Energy Down-Conversion Cs₃Cu₂Cl₅ Nanocrystals for Boosting the Efficiency of UV Photodetector

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Zero-dimension (0-D) lead halide perovskite nanocrystals (NCs) have attracted a sight of interest in the field of optoelectronic devices due to their outstanding properties, such as high photoluminescence quantum yield (PLQY) and size- and composition-controlled tunable emission wavelengths. However, the toxicity of lead (Pb) element in the lead perovskite NCs is the bottleneck for the commercial application of perovskite NCs. Herein, we report a facile ligand-assisted synthesis to achieve lead-free Cs₃Cu₂Cl₅ NCs with a high PLQY of ∼70% and good stability against environmental oxygen/moisture as a promising down-conversion material. It has good merits of high PLQY and large Stokes shift (∼300 nm) originated from the effect of Jahn–Teller distortion and self-trapped excitons (STEs). Furthermore, the Cs₃Cu₂Cl₅ NCs embedded composite films (NCCFs) were utilized to enhance the ultraviolet (UV) response of silicon (Si) photodetectors. External quantum efficiency (EQE) measurements show that the UV response can be greatly improved from 3.3 to 19.9% @ 295 nm based on NCCFs combined with Si photodiodes. Our work offers an effective approach to develop highly efficient and stable lead-free Cs₃Cu₂Cl₅ NCs for the application in the solar-blind UV photodetector.

Keywords: lead-free perovskite, all-inorganic perovskite nanocrystals, photoluminescence quantum yield, ligand-assisted, stability

INTRODUCTION

In the field of photodetection, silicon-based photodetectors have achieved great progress on account of their broadband spectral response, high responsivity, and cost-effective fabrication process, while the response of silicon-based photodetectors in the ultraviolet (UV) region has been limited by the shallow penetration depth of UV radiations (less than 20 nm @370 nm wavelength) and high reflection coefficient (An et al., 2013). One strategy is to enhance the UV detection of Si photodetectors by down-conversion materials such as all-inorganic lead halide perovskite nanocrystals (NCs) that transform the UV photons to visible lights (Zhang et al., 2018).
A good down-conversion layer needs to display high light absorption in the UV wavelength, good quantum yield, extraordinary transparency in the visible region, and large Stokes shift to reduce the self-reabsorption losses. Moreover, structural simplicity, low-cost fabrication, ambient operation, and long-term stability are also crucial figures of merit for real applications (Wang et al., 2017).

The origin of the Stokes shift is closely related to the excitonic finestructure, which consists of a manifold of closely spaced optical transitions arising from the quasi-degenerate bandedge electronic states (Voznyy et al., 2016), leading to the downconversion phenomenon of the nanocrystals (NCs). However, the relatively small Stokes of lead halide perovskite NCs’ shift brings about significant self-reabsorption loss. Moreover, the toxicity of lead (Pb) element in the lead perovskite NCs is the bottleneck for the commercial application of perovskite NCs (Chang et al., 2018). Recently, the work of lead-free perovskite NCs, including tin-based (Jellicoe et al., 2016; Wang et al., 2016; Zhang et al., 2016; Wang et al., 2017; Liang et al., 2020) and bismuth-based (Leng et al., 2016, 2018; Yang et al., 2017; Bekenstein et al., 2018; Zhou et al., 2018; Gao et al., 2019) perovskite NCs, has been targeted. Replacing lead with divalent non-toxic metal cations (e.g., tin and bismuth) or heterovalent metal cations (e.g., antimony and silver) has been developed to synthesize lead-free perovskite NCs. Unfortunately, the rapid oxidation of Sn$^{2+}$ to Sn$^+$ in air is detrimental to the photoluminescence quantum yield (PLQY) (Jellicoe et al., 2016; Wang et al., 2016, 2017; Zhang et al., 2016, 2017; Bekenstein et al., 2018; Chang et al., 2018; Leng et al., 2018; Zhou et al., 2018; Gao et al., 2019; Liang et al., 2020; Pan et al., 2020b). Therefore, it is of great interest to exploit novel NCs with lower toxicity and higher stability, especially for larger Stokes shift as down-conversion materials.

The copper halide NCs, such as Cs$_3$Cu$_2$X$_5$ (X = Cl, Br, and I) with large Stokes shift (~300 nm) and high PLQY in visible spectrum but sensitivity to UV radiation as well as excellent photo stability, appear to be ideal candidates as down-conversion materials for the UV photodetection application (Yang et al., 2018; Cheng et al., 2019; Li et al., 2020; Lian et al., 2020a,b; Luo et al., 2020; Wang et al., 2020). The tetrahedral structure of Cs$_3$Cu$_2$X$_5$ can be formed by Cu$^+$ coordinated with halide ions, which is different from octahedral structure of lead halide perovskite NCs. Unfortunately, the rapid oxidation of Sn$^{2+}$ to Sn$^+$ in air is detrimental to the photoluminescence quantum yield (PLQY) (Jellicoe et al., 2016; Wang et al., 2016, 2017; Zhang et al., 2016, 2017; Bekenstein et al., 2018; Chang et al., 2018; Leng et al., 2018; Zhou et al., 2018; Gao et al., 2019; Liang et al., 2020; Pan et al., 2020b). Therefore, it is of great interest to exploit novel NCs with lower toxicity and higher stability, especially for larger Stokes shift as down-conversion materials.

