was provided by the Liu Zhaoping group (Ningbo Institute of Material Technology and Engineering, CAS). All chemicals were used as-received without further purification. The water used throughout all experiments was deionized (DI) water from a Millipore-Q purification system (Millipore, USA) of resistivity 18.2 MO cm.

Synthesis of rGO-AgNP Hybrids

The rGO-AgNP hybrids were prepared based on a modified one-pot method.³¹ In a typical experiment, 20 mL of AgNO₃ solution (1.70 mg mL(cid:2)1), 1 mL of graphene oxide aqueous suspension (1 mg mL(cid:2)1) and 60 mL of DI water were mixed and heated rapidly to 90 °C in a water bath. Subsequently, 20 mL of trisodium citrate solution (10 mg mL(cid:2)1) was added under vigorous stirring. The resulting mixture was held at 90 °C for 24 h. After several washing and centrifugation steps, the resulting rGO-AgNP hybrids were re-dispersed in water (Ag, 1.08 mg mL(cid:2)1).

Sample Preparation for SERS Detection

The aqueous solutions of PAHs were prepared in concentration ranges from 10⁻⁷ M to 10⁻⁴ M. 0.1 mL of CTAB aqueous solution (2 mg mL(cid:2)1) was mixed with 0.9 mL of PAHs solution under vigorous stirring, followed by injection of 0.1 mL of an rGO-AgNP hybrid suspension. After sonication for 30 min, 10 mL of the resulting mixture was dropped on a cleaned glass slide, dried at ambient temperature, and measured by Raman spectrometry.

Characterization

X-Ray photoelectron spectroscopy (XPS) measurement was performed on an AXIS ULTRA DLD (Shimadzu, Japan) multifunctional X-ray photoelectron

spectroscopy with Mg or Al K α X-ray radiation as the X-ray source for excitation. Transmission electron microscope (TEM) measurements were performed on a Tecnai F20 TEM (FEI, USA) with an accelerating voltage of 200 kV. UV-visible absorption spectra were obtained on a Lambda 950 spectrophotometer (Perkin Elmer, American). XRD data were measured on a D8 advance X-ray diffractometer (Bruker, Germany) with a Cu K α X-ray radiation source. SERS spectra were collected with a Renishaw inVia model confocal microscopy Raman spectrometer (Renishaw, UK). A 532 nm laser was used for SERS excitation. To obtain comparable spectra, the test conditions of laser power and exposure time were all the same.

Results and Discussion

Characterization of the rGO-AgNP Hybrids

Fig. 1 [Figure 1: see original paper] presents the XRD patterns of GO and rGO-AgNP hybrids. The sharp peak at $2\theta = 11.21$ on the curve of GO was assigned to the (001) reflection of GO.³² After reduction with trisodium citrate, four diffraction peaks corresponding to the (111), (200), (220), and (311) crystalline planes of silver (JCPDS 04-0783) presented at $2\theta = 38.21, 44.31, 64.51,$ and 77.31 , respectively, which confirmed the existence of metallic Ag. In comparison with GO, the absence of the peak of GO(001) on the pattern of rGO-AgNP hybrids indicated that the regular stacks of GO were destroyed, which was caused by the insertion of AgNPs into layers of G sheets.^{33,34}

The UV-Vis spectra of GO and rGO-AgNP hybrid suspensions are shown in Fig. 2 [Figure 2: see original paper]. The formation of rGO-AgNP hybrids was further confirmed by UV-Vis measurement. The UV-Vis spectrum of the GO suspension exhibited a maximum absorption peak at 227 nm, which was attributed to π - π^* transitions of aromatic C=C bonds. This peak red shifted to 257 nm in the spectrum of rGO-AgNP hybrids, confirming the restoration of π -conjugation during the reduction procedure. A relatively narrow new peak appeared at 409 nm, which was due to the packed AgNPs on the surface of the G sheet. The above result indicated the formation of rGO-AgNP hybrids.

