Synthesis of Au or Pt@Perovskite Nanocrystals via Interfacial Photoreduction

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Abstract: The surface modification of perovskite nanocrystals (NCs) (i.e., their decoration with noble metals) holds great promise with respect to the tailoring of their properties but has remained a challenge because perovskite NCs are extremely sensitive to water and alcohols. In this study, Au or Pt@CsPbBr 3 NCs were successfully synthesized by photoreduction at the water/hexane interface. First, Cs 3 PbBr 6 NCs were synthesized through the hot-injection method. Then, Cs 3 PbBr 6 was transformed into CsPbBr 3 and subjected to noble metal modification, both at the interface. The synthesized CsPbBr 3 NCs exhibited a cubic perovskite phase and had an average size of approximately 13.5 nm. The deposited Au and Pt nanoparticles were crystalline, with a face-centered cubic lattice and average diameters of approximately 3.9 and 4.4 nm, respectively. The noble metal modification process had almost no effect on the steady-state photoluminescence (PL) emission wavelength but affected the charge-recombination kinetics of the CsPbBr 3 NCs. Time-resolved PL decay spectral analysis indicated that the fluorescence lifetimes of the Au and Pt@CsPbBr 3 NCs were shorter than those of the pure CsPbBr 3 NCs, probably owing to the quenching of the free charges because of electron transfer from the perovskite to the noble metal nanoparticles.

Keywords: perovskite nanocrystals; noble metal; interface; photoreduction

1. Introduction

Organic–inorganic lead halide perovskite (OIHP) materials with a general ABX 3 (A = organic cation, B = metal cation, and X = halide anion) formula have attracted significant attention in the field of optoelectronics because of their unique optical and semiconducting characteristics [1−3]. For instance, their use has rapidly increased the power conversion efficiency of solar cells from 3.81% [4] to more than 25% [5] within a few years. However, owing to the instability of OIHP, including its extreme sensitivity to oxygen and moisture, as well as its poor photo- and thermostability [6], a number of perovskite analogues have been developed. Replacing the A-site organic groups with inorganic cations to construct all-inorganic halide perovskites (IHPs) is a potential strategy for improving the stability of OIHP while maintaining its optical and electrical properties [7].

Cesium lead halide perovskite (CsPbX 3) nanocrystals (NCs), an IHP material, show great promise for use in a range of fields owing to their high photoluminescence quantum yield, narrow emission width, and tunable band gap, which covers the entire visible range [8]. They were first reported by the Kovalenko group in 2015 [9]. Since then, considerable progress has been made in the synthesis of CsPbX 3 NCs. For instance, the hot-injection method [10,11], the solvothermal method [12], ultrasonication [13], room-temperature precipitation [14], and chemical vapor deposition [15] have been employed to prepare CsPbX 3 NCs with controllable shapes and compositions.
Although the advances made in the synthesis of CsPbX$_3$ NCs have been impressive, it is known that heterostructured NCs formed by tailoring the properties of two or more dissimilar materials usually exhibit several interesting functionalities. An example is the surface modification of CsPbX$_3$ NCs at the individual particle level [16,17]. However, it remains a challenge to use conventional sol-gel methods for surface modification of CsPbX$_3$ NCs because of their water and alcohol intolerance. Recently, Hu et al. demonstrated an effective sol-gel process for modifying the surfaces of CsPbX$_3$ NCs at the interface of water and a nonpolar solvent and were able to produce CsPbX$_3$/metal oxide Janus NCs with improved stability [17]. Thus, interfacial synthesis is the key to preparing heterostructured NCs and is different from the other reported methods for preparing CsPbX$_3$/metal [18,19] and CsPbX$_3$/SiO$_2$ [20] heterostructures. Inspired by their report, herein, we performed photoreduction at the water/hexane interface to produce noble metal@CsPbBr$_3$ NCs. First, Cs$_4$PbBr$_6$ NCs were synthesized through the hot-injection method. Subsequently, the Cs$_4$PbBr$_6$ was transformed into CsPbBr$_3$ and subjected to noble metal modification, both at the interface. Owing to the localized surface plasmon resonance effect [21], high catalytic activity [22], and ease of charge separation [23] of the noble metal nanoparticles, the noble metal@CsPbX$_3$ NCs have the potential to play an important role in photodetectors, light-emitting diodes (LEDs), solar cells, and photocatalysts.