**EXPERIMENT SECTION**

**Materials**

Cesium carbonate (Cs$_2$CO$_3$, 99.9%), oleylamine (OAm, 80–90%), n-hexane (AR), and isopropanol (AR) were purchased from Macklin, cuprous chloride (CuCl, ≥99.95%) was purchased from Aladdin, oleic acid (OA, 90%) was purchased from Aldrich, and 1-octadecene (ODE, ≥90%) was purchased from General Reagent. All of the chemicals were used as received without further purification.

**Preparation of Cs-Oleate**

0.3258 g of Cs$_2$CO$_3$, 6.84 mL of 1-octadecene, and 3.16 mL of oleic acid were loaded into a 50-mL three-neck flask and dried for 1 h at 110°C, and then heated to 120°C under N$_2$ until all Cs$_2$CO$_3$ reacted with OA. The precursor has to be pre-heated to 100°C before injection.

**Synthesis of Cs$_3$Cu$_2$Cl$_5$**

In a typical synthesis of Cs$_3$Cu$_2$Cl$_5$ NCs, 0.1188 g of CuCl, 2 mL of OA, and 1 mL of OAm were mixed in a 50-mL three-neck flask and degassed under vacuum for 1 h at 110°C. After complete solubilization of CuCl, the temperature was raised to 160°C and kept for 5 min. Then, 6 mL of Cs-oleate solution was quickly injected and, 5 s later, the reaction mixture was cooled with an ice-water bath. The solution was filtered by 0.22-μm filter and then centrifuged at 6,000 rpm for 5 min. After centrifugation, the supernatant was discarded and the precipitate was dissolved in the mixture of n-hexane and isopropanol (1:1 by volume). The mixture was centrifuged at 600 rpm for 5 min to collect the precipitants.

**Preparation of QDs Embedded Organic Polymer Film**

The Cs$_3$Cu$_2$Cl$_5$ NCs powder was dispersed in the mixture of polydimethylsiloxane (PDMS) and curing agent (10:1 by weight), followed by heating at 90°C for 3 h under vacuum. For the fabrication of NCs embedded modified polyester (Ecoflex) film, Cs$_3$Cu$_2$Cl$_5$ NCs solution was mixed with polyester and stirred intensely. The QDs embedded film was obtained after static positioning for 30 min at room temperature in air.
Characterization
Absorption and PL spectra of QDs were measured by using the U-4100 and NIR-300 spectrophotometers, respectively. PLE, PLQY, and TRPL spectra were measured by an Edinburgh instruments FS5 spectrometer. Transmission electron microscopic (TEM) image of QDs was taken by the Tecnai G20. Si photodiodes were purchased from Beijing Light Sensing Technologies Ltd., part number LSSPD-5.9. The photoresponse of the detectors was characterized by Keithley 2400 source meter.

RESULTS AND DISCUSSION
The tunneling electron microscopic (TEM) image shows that Cs$_3$Cu$_2$Cl$_5$ NCs have a mean size of 3.4 nm with uniform size distribution (Figure 1A). Moreover, the clear lattice fringes are observed in the high-resolution TEM (HRTEM) image, indicating the good crystallinity of NCs (Figure 1B). The X-ray diffraction (XRD) patterns in Figure 1C confirm that the crystal phase of the obtained NCs coincides with the standard PXRD data of Cs$_3$Cu$_2$Cl$_5$. Both the characterization results of TEM and XRD confirm the successful synthesis of the lead-free Cs$_3$Cu$_2$Cl$_5$ NCs. As shown in Figure 1D, the Cs$_3$Cu$_2$Cl$_5$ NCs adopt an orthorhombic space group Pnma (Lian et al., 2020a). There are two types of Cu$^{+}$ sites in this crystal that form [Cu$_2$Cl$_5$]$^{3-}$ units, namely, a tetrahedral site and a triangular site. The framework of Cs$_3$Cu$_2$Cl$_5$ NCs consists of [Cu$_2$Cl$_5$]$^{3-}$ that are separated by Cs$^+$ cations (Luo et al., 2020).

