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# Design of gold nanoparticles-decorated SiO<sub>2</sub>@TiO<sub>2</sub> core/shell nanostructures for visible light-activated photocatalysis

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In this study, we designed core/shell nanostructures (CSNs) of silicon dioxide (SiO<sub>2</sub>)/titanium dioxide (TiO<sub>2</sub>), which were decorated with gold (Au) nanoparticles (NPs), to activate the visible light-driven photocatalytic reaction that is not feasible with only the SiO<sub>2</sub>@TiO<sub>2</sub> CSN because there is no absorption in the visible wavelengths. The CSNs were simply synthesized by hydrothermal method. In the core/shell structure, TiO<sub>2</sub> covered the SiO<sub>2</sub> core surface, which acts as a template. The attachment of additional Au NPs inside or outside the core/shell led to absorbance in visible wavelengths of light from the Sun, from which most radiation comes with a spectrum peak at yellow wavelength, owing to the localized surface plasmon resonance (LSPR) peak at approximately 550 nm. The SiO<sub>2</sub>@Au@TiO<sub>2</sub>@Au CSN (0.01 g) decomposed 1% of methyl orange (MO) solution in 15 mL deionized water in 1 h under the visible light. The visible light-driven photocatalytic reaction was ascribed to the LSPR around the Au NPs that were deposited inner or outer the mesoporous TiO<sub>2</sub> shell.

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## 1 Introduction

TiO<sub>2</sub> is a well-known mineral with many attractive characteristics such as nontoxicity, chemical stability to acidic and/or basic conditions, long-term stability, and good oxidizing ability.<sup>1,2</sup> Thus, many researchers have studied the usage of TiO<sub>2</sub> to extend its applications in different fields such as UV blocking pigments, self-cleaning materials, dye-sensitized solar cells, humidity sensors, lithium-ion storage, and photocatalysts.<sup>3–9</sup> TiO<sub>2</sub> NPs have been used as an effective photocatalyst to solve environmental problems by decomposing organic pollutants in water into CO<sub>2</sub> and H<sub>2</sub>O.<sup>9,10</sup> When the sun light illuminates TiO<sub>2</sub> in water, TiO<sub>2</sub> oxidizes H<sub>2</sub>O molecules into hydroxyl radicals (<sup>•</sup>OH), which can decompose the organic pollutants in water.<sup>10</sup>

However, TiO<sub>2</sub> has a limited photocatalytic activity only in the UV-light region (<400 nm) because of its wide bandgap (~3.2 eV [388 nm] for the anatase crystalline phase) and fast recombination of photo-generated electrons with holes.<sup>1</sup> UV light comprises a small fraction (approximately 4%) of the sun-light spectrum, whereas visible light comprises 44.4% intensity

of the sun-light. To use the abundant visible light as an energy source for photocatalytic activity, many different approaches have been made to narrow the TiO<sub>2</sub> bandgap for absorbing the light at visible wavelengths by doping the TiO<sub>2</sub> with transition metal dopants (tungsten, copper and iron)<sup>11–13</sup> and anionic nonmetal dopants (nitrogen and iodine).<sup>14,15</sup> However, these doping methods have many drawbacks such as a low doping concentration, reduced electron–hole lifetime, and low corrosion resistance.<sup>16,17</sup> Another method to generate the visible light responsive TiO<sub>2</sub> was achieved by using noble metals, *e.g.*, gold (Au), silver (Ag), and platinum (Pt).<sup>18–20</sup> Noble metals are successfully used because of their chemical stability and visible-light absorption due to the localized surface plasmon resonance (LSPR),<sup>21,22</sup> in which the conducting electrons on the noble metal surface undergo a collective oscillation induced by the light at a certain wavelength. Thus, at a plasmon resonant frequency the light absorption of noble metal NPs can be made. Because of this unique effect, noble metals assist the photocatalytic activity of semiconductors by absorbing the light at visible wavelengths and injecting the photo-generated electrons from the noble metal into the conduction band of the semiconductor.

Guo *et al.* reported that the Au NPs deposited at both sides of 5 nm thick TiO<sub>2</sub> sheet showed an enhanced near-field amplitude of localized surface plasmon resonance between the nearly touching Au NPs with a spacing of 5 nm, which could boost the photocatalytic activity of TiO<sub>2</sub> sheet.<sup>23</sup> Recently, Kim *et al.* synthesized Au NPs-decorated SiO<sub>2</sub>@amorphous TiO<sub>2</sub> core/shell structures and incorporated those into the photoanodes

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in dye-sensitized solar cells. Because of the plasmonic effect at the Au NPs, short-circuit current increased and power conversion efficiency was accordingly enhanced by 14%.<sup>24</sup> In addition, Hong *et al.* introduced 60 nm-sized Au NPs-attached TiO<sub>2</sub> nanofibers into an electron-transporting layer in perovskite solar cells and improved the conversion efficiency by 15% because of the LSPR effect.<sup>25</sup>

Until now, the location of Au NPs (inner or outer of the TiO<sub>2</sub> shell) and the number of Au NP layers within the CSNs, which can play a significant role in enhancing the optical properties in the visible-light region, have not been deeply studied. In this study, we designed various visible light-driven core/shell nanostructured (CSN) photocatalysts, in which the SiO<sub>2</sub> core was decorated with various arrangements of both Au NPs layer and TiO<sub>2</sub> shell. To investigate the LSPR effect of various CSNs on the photocatalytic activity, their optical properties and methyl orange (MO) decomposition abilities were compared among them, and with those of the commercially available mixed-phase of anatase and rutile TiO<sub>2</sub> (Degussa, P25).