2. Results and Discussion

We propose the following mechanism to explain the synthesis process (Figure 1). When the hexane suspension containing the Cs$_4$PbBr$_6$ NCs comes in contact with water, the Cs$_4$PbBr$_6$ is transformed into CsPbBr$_3$, which accompanies the stripping of CsBr through the hexane/water interface and removing partial hydrophobic capping ligands (i.e., oleic acid (OA) and oleylamine (OAm)) [17]. Subsequently, the noble metal precursor in contact with the interface reacts with the electrons photogenerated from CsPbBr$_3$ (methanol was used as a sacrificial agent). According to a previous report [8], the CsPbBr$_3$ NCs obtained through this water-triggered transformation process exhibit enhanced stability against moisture compared with those formed through the hot-injection method. Thus, Cs$_4$PbBr$_6$ NCs were chosen to prepare the Au or Pt@CsPbBr$_3$ NCs instead of CsPbBr$_3$ NCs presynthesized by the hot-injection method.

![Figure 1](image-url)  
**Figure 1.** Diagram of transformation of Cs$_4$PbBr$_6$ into Au or Pt@CsPbBr$_3$ at the water/hexane interface. NCs—nanocrystals.
To explore the morphologies of the synthesized Cs$_4$PbBr$_6$ and CsPbBr$_3$ NCs, transmission electron microscopy (TEM) imaging was performed. Figure 2a shows that the original Cs$_4$PbBr$_6$ NCs are quasispherical with an average diameter of approximately 13.9 nm. The high-resolution TEM (HRTEM) image in the inset clearly shows that the lattice spacing is 0.122 nm and is in good agreement with that of the (1 0 1) plane of rhombohedral Cs$_4$PbBr$_6$. The size distribution of the NCs was relatively narrow (Figure 2b), indicating that the NC growth process was well controlled. After the water-triggered transformation process, the NCs show a cube-like structure with an average edge length of approximately 13.5 nm. Further, the $d$-spacing of 0.152 nm, as measured using the HRTEM image (Figure 2c and its inset), could be assigned to the (1 1 1) plane of cubic CsPbBr$_3$. The size distribution of these NCs was similar to that of the Cs$_4$PbBr$_6$ NCs (Figure 2d).

Figure 2. TEM images and size distributions of Cs$_4$PbBr$_6$ (a) and CsPbBr$_3$ NCs (b).

The transformation was also monitored through ultraviolet (UV)-visible (vis) absorption and photoluminescence (PL) measurements. As shown in Figure 3a, two sharp peaks were observed at 230 and 314 nm (solid black line). These could be assigned to the pristine Cs$_4$PbBr$_6$ NCs [24]. The spectrum shows that the Cs$_4$PbBr$_6$ NC suspension did not exhibit any distinct absorption peaks in the visible-light region, in accordance with the colorless appearance of the suspension. After the reaction with water, the sharp peak at 314 nm disappeared and a new absorption peak emerged at approximately 510 nm (solid red line), indicating the transformation of Cs$_4$PbBr$_6$ into CsPbBr$_3$. The corresponding PL spectrum shows a sharp peak at 532 nm, with a small full-width-at-half-maximum value. Figure 3b shows the X-ray diffraction (XRD) pattern of the original Cs$_4$PbBr$_6$ NCs, as well as that of the product after the water-triggered transformation. The Cs$_4$PbBr$_6$ NCs exhibit...
a rhombohedral phase, and the diffraction peaks at approximately 22.4°, 27.8°, 28.7°, 29.0°, and 30.4° correspond to the (101), (610), (230), (121), and (330) planes, respectively [17]. On the other hand, the product formed after the water treatment exhibits a pure cubic CsPbBr₃ perovskite phase, and its diffraction pattern contains peaks related to the (100), (110), and (200) planes [17]. These results confirm that a change in the crystal structure from Cs₄PbBr₆ to CsPbBr₃ was successfully realized at the water/hexane interface.

Figure 3. (a) Absorption and photoluminescence (PL) spectra and (b) X-ray diffraction (XRD) patterns of Cs₄PbBr₆ and CsPbBr₃ NCs. The background of XRD patterns in (b) is shifted because of artificial translation of Y-axis.