We further investigated the photophysical properties of Cs$_3$Cu$_2$Cl$_5$ NCs. Figure 2A reveals that the photoluminescence (PL) peak of Cs$_3$Cu$_2$Cl$_5$ NCs is located at 525 nm, while the absorbance peak is located at 225 nm, indicating a relatively large exciton binding energy, consistent with previous reports (Li et al., 2020; Lian et al., 2020a). We improved the PLQY of Cs$_3$Cu$_2$Cl$_5$ NCs by optimizing the quantities of OAm during the synthesis. The QY of NCs is improved from 5.6 to 69.1% when the volume of OAm is increased from 1 to 2 mL, which is consistent with the variation of PL intensity (Figure 2B). The time-resolved PL (TRPL) decay spectra of Cs$_3$Cu$_2$Cl$_5$ NCs were performed to gain a deep insight into the exciton recombination dynamics. As shown in Supplementary Figure 2 and Supplementary Table 1, the lifetime of Cs$_3$Cu$_2$Cl$_5$ NCs is improved from 77.8 to 119.8 µs when the volume of OAm is increased from 1 to 2 mL. The prolonged exciton lifetime indicates the suppression of non-radiative recombination. Furthermore, the radiative and non-radiative decay rates were calculated based on the PLQY and the average lifetimes. The non-radiative recombination rate of 2 mL OAm is the lowest, quantitatively manifesting that non-radiative recombination can be suppressed by appropriate OAm. Broad emission with a long lifetime is a characteristic of

![FIGURE 1](attachment:image1.png)

**FIGURE 1** | (A) TEM image of Cs$_3$Cu$_2$Cl$_5$ NCs. (B) HRTEM image of Cs$_3$Cu$_2$Cl$_5$ NCs. (C) XRD pattern of Cs$_3$Cu$_2$Cl$_5$ NCs. (D) Crystal structure of Cs$_3$Cu$_2$Cl$_5$. 

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Cu(I)-complexed PL emissions involving Jahn–Teller distortion. And the Stokes shift is determined primarily by the energy difference between Cu(I)3d^{10} and Cu(II)3d^{9} (Jun et al., 2018). The stability of the Cs₃Cu₂Cl₅ NCs was measured under ambient environment and the PL intensity can be remained 85% of the initial intensity after 30 days (Figure 2C). The decrease of the PL intensity may be attributed to the aggregation of the NCs in the colloidal solution or the oxidation of Cu⁺ to Cu²⁺ on the NCs' surface (Lian et al., 2020b). It is noted that the PL intensity of NCs has an obvious enhancement after 24 h, ascribing to the passivation of surface defect states by water molecules.

To shed light on the emission mechanism of Cs₃Cu₂Cl₅ NCs, the relationship between the emission and the excitation intensity is characterized to explore whether the PL emission is originated from permanent defects. As shown in Figure 2D, the PL intensity increases linearly with the increment of the excitation intensity, without saturation of PL, indicating that the PL emission arises from the intrinsic emitting states of material other than the permanent defect (Lian et al., 2020a). We also measured the emission wavelength-dependent photoluminescent excitation (PLE) spectra and excitation wavelength-dependent PL spectra of Cs₃Cu₂Cl₅ NCs. The normalized PLE spectra kept the uniform shapes with different detected emission wavelengths ranging from 480 to 580 nm. Similarly, the normalized PL spectra kept the identical features when the excitation wavelength increased from 250 to 340 nm (Supplementary Figure 3). Such uniform features of PLE and PL spectra with different detected emission or excitation wavelengths reveal that the green PL emission of Cs₃Cu₂Cl₅ NCs originates from the relaxation of the same excited states instead of ion luminescence. All the characterizations above indicate that the STEs obviously dominate the emission mechanism of the Cs₃Cu₂Cl₅ NCs.

Considering that the transmittance of NCCFs is vital for the performance of photodetectors coated with NCCFs, the transmittance of different organic polymer films was investigated. Figure 3A shows that the PDMS films are highly transparent with an average transmittance of 70% @400–800 nm, while the transmittance of modified polyester (Ecoflex) films decreases almost linearly from 44.48% @800 nm to 16.5% @ 400 nm. Figure 3B shows that the PL and Absorbance spectra of NCCFs are consistent with those of NCs, indicating that the superior optical property of Cs₃Cu₂Cl₅ NCs is well preserved. The high PLQY, good transmittance, and outstanding stability make NCCFs an attractive candidate for down-conversion application to enhance the UV detection of Si photodetectors. As shown in Figure 3C, the NCCFs were directly coated onto the surface of the Si photodiode, covering the whole detection area.