## 2 Experimental

### 2.1 Materials and reagents

All chemical reagents were of analytical grade and used without further purification: ethanol (CH<sub>3</sub>CH<sub>2</sub>OH, anhydrous, Samchun Chem.), tetraethyl orthosilicate (TEOS, Si(OC<sub>2</sub>H<sub>5</sub>)<sub>4</sub>, 99%, Sigma-Aldrich Co., Ltd), ammonia hydroxide solution (NH<sub>4</sub>OH, 28–30%, Sigma-Aldrich Co., Ltd), titanium butoxide (TBT, Ti(OCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CH<sub>3</sub>)<sub>4</sub>, 97%, Sigma-Aldrich Co., Ltd), (3-aminopropyl)trimethoxysilane (APTMS, H<sub>2</sub>N(CH<sub>2</sub>)<sub>3</sub>Si(OCH<sub>3</sub>)<sub>3</sub>, 97%, Sigma-Aldrich Co., Ltd), gold(III) chloride hydrate (HAuCl<sub>4</sub>·xH<sub>2</sub>O, 99.999%, Sigma-Aldrich Co., Ltd), sodium citrate tribasic dehydrate (HOC(COONa)(CH<sub>2</sub>COONa)<sub>2</sub>·2H<sub>2</sub>O, 99.9%, Sigma-Aldrich Co., Ltd), and P25. Commercial MO (C<sub>14</sub>H<sub>14</sub>N<sub>3</sub>·NaO<sub>3</sub>S, 0.1%, Sigma-Aldrich Co., Ltd) was used as a target dye. All aqueous solutions were prepared with deionized (DI) water, filtered using a Milli-Q plus system ( $\rho \geq 18 \text{ M}\Omega \text{ cm}$ ).

### 2.2 Synthesis of SiO<sub>2</sub>@Au CSN

SiO<sub>2</sub> spheres were produced using the well-known Stöber method.<sup>26</sup> 0.3 mM TEOS in 80 mL ethanol as a SiO<sub>2</sub> precursor was mixed with 3 mL ammonia hydroxide and 5 mL DI water, and stirred at room temperature for 12 h. Amine functionalization at the SiO<sub>2</sub> sphere surface is necessary for the attachment of Au NPs by electrostatic interaction. 1 mL APTMS was slowly added into the prepared SiO<sub>2</sub> spheres-dispersed DI solution under mild stirring (400 rpm), and the solution was placed in a conventional oven at 70 °C for 4 h to initiate the amine functionalization at the Au NP surface. After the reaction, the amine functionalized SiO<sub>2</sub> spheres were thoroughly washed three times using a centrifuge (5000 rpm for 30 min). The collected amine-functionalized SiO<sub>2</sub> spheres were mixed with the Au NP solution, which was prepared using the Turkevich method;<sup>27</sup> 0.5 mM gold(III) chloride hydrate was added into 30 mL boiling DI water (hot-plate temperature: 300 °C) with vigorous stirring (800 rpm). 100 mM sodium citrate DI solution

was added into the boiling Au precursor solution for 15 min until the solution changed from colourless to red violet.

### 2.3 Synthesis of SiO<sub>2</sub>@Au@TiO<sub>2</sub>@Au CSN

To obtain the SiO<sub>2</sub>@Au@TiO<sub>2</sub> CSN, 15 mM TBT in ethanol was injected using a syringe pump into 50 mL ethanol solution having the dispersed SiO<sub>2</sub>@Au CSN under mild stirring (300 rpm). After adding 4 mL DI water, the solution was transferred to a Teflon autoclave and placed in an oven at 130 °C for 12 h. After cooling to room temperature, the precipitate was washed with ethanol 3 times using a centrifuge. The washed precipitate of SiO<sub>2</sub>@Au@TiO<sub>2</sub> CSN was dispersed in ethanol and mixed with APTMS for further amine functionalization on the TiO<sub>2</sub> shell surface as previously mentioned. After the amine functionalization, the Au NP solution was added into the SiO<sub>2</sub>@Au@TiO<sub>2</sub> CSN-dispersed DI solution for the final Au NP attachment to the outer shell of TiO<sub>2</sub>.

### 2.4 Photocatalytic experiments

Three different CSNs (SiO<sub>2</sub>@Au@TiO<sub>2</sub>, SiO<sub>2</sub>@TiO<sub>2</sub>@Au, and SiO<sub>2</sub>@Au@TiO<sub>2</sub>@Au) were prepared to measure the photocatalytic activity, and to compare their activity with that of the widely used P25 (TiO<sub>2</sub>, Degussa). The photocatalytic activity was measured in ambient atmosphere at room temperature by degrading the methyl orange (MO) solution with illumination with an AM 1.5 simulated sunlight source (SANNEI solar simulator, Class A). The UV light ( $\lambda < 400 \text{ nm}$ ) was cut off using a UV shield film (SK-2, Sunnano) to observe the photocatalytic activity under only visible light at a power density of  $80 \pm 2.5 \text{ mW cm}^{-2}$ . Each CSN powder (0.01 g) was added into 20 mL DI water with 1 vol% of MO. The CSN suspensions were vigorously stirred in the dark for 30 min to reach adsorption/desorption equilibrium of MO molecules on the CSN surface. At every 15 min, 1 mL suspension was sampled and filtered to measure the resident MO concentration using a UV-visible spectrometer (AvaSpec-ULS2048L-USB2 Spectrometer, Jinyoung tech Inc.).