To verify the transformation from GO to G, the corresponding XPS spectra of the GO sheets and the rGO-AgNP hybrids were carried out. As shown in Fig. 3A [Figure 3: see original paper], three component peaks corresponding to carbon atoms in different chemical environments were distinguished. These peaks at 284.6 eV, 286.8 eV, and 288.1 eV originated from C-C, C-O, and C=O bonds, respectively.³⁵ Compared with GO, the peak intensities of C-O and C=O bonds decreased obviously relative to that of the C-C bond in the C1s XPS spectrum of rGO-AgNP, which indicated that most of the oxygen containing functional groups had been removed after reduction.

The morphology of rGO-AgNP hybrids was investigated by TEM. Fig. 4 [Figure 4: see original paper] shows the representative TEM images of the hybrid at different magnifications. It is seen clearly that the almost transparent graphene

sheet was decorated with a large amount of silver nanoparticles. From the overlap of silver nanoparticles it was inferred that the nanoparticles were loaded onto both sides of the G sheet. The higher magnification image (Fig. 4B) shows that the silver nanoparticles are almost all in a roughly spherical morphology with an average diameter of about 20 nm.

Effect of CTAB

To confirm the effect of CTAB, a contrast experiment was performed. Fig. 5 [Figure 5: see original paper] shows the SERS spectra of pyrene on the rGO-AgNP hybrids substrate with CTAB and without CTAB. When CTAB was applied, a series of primary vibration bands were observed in the SERS spectrum of pyrene on the rGO-AgNP hybrids, which was confirmed according to a reported work.³⁶ The bands at 408 cm^{-1} and 591 cm^{-1} were assigned to the skeletal stretching vibration mode, and the band at 1064 cm^{-1} was attributed to the C-H in-plane bending mode. The band at 1239 cm^{-1} originated from the C-C stretching and C-H in-plane bending modes. The band at 1405 cm^{-1} was assigned to the C-C stretching coupled with ring breathing modes. The bands at 1593 cm^{-1} and 1627 cm^{-1} were attributed to the C-C stretching mode. In the absence of CTAB, no characteristic band of pyrene on the rGO-AgNP hybrids was found.

The above results indicate that CTAB plays a role in the detection of pyrene. CTAB is a kind of cationic surfactant with a long alkyl chain, which forms micelles in the water and solubilizes the hydrophobic molecules. The hydrophobic pyrene molecules are enclosed by the micelles consisting of CTAB. The micelles adsorb to the surface of the rGO-AgNP hybrid due to electrostatic adsorption. This step is similar to the self-assembly of a thiolate-aliphatic monolayer on noble metal nanoparticles. The difference is that several processes of surface modification, enrichment and bringing analyte molecules close to the plasmonic electric field were combined into one step in our work. This method makes it possible to obtain stable aqueous suspensions of functionalized graphene with alkyl chains.

Optimization of the SERS Substrate

The amount of CTAB was optimized to achieve a stronger Raman signal of the analyte molecule with fewer disturbing signals from CTAB. As shown in Fig. 6 [Figure 6: see original paper], the Raman intensity of perylene increased with the increase of the concentration of CTAB (0.002-0.5 mg mL^{-1}). When the concentration of CTAB was lower than 0.002 mg mL^{-1} , no peak of pyrene was observed in the spectrum. If the concentration of CTAB was over 0.5 mg mL^{-1} , the characteristic peaks of CTAB became stronger and interfered with the signal of the target molecules. Based on an overall consideration of these factors, the optimal concentration of CTAB was 0.2 mg mL^{-1} .

SERS Detection of PAHs

The SERS spectra of pyrene with different concentrations are shown in Fig. 7A [Figure 7: see original paper]. As observed, the intensity of bands decreased as the concentration of pyrene decreased. When the concentration of pyrene was below 10(cid:2)6 M, its characteristic bands were hardly recognized.

As shown in Fig. 7B, the SERS spectra of perylene exhibited five major peaks at 548 cm(cid:2)1 (C-C-C bending), 978 cm(cid:2)1 (C-C stretching), 1298 cm(cid:2)1 (C-C stretching), 1374 cm(cid:2)1 (C-C stretching), and 1567 cm(cid:2)1 (C-C stretching), respectively.³⁷ Raman spectra of the solid perylene are also presented for comparison. The detection limit of perylene was 10(cid:2)7 M. Generally speaking, our strategy was more sensitive in detecting perylene than pyrene.

