Quantitative SERS sensors for environmental analysis of naphthalene

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In the investigation of chemical pollutants, such as PAHs (Polycyclic Aromatic Hydrocarbons) at low concentration in aqueous medium, Surface-Enhanced Raman Scattering (SERS) stands for an alternative to the inherent low cross-section of normal Raman scattering. Indeed, SERS is a very sensitive spectroscopic technique due to the excitation of the surface plasmon modes of the nanostructured metallic film. The surface of quartz substrates was coated with a hydrophobic film obtained by silanization and subsequently reacted with polystyrene (PS) beads coated with gold nanoparticles. The hydrophobic surface of the SERS substrates pre-concentrates non-polar molecules such as naphthalene. Under laser excitation, the SERS-active substrates allow the detection and the identification of the target molecules localized close to the gold nanoparticles. The morphology of the SERS substrates based on polystyrene beads surrounded by gold nanoparticles was characterized by scanning electron microscopy (SEM). Furthermore, the Raman fingerprint of the polystyrene stands for an internal spectral reference. To this extent, an innovative method to detect and to quantify organic molecules, as naphthalene in the range of 1 to 20 ppm, in aqueous media was carried out. Such SERS-active substrates tend towards an application as quantitative SERS sensors for the environmental analysis of naphthalene.

Introduction

High quality factors of surface plasmon modes lead Surface-Enhanced Raman Scattering (SERS) as a very sensitive spectroscopic technique. Huge Raman enhancements, up to $10^{14}$-fold\textsuperscript{1–5} the normal Raman signal of non-adsorbed molecules, have been observed when molecules are adsorbed near metallic surfaces. Latter, as silver, gold, copper, exhibit high optical reflectivity.\textsuperscript{a} The short-range chemical model and the long-range electromagnetic model\textsuperscript{7–10} help to reveal the Raman enhancement origin. The first one is related to a charge transfer between the roughened surface and the target molecule via an increase in adsorbed molecule polarisability. The second one is linked to the excitation of surface plasmons leading to an increase in the local electromagnetic field. A final Raman enhancement of an adsorbed molecule can be considered as the coupling of both mechanisms.

Many studies were realized on polystyrene (PS) beads chemically modified for the purpose of grafting metal nanoparticles. Functionalization of the polystyrene beads can occur thanks to a modification of the surface electric charge,\textsuperscript{11–13} a thiolation\textsuperscript{14} or an amination\textsuperscript{15} of the PS beads. In both latter cases, the functional groups –SH or –NH\textsubscript{2} have an affinity for metallic particles: gold is bound to the surface through covalent interactions.\textsuperscript{16–18}

The fabrication of SERS-active substrates through PS beads coated with metal nanoparticles has been lately employed in a wide variety of applications. For example, the crystal violet dye as a probe has been used,\textsuperscript{19} the identification of benzenethiol adsorbed on Ag films,\textsuperscript{20} the detection of streptavidin molecules after biotin grafting\textsuperscript{21} and the investigation for chemical components in saliva.\textsuperscript{22} In our case, a facile method based on Debye adsorption – the interaction between a permanent dipole and an induced dipole – is employed to coat non-polar PS beads with gold nanoparticles. Although silver presents a higher SERS response with an enhancement factor 10–100-fold greater than gold, Au was chosen here on account of AgCl formation on silver surfaces.\textsuperscript{23} Such a facile method used in core–shell fabrication\textsuperscript{24} has been chosen here in order to avoid the addition of organic matter with terminal functional groups whose Raman fingerprints may represent a drawback for the SERS spectra exploitation. Subsequently, PS beads decorated with gold nanoparticles were immobilized onto a quartz substrate previously functionalized with 3-(mercaptopropyl)trimethoxysilane (MPTMS). The MPTMS film and the PS beads stand for the hydrophobic part allowing the pre-concentration of apolar molecules such as PAHs (Polycyclic Aromatic Hydrocarbons).

Herein the aim of the study is to give an innovative method to detect and to quantify organic molecules as PAHs at low concentrations in aqueous medium.
concentration in aqueous media. Such target molecules are of interest for the exploitation and exploration of oceanic and energetic resources. In addition, the detection of PAHs, such as naphthalene, which is a chemical pollutant, presents a societal interest (DCE Directive Cadre sur l’Eau 200/60/CE). Nowadays, there is a lack of a relevant technique that allows qualitative, quantitative, fast and in situ analysis of PAHs in aqueous media.