## 3 Result and discussion

### 3.1 Structure observation

We synthesized the SiO<sub>2</sub>@Au@TiO<sub>2</sub>@Au CSN by coating the Au NPs and TiO<sub>2</sub> on the SiO<sub>2</sub> core in sequence as depicted in Fig. 1. The amine (NH<sup>3+</sup>) functionalization facilitated the attachment of Au NPs onto the both SiO<sub>2</sub> core and TiO<sub>2</sub> shell surface. Therefore, uniform amine functionalization was the most important step to achieve the dense coverage of Au NPs. To obtain the uniform amine functionalization, dispersion of the SiO<sub>2</sub> spheres in ethanol at a low concentration was necessary. The Au NPs prepared using the Turkevich method were surrounded with negatively charged citrate ions (C<sub>3</sub>H<sub>5</sub>O(COO)<sub>3</sub><sup>3-</sup>), which improved the dispersion and stabilization of Au NPs in the solution owing to the electrostatic repulsion between them. Meanwhile, those negatively charged citrate ions electrostatically attracted the positively charged NH<sup>3+</sup> functionalized SiO<sub>2</sub> spheres to form the Au NPs coating. Scanning electron microscope (FE-SEM, JEOL 2010F) images of the SiO<sub>2</sub>, SiO<sub>2</sub>@Au,



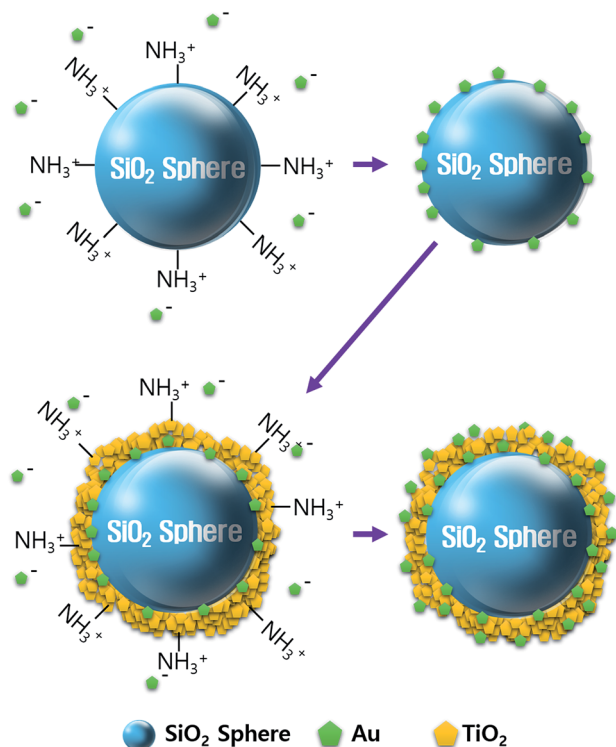


Fig. 1 Schematic representation of the  $\text{SiO}_2\text{@Au@TiO}_2\text{@Au}$  CSN synthesis.

$\text{SiO}_2\text{@Au@TiO}_2$ , and  $\text{SiO}_2\text{@Au@TiO}_2\text{@Au}$  CSNs are shown in Fig. 2a–d. Fig. 2a shows the 200 nm-sized  $\text{SiO}_2$  core with a smooth surface. In contrast, the  $\text{SiO}_2\text{@Au}$  has a rough surface with the Au NPs (Fig. 2b). The high-resolution image of it (inset of Fig. 2b) shows the average size of Au NPs of 15 nm. Fig. 2c shows two  $\text{SiO}_2\text{@Au@TiO}_2$  CSNs after complete coverage with the  $\text{TiO}_2$  shell. In this case, the Au NPs are invisible after coating the 30 nm-thick  $\text{TiO}_2$  shell, which was composed of a few nanometer-sized  $\text{TiO}_2$  NPs with a relatively smooth  $\text{SiO}_2\text{@Au@TiO}_2$  CSN surface.

The  $\text{TiO}_2$  shell was formed by two different mechanisms above 130 °C: intermediate titanium hydroxide ( $\text{Ti}(\text{OH})_4$ ) was formed in the hydrolysis reaction (i) and converted into  $\text{TiO}_2$  by condensation reaction (ii) or TBT decomposition by reaction with the intermediate  $\text{Ti}(\text{OH})_4$  into  $\text{TiO}_2$  and by-product of butanol (iii):<sup>28</sup>

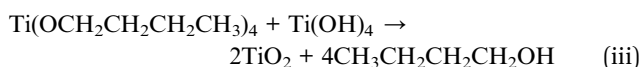
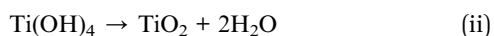
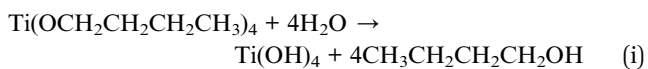


Fig. 2d shows the Au NPs that were coated onto the entire surface of a  $\text{SiO}_2\text{@Au@TiO}_2$  CSN. For the Au NP attachment, the  $\text{NH}_3^+$  functionalization treatment was also performed to the  $\text{TiO}_2$  shell. Fig. 2e shows a high-angle annular dark-field (HAADF) image of the  $\text{SiO}_2\text{@Au@TiO}_2$  CSN and the