![FIGURE 2](image-url) | (A) PL and absorption spectra of Cs₃Cu₂Cl₅ NCs solution. (B) PL spectra of different quantities of OAm-capped NCs. (C) Air-stability test of Cs₃Cu₂Cl₅ NCs. (D) Dependence of the emission intensity in Cs₃Cu₂Cl₅ NCs.
FIGURE 3 | (A) Transmittance spectra of the Ecoflex and PDMS. (B) Abs spectra and PL spectra of Cs$_3$Cu$_2$Cl$_5$ NCs and NCCFs. (C) Schematic diagram of the configuration of the UV-enhanced Si photodiode and the down-conversion process of NCCFs.

enhancement of UV detection of the Si photodetector with NCCFs can be analyzed by considering the light converting process as displayed in the right of Figure 3C. The incident photons with energy above the bandgap of the NCCFs can be transformed to visible photons by the down-conversion of film due to the high PLQY of Cs$_3$Cu$_2$Cl$_5$ NCs. Then most of the visible photons can pass through the NCCFs, followed by reception by the Si detector.

Subsequently, we characterized the Si photodetector coated with NCCFs to evaluate the effect of Cs$_3$Cu$_2$Cl$_5$ NCs on device performance. The photoresponsivity (R), specific detectivity ($D^*$), and external quantum efficiency (EQE) are evaluated according to the following equations (Li et al., 2017):

\[ R = \frac{I_p - I_d}{P \cdot S} \]  

\[ D^* = \frac{A^{1/2}R}{(2eI_d)^{1/2}} \]  

\[ \text{EQE} = \frac{I_p/e}{P/h\nu} = \frac{hc}{e\lambda}. \]

Where $I_p$, $I_d$, $S$, $A$, $h$, and $c$ is the photocurrent, dark current, effective illuminated area, detection area, Planck's constant, and light speed, respectively.

The EQE characteristics reveal that the bare Si photodetector was not sensitive to UV radiation ranging from 257 to 300 nm, while the response of the NCCFs-coated Si photodetector can be extended to 257 nm (Figure 4A). The maximum EQE of the NCCFs-coated Si photodetector achieves 19.66% @295 nm compared with 3.3% of the bare photodetector, indicating the efficient down-conversion of NCCFs for the Si photodetector. The EQE reduction of the photodetector in the visible spectrum can be ascribed to the limited transmittance of NCCFs and the self-absorption-induced loss. As shown in Figure 4B, the letter “U” was imaged by the NCCFs-coated Si photodetector equipped with a slide rail controlled by a stepping motor.

The response speed is a key parameter for a photodetector that reflects the stability to catch a fast-transforming optical signal. We used a signal generator to generate incident light with different frequencies and recorded the photocurrent through an oscilloscope in real time. The current–time (I–T) curves of the NCCFs-coated Si photodetector are recorded under the incident light with the frequency ranging from 100 to 3,000 Hz (Figure 4C and Supplementary Figure 5). It is obvious that the NCCFs-coated Si photodetector can operate at a relatively wide frequency with outstanding stability and reproducibility. The relative balance [($I_{max} - I_{min}$)/$I_{max}$] of the NCCFs-coated Si photodetector is maintained at 97.2% at a frequency of 3,000 Hz which indicates the potential application at high frequency and capacity to monitor ultrafast optical signals (Yan et al., 2019; Yu et al., 2019). Moreover, on the condition of 1,000 Hz incident light, the rise time of the device (from 10 to 90% of the saturated value) is about 28 μs, while the decay time is 40 μs (from 90 to 10%), indicating the fast response speed of the device (Figure 4D).
CONCLUSION

In conclusion, 0-D colloidal Cs$_3$Cu$_2$Cl$_5$ NCs have been successfully synthesized via a hot-injection method. The prepared Cs$_3$Cu$_2$Cl$_5$ NCs exhibit a high PLQY of ~70% with the PL peak located at 525 nm, a large Stokes shift, and an extremely long lifetime, which is attributed to the STEs’ effect. Moreover, the as-prepared NCs exhibited excellent air stability over 30 days. These features make them very suitable for down-conversion applications. By applying the NCCFs on to the top of the Si photodetector, we successfully fabricated the prototype devices of solar-blind UV-enhanced Si photodetectors. The NCCFs-coated Si photodiodes extended the response spectra to UV region of 257–300 nm and improved the EQE from 3.3 to 19.66% @295 nm. Our work offers an effective and low-cost approach to achieve UV-enhanced Si photodetectors toward broadband and solar-blind light detection.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author/s.

AUTHOR CONTRIBUTIONS

WL and JP led the device physics work. JC led the materials chemistry work. CL synthesized and optimized the QD materials and carried out the device fabrication and characterizations. ZhZ and FF conducted the optical measurements and analyzed the optical properties of QDs. CL and LW wrote the first draft of the manuscript. JC, WL, XZ, and JP provided major revisions. JC and WL supervised the project. All authors discussed the results and commented on the manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmats.2021.682833/full#supplementary-material
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**Conflict of Interest:** CL was employed by the company China Nuclear Power Engineering Co., Ltd.

The remaining authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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