The low solubility of PAHs in the marine medium requires the development of sensitive substrates. Hydrophobic SERS-active substrates offer the expected solution.\textsuperscript{25–29} Several papers present the SERS effect as a quantitative analytical tool using: the quantitative control of the substrate-withdrawal method onto adsorbed sodium benzoate,\textsuperscript{30} the partial least-squares (PLS) algorithm to determine glucose detection,\textsuperscript{31} self-assembled monolayers (SAMs) as an internal standards and PLS,\textsuperscript{32} isotopically labeled creatinine as internal standard and PLS,\textsuperscript{33} nicotine and pyridine respective deuterated isotopomers as internal standards in a microfluidic system.\textsuperscript{34} To the best of our knowledge, this paper reports for the first time SERS measurements of adsorbed sodium benzoate, quantitatively fast and in situ analysis of PAHs in aqueous media.

Experimental

Instrumentation

SERS spectra of naphthalene solutions were recorded at room temperature with a Raman spectrometer (Labram HR800 – Horiba Jobin Yvon) using the 632.8 nm excitation line of a He–Ne laser (power 10 μW at the sample) in backscattering geometry. The laser beam was focused through a water immersion lens (×100, NA = 1) to a spot of around 1 μm\textsuperscript{2}. Scattered radiation was collected at 180° relative to the excitation beam and detected with an Andor CCD cooled by the Peltier effect. A 300 g mm\textsuperscript{-1} grating coupled to an 800 mm spectrograph allows a spectral resolution of around 2 cm\textsuperscript{-1}. Spectral calibration was performed on silicon sample (at 520 cm\textsuperscript{-1}). Two stage edge filters allow Raman Stokes studies. The accumulation times were 2 × 10 s.

Scanning Electron Microscopy (SEM) measurements were done using a field emission scanning electron microscope (FEI-Quanta200). Before loading into the observation chamber, the samples were coated with a thin Au film (50 nm) by a field emission scanning electron microscope (FEI-Quanta200). Before loading into the observation chamber, the samples were coated with a thin Au film (50 nm) by sputtering to avoid a surface charging effect.

Chemicals and materials

HAuCl\textsubscript{4}·3H\textsubscript{2}O was purchased from Acros Organics. Polystyrene (PS) beads with 800 nm diameter (±50 nm), hydrochloric acid (37%) and (3-mercaptopropyl)trimethoxysilane (MPTMS, 95%) were obtained from Sigma-Aldrich. Trisodium citrate was purchased from Prolabo, nitric acid (65%) was from Merck, hydrogen peroxide (30%) was from J.T. Baker, ammonium hydroxide (33%) was from Riedel-de-Haën, methanol and ethanol were of analytical grade from Fisher. Napthalene from Fisher was used as the analytical molecular reference.

The quartz substrates were tailored with 8 mm diameter and 0.8 mm thickness from Heraeus. The quartz was cleaned using \textit{aqua regia} (25% HCl, 25% HNO\textsubscript{3}, 50% distilled water v/v) and afterwards a base ‘piranha’ solution (75% NH\textsubscript{4}OH, 25% H\textsubscript{2}O\textsubscript{2} v/v) to eliminate organic compounds and to promote hydroxylated surfaces.

Polystyrene beads coated with gold nanoparticles and naphthalene solutions

The solution of PS beads coated with gold nanoparticles was prepared taking inspiration from the method described by Frens.\textsuperscript{35} 20 μL of PS beads of the commercial solution (3.6 × 10\textsuperscript{11} particles per milliliter) dispersed in 5 mL of distilled water were added to 5 mL of an ethanolic solution of tetrachloroauric acid at 2 × 10\textsuperscript{-3} M under stirring. The reactor was placed on a hot-plate and magnetically stirred. Temperature mixture was controlled with a thermometer (−90 °C). Then, 10 mL of sodium tricitrate at 4 × 10\textsuperscript{-3} M was slowly added. After the addition of the reducing agent, the gold particles start to form during a process known as nucleation and subsequently grow. The initially dark-grey solution turned to a red-violet hue after thirty minutes of continued boiling. After one hour of boiling, the mixture was cooled at room temperature under stirring.

The naphthalene solution was prepared by dissolving PAH in methanol, wherein it is more soluble, and diluted in distilled water. Naphthalene in pure water at 25 °C is soluble at 31.8 mg L\textsuperscript{-1}. A 20 ppm naphthalene solution was prepared from 62 μL of 0.05 mol L\textsuperscript{-1} naphthalene methanol solution and the corresponding blank composed of 62 μL of methanol completed by 20 mL of distilled water.