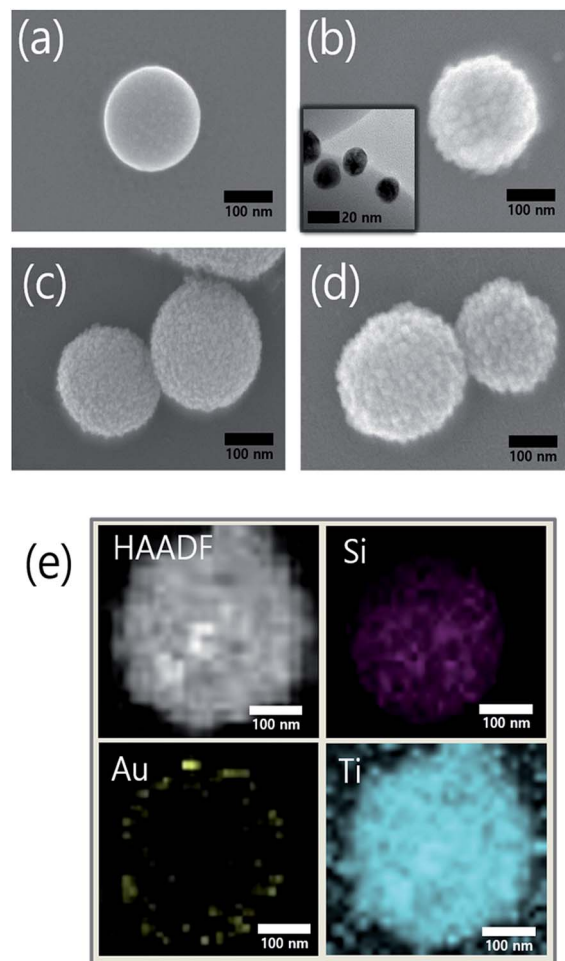


Fig. 2 SEM images of (a) bare  $\text{SiO}_2$ , (b)  $\text{SiO}_2\text{@Au}$ , (c)  $\text{SiO}_2\text{@Au@TiO}_2$ , and (d)  $\text{SiO}_2\text{@Au@TiO}_2\text{@Au}$ . (e) HAADF-STEM image and the corresponding EDS element mapping images of silicon (Si, purple), gold (Au, yellow) and titanium (Ti, skyblue) for the  $\text{SiO}_2\text{@Au@TiO}_2$  CSN. The inset of (c) is the high-resolution TEM image of Au NPs on the  $\text{SiO}_2$  sphere surface.

corresponding elemental mappings of Si, Au and Ti. It is shown that the Au NPs coated the  $\text{SiO}_2$  core surface and were then thoroughly covered by the  $\text{TiO}_2$  shell.

Fig. 3 shows the X-ray diffraction (XRD) patterns from three different samples using Cu K-alpha radiation (40 kV, 100 mA, Rigaku D/max-2400). The broad XRD pattern between 20° and 30° comes from the amorphous  $\text{SiO}_2$  sphere (JCPDS 29-0085, ICSD data).<sup>29</sup> The  $\text{SiO}_2\text{@Au}$  powder revealed two sharp peaks at 38.18° and 44.38°, which correspond to the Au cubic crystal (111) and (200) peaks, respectively (JCPDS 65-2870).<sup>30</sup> The diffraction peaks of the anatase  $\text{TiO}_2$  crystal planes of (101), (200), (105) and (211) were observed from the  $\text{SiO}_2\text{@Au@TiO}_2\text{@Au}$  powder (JCPDS-521-1272, ICSD data),<sup>31</sup> which were not shown in the  $\text{SiO}_2\text{@Au}$  diffraction pattern. These peaks indicate that the synthesized  $\text{TiO}_2$  shell had anatase crystallinity, which has been reported to show the best photocatalytic activity among various  $\text{TiO}_2$  crystallinities (rutile, brookite, and anatase).<sup>32</sup> Only Au,  $\text{TiO}_2$ , and  $\text{SiO}_2$  diffraction





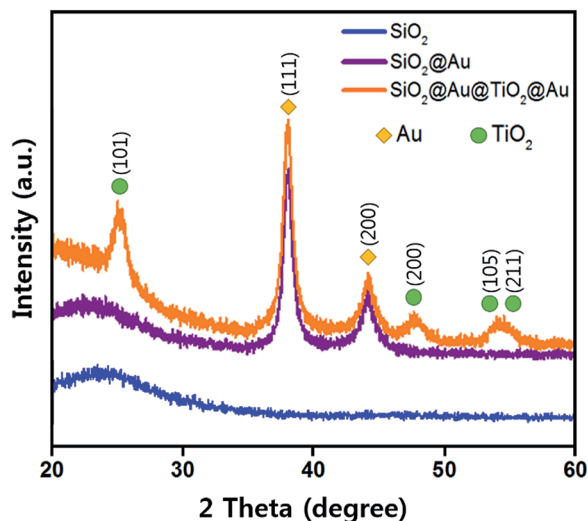


Fig. 3 X-ray diffraction patterns of as-prepared  $\text{SiO}_2$ ,  $\text{SiO}_2@Au$  and  $\text{SiO}_2@Au@TiO_2@Au$  CSNs.

peaks were observed, demonstrating that no impurities remained in the synthesized CSNs.

### 3.2 Optical characteristics of core/shell materials

The UV-Vis. absorption spectra of ethanol solutions (4 mL) that contained 0.001 g of each synthesized  $\text{SiO}_2@TiO_2$ ,  $\text{SiO}_2@Au@TiO_2$ ,  $\text{SiO}_2@TiO_2@Au$ , and  $\text{SiO}_2@Au@TiO_2@Au$  CSN are shown in Fig. 4. The four  $\text{SiO}_2@TiO_2$ -based CSNs have an absorption in the UV range and absorption edge at 313 nm. After the Au NPs attachment, the  $\text{SiO}_2@TiO_2@Au$  sample had an absorption peak at 550 nm, which is ascribed to the collective oscillation of metal conduction electrons driven by a specific wavelength (LSPR). This peak was red-shifted to 572 nm in the  $\text{SiO}_2@Au@TiO_2$  sample. Because the LSPR is sensitive to the complex dielectric environments,<sup>33</sup> the number of neighboring Au NPs,<sup>34</sup> and interparticle spacing of Au NPs,<sup>35</sup> the LSPR peak from the Au NPs can shift depending on their arrangement and location. It is reported that when the interparticle spacing becomes smaller lithographically, the

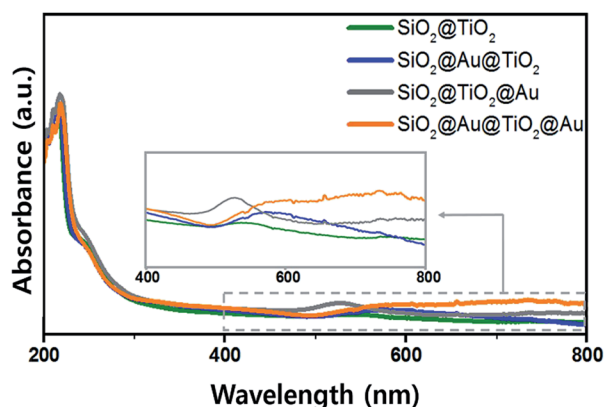


Fig. 4 UV-visible absorption spectra of various  $\text{SiO}_2@TiO_2$  CSNs.

absorption peak from the Au NPs is broadened and shifted to the red.<sup>35</sup> In our case, when the Au NPs were located at both the inner and outer  $TiO_2$  shell, the LSPR absorption band was interestingly broadened through visible wavelengths and even extended to the infrared region, which could be caused by the plasmonic coupling between the two Au NPs layers with a gap of 30 nm-thick  $TiO_2$  dielectric shell. We made three identical sample sets, showing the similar absorption spectra tendency among the samples.