Figure 4 shows the TEM images of the Au or Pt@CsPbBr₃ hybrid NCs prepared by reacting Cs₄PbBr₆ NCs with the corresponding noble metal precursor under light
irradiation at the water/hexane interface. These hybrid NCs were similar in shape and size to the pure CsPbBr$_3$ NCs shown in Figure 1c. Moreover, the Au and Pt nanoparticles were deposited randomly on the surfaces of the CsPbBr$_3$ NCs, and their sizes were relatively small, at 3.9 and 4.4 nm, respectively. The HRTEM images show that the formed Au and Pt nanoparticles are crystalline with $d$-spacings of 0.106 and 0.090 nm, respectively, which correspond to the face-centered cubic (fcc) Au (400) (JCPDS Card No. 04–0784) and Pt (331) planes (JCPDS Card No. 70–2057), respectively. These results confirmed that the photoreduction reaction had been carried out successfully at the water/hexane interface. The X-ray photoelectron spectroscopy (XPS) survey scans of Au@CsPbBr$_3$ and Pt@CsPbBr$_3$ NCs are presented in the left of Figure 5, in which the presence of Cs, Pb, Br, and Au (or Pt) elements is confirmed. In the high-resolution XPS spectra (the right part of Figure 5), doublet peaks containing a low energy band (Au4f$_{7/2}$) and a high energy band (Au4f$_{5/2}$) are observed at 84.0 and 87.7 eV for Au@CsPbBr$_3$, respectively, and 71.0 (Pt4f$_{7/2}$) and 74.3 eV (Pt4f$_{5/2}$) for Pt@CsPbBr$_3$, which are assigned to metallic Au(0) or Pt(0) [25,26]. The results demonstrate that the noble metal precursors can be effectively reduced through the interfacial photoreduction.

![TEM images](image)

**Figure 4.** TEM images of (a) Au@CsPbBr$_3$ and (b) Pt@CsPbBr$_3$ NCs.
Figure 5. X-ray photoelectron spectroscopy (XPS) data of (a) Au@CsPbBr$_3$ and (b) Pt@CsPbBr$_3$ NCs.

Successful deposition of the Au or Pt nanoparticles was also verified through UV-vis absorption measurements. Figure 6a shows the absorption spectra of suspensions containing the different NCs. Compared with the peaks of the pure CsPbBr$_3$ NCs, the absorption edges of the Au@CsPbBr$_3$ and Pt@CsPbBr$_3$ NCs exhibited a red shift. Furthermore, the Pt@CsPbBr$_3$ NCs showed enhanced absorption for wavelengths greater than 520 nm, while the Au@CsPbBr$_3$ NCs showed additional weak absorption peaks at 420–450 nm. These differences in the absorption spectra may be attributed to the surface plasmonic resonance effect of the deposited noble metal nanoparticles [16]. All the NCs exhibited PL emission peaks at 532 nm, and their full-width-at-half-maximum values were also similar (Figure 6b). A bright-green PL emission was observed under UV-light irradiation (Figure 6c). Time-resolved PL (TRPL) measurements were performed to study charge transfer from the perovskite NCs to the noble metal nanoparticles. The TRPL decay data (Figure 6d) were fitted using a biexponential function of time, $I(t)$, as shown in Equation (1), and the fitted data are listed in Table 1.

$$I(t) = A_1e^{-t/\tau_1} + A_2e^{-t/\tau_2}$$  \hspace{1cm} (1)
quenching of the free charges because of electron transfer from the perovskite to the noble metal nanoparticles.

![Absorption and photoluminescence (PL) spectra and photographs and time-resolved PL (TRPL) decay curves](image)

Figure 6. (a) Absorption and (b) photoluminescence (PL) spectra and (c) photographs and (d) time-resolved PL (TRPL) decay curves of different NC suspensions. Photographs were taken under daylight (top) and UV-light illumination (bottom).

Table 1. Fitting parameters for TRPL spectra shown in Figure 6d.

| Sample                  | $\tau_1$ (ns) | $A_1$   | $\tau_2$ (ns) | $A_2$   |
|-------------------------|---------------|---------|---------------|---------|
| CsPbBr$_3$ NCs          | 14.8          | 0.94    | 59.4          | 0.06    |
| Au@CsPbBr$_3$ NCs       | 7.03          | 0.95    | 28.2          | 0.05    |
| Pt@CsPbBr$_3$ NCs       | 2.49          | 0.96    | 9.97          | 0.04    |

3. Materials and Methods

Materials: Cesium carbonate (Cs$_2$CO$_3$, 99.99%), lead (II) bromide (PbBr$_2$, 99.99%), OA (90%), OAm (80%), 1-octadecene (ODE, 90%), and dihydrogen hexachloroplatinate(IV)
hexahydrate (H$_2$PtCl$_6$·6H$_2$O, AR) were purchased from Shanghai Aladdin Biochemical Technology Co., Ltd. (Shanghai, China). Hexane and methanol (99.5%) were purchased from Sinopharm Chemical Reagent Co., Ltd. (Shanghai, China). Gold(III) chloride trihydrate (HAuCl$_4$·3H$_2$O, 99.9%) was obtained from China National Pharmaceutical Group Co., Ltd. (Beijing, China). All chemicals were used without further purification.