The identification of a complex system that contains both perylene and pyrene was also performed. Fig. 8 [Figure 8: see original paper] shows the SERS spectra obtained from a mixture of 10(cid:2)5 M pyrene and 10(cid:2)5 M perylene (black), 10(cid:2)6 M pyrene and 10(cid:2)6 M perylene (red), and 4 (cid:4) 10(cid:2)7 M pyrene and 4 (cid:4) 10(cid:2)7 M perylene (blue) on rGO-AgNP hybrids substrate. The peaks marked with a “star” are assigned to PYR and the peaks marked with a “rhombus” are assigned to PER. The main peaks of individual PAHs were clearly observed and distinguished. The signal of perylene was much stronger than that of pyrene at the same concentration, because the method was more sensitive with perylene. The strong peaks of perylene won't overlap the weak peaks of pyrene and their detection limit was almost the same as the single component detection result, indicating that competitive adsorption between PAHs could be ignored in our work.

Stability of SERS Spectra

Stable signals are significant to obtain accurate analytical results. In order to inspect the temporal stability of the substrate, the SERS spectra of perylene on rGO-AgNP were collected under continuous laser irradiation in air for 1 h. The spectra were recorded every 4 min at the same point of the rGO-AgNP hybrids. As shown in Fig. 9 [Figure 9: see original paper], there is no obvious change in intensity and overall shape of the Raman bands during the laser irradiation. The relative standard deviation of the intensity of the band at 1567 cm(cid:2)1 is 7.0%, which shows that the substrate possessed good stability under long time laser exposure.

Conclusions

This work provides a method to detect PAHs with CTAB-assisted rGO-AgNP hybrids as a SERS substrate. CTAB micelles acted as the host material to capture PAHs molecules by means of hydrophobic interaction, and were adsorbed to the surface of rGO-AgNP hybrids, which increased the affinity between PAHs

molecules and the SERS substrate. After the optimization of the components of the substrate, the detection limits of pyrene and perylene were 10(cid:2)6 M and 10(cid:2)7 M, respectively. The stability test result shows little signal reduction under long time laser exposure, which indicated the good stability of our substrate.

Conflicts of Interest

The authors declare no competing financial interest.

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Fig. 1 XRD patterns of the GO (I) and rGO-AgNP (II) hybrids.

Fig. 2 UV-Vis absorption spectra of aqueous dispersions of the GO sheets and rGO-AgNP hybrids.

Fig. 3 C1s XPS spectra of the GO (A) and G-AgNP hybrids (B).

Fig. 4 TEM images of the rGO-AgNP hybrids at different magnifications.

Fig. 5 Raman spectrum of solid pyrene (blue), solid CTAB (green) and SERS spectra of pyrene (10(cid:2)4 M) on the rGO-AgNP hybrids substrate with CTAB (red), and without CTAB (black).

Fig. 6 SERS spectra of pyrene (10(cid:2)4 M) obtained on rGO-AgNP hybrids with different contents of CTAB.

Fig. 7 Raman spectrum of pyrene and the SERS spectra of different concentrations of pyrene on the rGO-AgNP hybrid substrate (A), Raman spectrum of perylene and SERS spectra of different concentrations of perylene on the rGO-AgNP hybrid substrate (B).

Fig. 8 SERS spectra obtained from the mixture of 10(cid:2)5 M pyrene and 10(cid:2)5 M perylene (black), 10(cid:2)6 M pyrene and 10(cid:2)6 M perylene (red), 4 (cid:4) 10(cid:2)7 M pyrene and 4 (cid:4) 10(cid:2)7 M perylene (blue) on rGO-AgNP hybrids substrate. The peaks marked with a “star” are assigned to PYR and the peaks marked with a “rhombus” are assigned to PER.

Fig. 9 The stability of the rGO-AgNP hybrid substrate probed with perylene (10(cid:2)6 M) under continuous laser irradiation.

Scheme 1 The protocol for the fabrication of rGO-AgNP hybrids and their application in SERS detection of PAHs.

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