### 3.3 Photocatalytic activity of core/shell materials under visible light

In order to test the visible light-driven photocatalytic ability of the synthesized CSNs, the MO solution was photocatalytically degraded by illuminating the MO solution containing the dispersed CSN powders with visible light. MO is widely used by chemists to indicate the titration of acid by observing its colour change below pH 3.1. Because of its mutagenic properties and acidity (pH 3.47), direct contact should be avoided for health concern.<sup>36</sup> Therefore, the decomposition of MO is required for health concern and environment safety.

At first, the MO solution without any photocatalyst was irradiated to check the photolysis of MO itself under visible wavelengths with a simulated sunlight source at an intensity of  $80 \pm 2.5 \text{ mW cm}^{-2}$ . But, no photolytic reaction of the only MO solution to the visible light was observed. Fig. 5a–d compare the absorbance variations of MO after photodegradation under visible light. Compared to the commercially available P25, all Au-decorated CSN samples obviously showed enhanced visible light-driven photocatalytic activities. The intensity of MO absorption peak ( $\lambda = 485 \text{ nm}$ ) decreased with the irradiation time. Meanwhile, in the case of P25 sample, the MO was not decomposed at all with the irradiation time because the visible light cannot provide sufficient energy to excite the electrons over its large bandgap to generate electron–hole pairs. However, in the case of Au NPs-coated CSN samples ( $\text{SiO}_2@Au@TiO_2$ ,  $\text{SiO}_2@TiO_2@Au$ , and  $\text{SiO}_2@Au@TiO_2@Au$ ), electrons are generated under visible light because of the LSPR at the Au NPs and transferred to the conduction band of  $TiO_2$  within femto seconds as shown in Fig. 6.<sup>37</sup> The transferred electrons can generate hydroxyl radicals ( $\cdot OH$ ) at the  $TiO_2$ /water interface,<sup>38,39</sup> which eventually decompose the MO in the solution into non-hazardous by-products. Meanwhile, after electrons' transfer to the  $TiO_2$ , the holes remained in the Au NPs oxidized the interfacial water to produce the hydroxyl radicals, which additionally decomposed the MO molecules.<sup>40–44</sup>

The  $\text{SiO}_2@Au@TiO_2@Au$  sample had the best photocatalytic performance among the CSN samples. In this case, both sides of the  $TiO_2$  shell were coated by the Au NPs, which generated more visible light-driven electrons and consequently hydroxyl radicals at the  $TiO_2$ /water interface.

Fig. 5e shows the degradation efficiency of the photocatalyst by measuring the change in MO concentration ( $C/C_0$ ) with irradiation time ( $C_0$  is the initial concentration and  $C$  is the MO concentration after a certain irradiation time). The degradation efficiency is defined as follows:



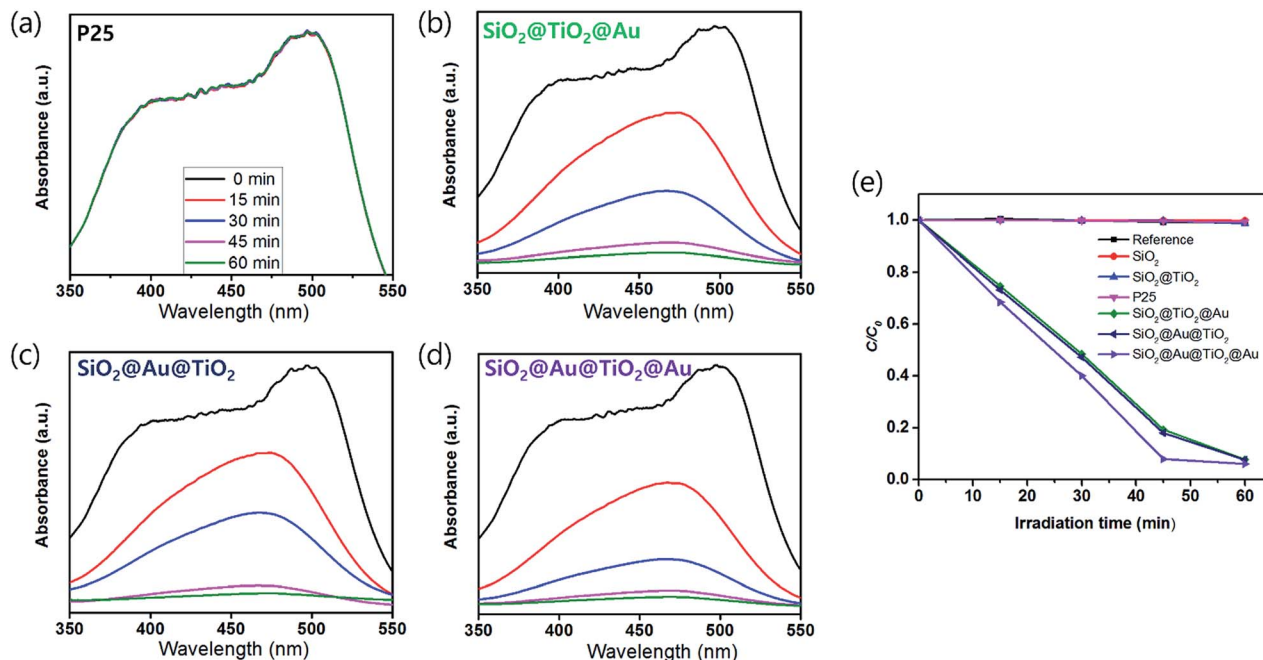


Fig. 5 UV-vis absorption spectra showing the degradation of MO solution with varying irradiation time under visible light by (a) P25, (b)  $\text{SiO}_2/\text{TiO}_2/\text{Au}$ , (c)  $\text{SiO}_2/\text{Au}/\text{TiO}_2$ , and (d)  $\text{SiO}_2/\text{Au}/\text{TiO}_2/\text{Au}$ . (e) A comparison of the photocatalytic activity of reference,  $\text{SiO}_2$ ,  $\text{SiO}_2/\text{TiO}_2$ , the Au NPs-decorated CSN photocatalysts and P25 under the visible light.

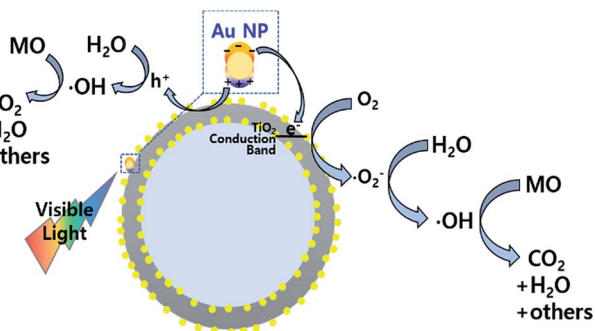


Fig. 6 Photocatalytic mechanism of the Au NPs-decorated  $\text{SiO}_2/\text{Au}/\text{TiO}_2$  CSN under the visible light irradiation.

$$\text{Degradation (\%)} = (1 - C/C_0) \times 100\%$$

There was no photocatalytic activity with the  $\text{SiO}_2$ ,  $\text{SiO}_2/\text{TiO}_2$  and P25 samples, the Au-decorated CSNs decomposed nearly the entire MO quantity in an hour under visible light, illustrating that the visible light was absorbed at the Au NPs due to LSPR. The photodegradation rate is expressed as follows:<sup>45</sup>

$$\ln(C/C_0) = -kt$$

where,  $k$  ( $\text{min}^{-1}$ ) is the degradation rate constant and is calculated by fitting the experimental data curve with an exponential function. The  $k$  values of the  $\text{SiO}_2/\text{TiO}_2/\text{Au}$ ,  $\text{SiO}_2/\text{Au}/\text{TiO}_2$ , and  $\text{SiO}_2/\text{Au}/\text{TiO}_2/\text{Au}$  CSNs are  $4.74 \times 10^{-2}$ ,  $4.45 \times 10^{-2}$ , and  $5.69 \times 10^{-2}$ , respectively. Considering the 30 nm-thick  $\text{TiO}_2$

shell around the 200 nm diameter  $\text{SiO}_2$ , the amount of  $\text{TiO}_2$  material is minimal (roughly only 0.0017 g in each suspension sampling, which was simply calculated from the volume ratio and density of the  $\text{SiO}_2$  and  $\text{TiO}_2$  with ignorance of the weight of Au NPs). However, in comparison to the P25 sample, the Au-decorated CSN samples ( $\text{SiO}_2/\text{Au}/\text{TiO}_2$ ,  $\text{SiO}_2/\text{TiO}_2/\text{Au}$ , and  $\text{SiO}_2/\text{Au}/\text{TiO}_2/\text{Au}$  CSN) demonstrated an active photocatalytic ability under the visible light.

### 3.4 Porosity study of $\text{SiO}_2/\text{TiO}_2$ core/shell

To evidence the infiltration of MO solution to the inner Au NPs layer, the porosity of  $\text{TiO}_2$  shell was checked by the Brunauer–Emmett–Teller (BET) and Barrett–Joyner–Halenda (BJH) methods. A TEM image of Fig. 7a depicts the  $\text{SiO}_2$  spheres coated with the mesoporous  $\text{TiO}_2$  shell, which is composed of a few nm-sized nanoparticles as shown in inset figure. Adsorption/desorption of nitrogen ( $\text{N}_2$ ) analysis was performed with a ASAP 2020 (Micromeritics, USA) at 77.30 K. Prior to performing the BET experiment, the  $\text{SiO}_2$  and  $\text{SiO}_2/\text{TiO}_2$  powder samples were degassed at 150 °C. In Fig. 7b, the  $\text{N}_2$  gas adsorption/desorption isotherm curves for  $\text{SiO}_2$  and  $\text{SiO}_2/\text{TiO}_2$  samples follow the type (IV) isotherm curve with a hysteresis loop, representing the mesoporosity of both samples.<sup>46</sup> The BET surface area of  $\text{SiO}_2/\text{TiO}_2$  and  $\text{SiO}_2$  were  $12.83 \text{ m}^2 \text{ g}^{-1}$  and  $134.88 \text{ m}^2 \text{ g}^{-1}$ , respectively. Hence, the  $\text{SiO}_2/\text{TiO}_2$  powder absorbed much larger amount of  $\text{N}_2$  gas than that of the  $\text{SiO}_2$  powder owing to the mesoporous  $\text{TiO}_2$  shell. In comparison, the previously synthesized  $\text{SiO}_2/\text{TiO}_2$  core/shell had a surface area of only  $14.6 \text{ m}^2 \text{ g}^{-1}$ .<sup>47</sup> Owing to the higher  $\text{TiO}_2$  shell mesoporosity and closely packed Au NPs of our core/shell structure, it