SERS substrates

The strategy for obtaining SERS-active substrates was to graft silanes with terminal –SH groups. The silanes are bound chemically to the substrate \textit{via} hydroxyl groups existing at the quartz surface.\textsuperscript{36} With MPTMS silanization, the quartz surface properties are modified and the non-polar –CH\textsubscript{2} groups give hydrophobic surfaces. The non-polar PS beads decorated with gold nanoparticles were strongly grafted to the functionalized quartz surface \textit{via} covalent bonds between –SH and Au.\textsuperscript{37}

Clean disk quartz substrates were immersed in a 5% (v/v) methanol solution of MPTMS for four hours according to the results obtained by Seitz et al.\textsuperscript{38} After silanization, the substrates were rinsed several times in methanol, sonicated in methanol for twenty minutes to remove excess silane and dried for one hour at 100 °C in an air oven. Then, the silanized quartz were immersed overnight in the gold coated PS bead solution. Finally, SERS substrates were dried for ten minutes at 100 °C. Suitable substrates for adsorbed hydrophobic molecule species and SERS measurements were thus synthesized. Fig. 1 presents a schematic representation of the SERS substrate fabrication procedure.

Results and discussion

Within the scope of characterizing the morphology of the SERS substrates, Scanning Electron Microscopy (SEM) measurements were carried out. Fig. 2(a) shows an overview of the SERS substrate surfaces. The PS beads, with a measured diameter of around 800 nm, are deployed onto the substrate surface as a monolayer. The SEM images clearly show that the PS bead surfaces are partially coated; gold nanoparticles were locally
grafted to the PS bead surface. Besides gold nanoparticles are also present on the MPTMS. Gold nanoparticles have an estimated diameter of \( \frac{C_2}{50} \) nm (Fig. 2(b)). Nevertheless, such a gold coating of PS beads is sufficient to generate the SERS effect. In order to limit the contribution of the SERS effect due to the gold nanoparticles localized onto the MPTMS, all the SERS measurements were performed with the laser beam focused onto the PS beads.

Fig. 3(a) presents the SERS spectrum of the PS beads through the blank and the Raman spectrum of PS solid for comparison. The Raman peaks of the PS solid are almost all present in the SERS spectrum of the PS beads through the blank. The unassigned peaks, in particular in the region 1500–1700 cm\(^{-1}\), can correspond to the MPTMS organic part and the H–O–H bending (1600–1630 cm\(^{-1}\)) of water. The main Raman peak of polystyrene (1004 cm\(^{-1}\)) was identified as the ring vibration. It was located with an arrow in the figure and used as an internal spectral reference in the following quantitative study.

Fig. 3(b) shows the SERS spectra of naphthalene solutions from 1 to 20 ppm. PS beads through the blank, PS beads through the blank after ethanol rinsing and for comparison the Raman spectrum of solid-state naphthalene is also given. Integration time of \( 2 \times 10 \) s.

Concerning the quantitative proof, Fig. 4 represents the normalized intensity \( I_{\text{Napht.}}/I_{\text{PS}} \) with regard to the naphthalene concentration in the range of 1–20 ppm. The normalized intensity \( I_{\text{Napht.}}/I_{\text{PS}} \) is the ratio between the band intensity of the naphthalene at 1380 cm\(^{-1}\) and the intensity of the internal spectral reference (1004 cm\(^{-1}\)). In the range of 1–20 ppm naphthalene concentration, a linear fit was plotted. Nevertheless,
The reciprocal fit allowing the use of the sensor in real conditions of application was plotted (not presented here). For example, a solution with an unknown naphthalene concentration can be determined with an uncertainty of 30% insofar as the naphthalene concentration is 12 ppm, i.e. an uncertainty of ±3.6 ppm.

For the purposes of studying the repeatability of SERS detection of naphthalene, Fig. 5 gathers the normalized intensity (I_{Napht}/I_{PS}) of a 10 ppm naphthalene solution with regard to the different analysed areas (from A to E) of the samples.

As a result, the SERS response of the sensor was quite repeatable (SD = 0.47; ca. 16%) in the naphthalene detection whatever the tested area of the sample and synthesis to synthesis. To this extent, the repeatability of the experimental measurements with the SERS substrate based on gold coated PS beads was proved. In addition, Fig. 5 displays that 11 rinsing steps with ethanol and water (one step between each spectrum) on sample 1 and 6 rinsing steps on sample 2 do not make the sensor worse according to the homogeneity of the response.

Conclusion

SERS-active substrates based on PS beads coated with gold nanoparticles were achieved with success. The morphology of the SERS sensors was described as a monolayer of PS beads partially coated with gold nanoparticles. The originality and the advantage of these SERS-active substrates reside in the pre-concentration of naphthalene onto a hydrophobic surface coupled to the use of an internal spectral reference via the Raman fingerprint of the PS beads. This study suggests that a determination of the concentration of a naphthalene solution was feasible in the range of 1–20 ppm with a high repeatability. Such SERS-active substrates tend towards an application as quantitative SERS sensors for environmental analysis. In the near future, in situ quantitative SERS measurements can be planned on hydrophobic molecules of environmental interest such as PAHs.

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