Synthesis of Cs$_4$PbBr$_6$ NCs: The Cs$_4$PbBr$_6$ NCs were prepared using the hot-injection method [17]. A cesium oleate solution was prepared by mixing 0.32 g of Cs$_2$CO$_3$ (0.98 mmol), 1 mL of OA, and 16 mL of ODE in a 100-mL three-neck flask. The solution was dried for 1 h at 120 °C under vacuum and then heated in an Ar atmosphere to 150 °C until all the Cs$_2$CO$_3$ had reacted with OA. During a typical run to synthesize the Cs$_4$PbBr$_6$ NCs, OAm (2 mL), OA (2 mL), ODE (20 mL), and PbBr$_2$ (0.4 mmol) were added to a 100-mL three-neck flask and dried under vacuum for 1 h. The reaction mixture was then heated to 140 °C. Next, 8.8 mL of a hot cesium oleate solution was rapidly injected into the PbBr$_2$ solution. After 6 s, the reaction mixture was immersed in an ice-water bath for immediate cooling. The product was centrifuged at 12,000 rpm for 5 min to remove the ODE and unreacted OA as well as the OAm. The precipitate was collected and redispersed in hexane and then centrifuged at 3000 rpm for 5 min to remove the oversized Cs$_4$PbBr$_6$ NCs. The concentration of the Cs$_4$PbBr$_6$ NC suspension was adjusted to 1 mg/mL for further synthesis.

Synthesis of Pt@CsPbBr$_3$ NCs: During a typical synthesis process, 0.1 mL of methanol was mixed with 0.4 mL of an aqueous H$_2$PtCl$_6$·6H$_2$O solution (1.448 mM), and the mixture was injected into 10 mL of the Cs$_4$PbBr$_6$ NC suspension. The system was irradiated with a halogen lamp (~20 mW/cm$^2$) for 1 h under vigorous stirring and then kept undisturbed under the ambient conditions for 12 h. The product was centrifuged at 6000 rpm for 5 min. The precipitates were discarded, and the supernatant was collected for characterization.

Synthesis of Au@CsPbBr$_3$ NCs: The synthesis procedure was similar to that for Pt@CsPbBr$_3$ NCs, except that an aqueous HAuCl$_4$·3H$_2$O solution (1.904 mM) was used.

Synthesis of CsPbBr$_3$ NCs. The synthesis procedure was similar to that of the Pt@CsPbBr$_3$ NCs, except that pure water without a metal precursor or methanol was used.

Characterization: The UV-vis absorption spectra were recorded using a Shimadzu UV-vis spectrophotometer (UV3600) (Shimadzu, Kyoto, Japan). The morphologies of the NCs were characterized by means of TEM (JEOL JEM-2100HR) (JEOL Ltd., Tokyo, Japan). For the XRD measurements, the NC suspensions were cast on cleaned glass substrates and dried. The measurements were performed using a Rigaku diffractometer (D/max 2500 PC) (Rigaku Corporation, Akishima, Tokyo, Japan) and Cu Kα radiation. XPS analysis was carried out using PHI Quantera II (Ulvac-PHI, Kanagawa, Japan). The PL and TRPL decay measurements were performed using a Horiba DeltaFlex PL system (Horiba, Tokyo, Japan) with a 520-nm pulsed laser for excitation at approximately 25 °C.

4. Conclusions

In this study, Au and Pt@CsPbBr$_3$ NCs were successfully synthesized via a reaction at the water/hexane interface, which involved the transformation of Cs$_4$PbBr$_6$ into CsPbBr$_3$ and its subsequent modification with a noble metal. The prepared CsPbBr$_3$ NCs exhibited a cubic perovskite phase and an average size of approximately 13.5 nm. The deposited Au and Pt nanoparticles were crystalline, with an fcc lattice and average diameters of approximately 3.9 and 4.4 nm, respectively. The noble metal modification process affected the charge recombination kinetics of the CsPbBr$_3$ NCs, probably owing to the quenching of the free charges because of electron transfer from the perovskite to the noble metal nanoparticles. This simple strategy for designing and synthesizing perovskite/noble metal nanostructures offers new opportunities to increase their applicability in photodetectors, LEDs, solar cells, photocatalysts, and other such devices. With respect to solar cells, CsPbBr$_3$ NCs modified with a noble metal may be used to construct perovskite quantum dot-based solar cells. Compared with bare perovskite NCs, those modified with a noble metal can enhance the light-harvesting properties of devices owing to their localized surface plasmon resonance effect. In the case of photocatalysts, a composite of Au or Pt@CsPbBr$_3$ NCs and
reduced graphene oxide or TiO₂ would be suitable for use as a visible-light-responsive photocatalyst for H₂ evolution in aqueous HI solutions. Modification with a noble metal has the potential to enhance the photocatalytic activity of CsPbBr₃ NCs. Further research in these directions is underway in our laboratory.

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