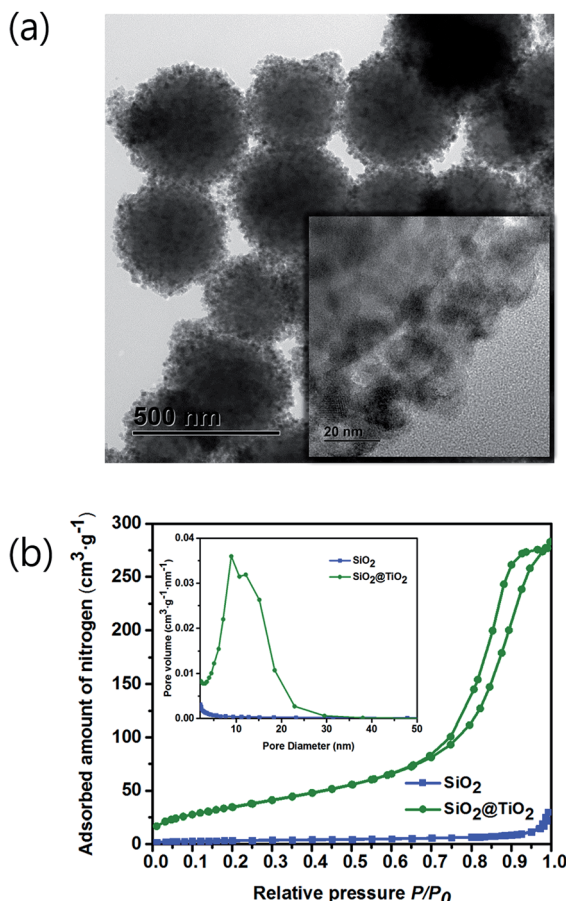


Fig. 7 (a) TEM image of  $\text{SiO}_2@\text{TiO}_2$ . Inset is the high-resolution TEM image, showing the coverage of  $\text{SiO}_2$  core surface with the  $\text{TiO}_2$  NPs. (b)  $\text{N}_2$  adsorption/desorption isotherms of  $\text{SiO}_2$  and  $\text{SiO}_2@\text{TiO}_2$ . Inset shows the corresponding pore size distributions.

could show a better photocatalytic activity to decompose more pollutants.

The desorption isotherm was used to estimate the BJH pore-size distribution. The average pore diameters of  $\text{SiO}_2$  and  $\text{SiO}_2@\text{TiO}_2$  were calculated to be 7.10 nm and 12.55 nm, respectively. Inset image in Fig. 7b shows that the  $\text{TiO}_2$  shell has a larger pore size and a pore volume than those of  $\text{SiO}_2$  core. Therefore, the MO solution could infiltrate into the inner Au NPs layer through the mesoporous  $\text{TiO}_2$  shell and be decomposed by the visible light irradiation.

## 4 Conclusions

$\text{SiO}_2@\text{TiO}_2$  core/shell nanostructures (CSNs) were hydrothermally synthesized to explore them as photocatalysts for the degradation of the hazardous pollutants in water. However, the visible wavelengths of the Sun cannot excite the  $\text{TiO}_2$  to generate electrons under visible light because of its large bandgap. Thus, Au NPs were attached to the inner or outer  $\text{TiO}_2$  shell by amine functionalization and electrostatic interaction. The Au NPs uniformly covered the  $\text{SiO}_2$  core and the  $\text{TiO}_2$  shell, producing the  $\text{SiO}_2@\text{Au}@\text{TiO}_2@\text{Au}$  CSN. The Au NPs-decorated CSNs

revealed light absorption in visible wavelengths because of the localized surface plasmonic resonance at the Au NPs and even in infrared wavelengths in the case of  $\text{SiO}_2@\text{Au}@\text{TiO}_2@\text{Au}$  CSN. Therefore, the Au NPs-decorated CSNs decomposed MO solution under the visible light. The plasmonic effect of Au NPs led to electron generation under the visible light, which could not occur for P25. In 1 h, 0.01 g Au NPs-decorated CSNs ( $\text{SiO}_2@\text{Au}@\text{TiO}_2$ ,  $\text{SiO}_2@\text{TiO}_2@\text{Au}$ ,  $\text{SiO}_2@\text{Au}@\text{TiO}_2@\text{Au}$ ) decomposed 1 vol% MO in 20 mL DI water.

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## Notes and references

- 1 R. Asashi, T. Morikawa, T. Ohwaki, K. Aoki and Y. Taga, *Science*, 2001, **293**, 269.
- 2 S. M. Gupta and M. Tripathi, *Chin. Sci. Bull.*, 2011, **56**, 1639.
- 3 H. Yang, S. Zhu and N. Pan, *J. Appl. Polym. Sci.*, 2004, **92**, 3201.
- 4 H. Yaghoubi, N. Taghavinia and E. K. Alamdari, *Surf. Coat. Technol.*, 2010, **204**, 1562.
- 5 A. Burke, S. Ito, H. Snaith, U. Bach, J. Kwiakowski and M. Grätzel, *Nano Lett.*, 2008, **8**, 977.
- 6 P.-G. Su and L. N. Huang, *Sens. Actuators, B*, 2007, **123**, 501.
- 7 M.-C. Yang, Y.-Y. Lee, B. Xu, K. Powers and Y. S. Meng, *J. Power Sources*, 2012, **207**, 166.
- 8 C.-C. Lin, J.-W. Liao and W.-Y. Li, *Ceram. Int.*, 2013, **39**, S733.
- 9 K. Nakata and A. Fujishima, *J. Photochem. Photobiol., C*, 2012, **13**, 169.
- 10 J. Schneider, M. Matsuoka, M. Takeuchi, J. Zhang, Y. Horiuchi, M. Anpo and D. W. Bahnemann, *Chem. Rev.*, 2014, **114**, 9919.
- 11 N. Couselo, E. F. S. García, R. J. Candal and M. Jobbágy, *J. Phys. Chem. C*, 2008, **112**, 1094.
- 12 L. S. Yoong, F. K. Chong and B. K. Dutta, *Energy*, 2009, **34**, 1652.
- 13 K. C. Christoforidis, A. Iglesias-Juez, S. J. A. Figueroa, M. D. Michiel, M. A. Newton and M. Fernández-García, *Catal. Sci. Technol.*, 2013, **3**, 626.
- 14 G. Yang, Z. Jiang, H. Shi, T. Xiao and Z. Yan, *J. Mater. Chem.*, 2010, **20**, 5301.
- 15 Q. Zhang, Y. Li, E. A. Ackerman, M. Gajdardziska-Josifovska and H. Li, *Appl. Catal., A*, 2011, **400**, 195.
- 16 Y. Wang, H. Cheng, Y. Hao, J. Ma, W. Li and S. Cai, *Thin Solid Films*, 1999, **349**, 120.
- 17 H. Yamashita, M. Honda, M. Harada, Y. Ichihashi, M. Anpo, T. Hirao, N. Itoh and N. Iwamoto, *J. Phys. Chem. B*, 1998, **102**, 10707.
- 18 A. Pandikumar, S. Murugesan and R. Ramaraj, *ACS Appl. Mater. Interfaces*, 2010, **2**, 1912.
- 19 D.-H. Yu, X. Yu, C. Wang, X.-C. Liu and Y. Xing, *ACS Appl. Mater. Interfaces*, 2012, **4**, 2781.



- 20 H. Chen, S. Chen, X. Quan, H. Yu, H. Zhao and Y. Zhang, *J. Phys. Chem. C*, 2008, **112**, 9285.
- 21 P. K. Jain, X. Huang, I. H. El-Sayed and M. A. El-Sayed, *Acc. Chem. Res.*, 2008, **41**, 1578.
- 22 T. Y. Jeon, D. J. Kim, S. G. Park, S. H. Kim and D. H. Kim, *Nano Convergence*, 2016, **3**, 18.
- 23 H. Wang, T. You, W. Shi, J. Li and L. Guo, *J. Phys. Chem. C*, 2012, **116**, 6490.
- 24 Y. H. Jang, Y. J. Jang, S. T. Kochuveedu, M. Byun, Z. Lin and D. H. Kim, *Nanoscale*, 2014, **6**, 1823.
- 25 S. S. Mali, C. S. Shim, H. Kim, P. S. Patil and C. K. Hong, *Nanoscale*, 2016, **8**, 2664.
- 26 W. Stöber, A. Fink and E. Bohn, *J. Colloid Interface Sci.*, 1968, **26**, 62.
- 27 J. Turkevich, P. C. Stevenson and J. Hillier, *Discuss. Faraday Soc.*, 1951, **11**, 55.
- 28 H. Li, S. G. Sunol and A. K. Sunol, *Nanotechnology*, 2012, **23**, 294012.
- 29 H. Wang, M. Yu, C. K. Lin and J. Lin, *J. Colloid Interface Sci.*, 2006, **300**, 176.
- 30 L. Shang, F. Zhao and B. Zeng, *Electroanalysis*, 2013, **25**, 453.
- 31 W. Huang, X. Tang, Y. Wang, Y. Koltypin and A. Gedanken, *Chem. Commun.*, 2000, **48**, 1415.
- 32 J. Zhang, P. Zhou, J. Liu and J. Yu, *Phys. Chem. Chem. Phys.*, 2014, **16**, 20382.
- 33 K. L. Kelly, E. Coronado, L. L. Zhao and G. C. Schatz, *J. Phys. Chem. B*, 2003, **107**, 668.
- 34 Y. Yan, H. Shan, M. Li, S. Chen, J. Liu, Y. Cheng, C. Ye, Z. Yang, X. Lai and J. Hu, *Sci. Rep.*, 2015, **5**, 16715.
- 35 C. Mu, J. P. Zhang and D. Xu, *Nanotechnology*, 2009, **21**, 015604.
- 36 Y. Badr and M. A. Mahmoud, *J. Phys. Chem. Solids*, 2007, **68**, 413.
- 37 A. Furube, L. Du, K. Hara, R. Katoh and M. Tachiya, *J. Am. Chem. Soc.*, 2007, **129**, 14852.
- 38 Q. Zhang, D. Q. Lima, I. Lee, F. Zaera, M. Chi and Y. Yin, *Angew. Chem.*, 2011, **123**, 7226.
- 39 Z. Bian, J. Zhu, F. Cao, Y. Lu and H. Li, *Chem. Commun.*, 2009, **25**, 3789.
- 40 J. Xue, S. Ma, Y. Zhou and Q. Wang, *RSC Adv.*, 2015, **5**, 88249.
- 41 Y. Tian and T. Tatsuma, *J. Am. Chem. Soc.*, 2005, **127**, 7632.
- 42 S. A. Ansari, M. M. Khan, M. O. Ansari and M. H. Cho, *New J. Chem.*, 2015, **39**, 4708.
- 43 S. T. Kochuveedu, Y. H. Jang and D. H. Kim, *Chem. Soc. Rev.*, 2013, **42**, 8467.
- 44 C. Wang and D. Astruc, *Chem. Soc. Rev.*, 2014, **43**, 7188.
- 45 W. Xia, H. Wang, X. Zeng, J. Han, J. Zhu, M. Zhou and S. Wu, *CrystEngComm*, 2014, **16**, 6841.
- 46 K. S. W. Sing, D. H. Everett, R. A. W. Haul, L. Moscou, R. A. Pierotti, J. Rouquerol and T. Siemieniowska, *Pure Appl. Chem.*, 1985, **57**, 603.
- 47 M. Ye, H. Zhou, T. Zhang, Y. Zhang and Y. Shao, *Chem. Eng. J.*, 2013, **226**, 209.

