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Green synthesis of highly stable CsPbBr₃ perovskite nanocrystals using natural deep eutectic solvents as solvents and surface ligands†

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Perovskite nanocrystals (PNCs) have attracted widespread attention as promising materials for the optoelectronic field due to their remarkable photophysical properties and structural tunability. However, their poor stability and the use of toxic organic solvents in the preparation process have severely restricted their practical applications. Herein, a facile, rapid and toxic organic solvent-free synthesis strategy of CsPbBr₃ PNCs was developed for the first time via the ligand-assisted reprecipitation (LARP) method using natural deep eutectic solvents (NADESs) as solvents and surface ligands. In this method, the NADESs not only functioned as solvents for green synthesis, but also served simultaneously as surface ligands of CsPbBr₃ PNCs to significantly improve their optical properties and stability. The as-synthesized CsPbBr₃ PNCs exhibited high photoluminescence quantum yield (PLQY, ~96.8%), narrow full width at half-maximum (FWHM, ~18.8 nm) and a high stability that retained 82.9% of PL intensity after 70 days. This work provides a new strategy for the green synthesis of PNCs, which promises feasibility for the industrial large-scale synthesis of high-quality PNCs.

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Introduction

In recent years, perovskite nanocrystals (PNCs) have received much attention because of their excellent optoelectronic properties, such as narrow full width at half-maximum (FWHM), high photoluminescence quantum yield (PLQY), low exciton binding energy, tunable band gap, and low defect density.^{1–3} They show great potential for application in the fields of light-emitting diodes (LED),^{4,5} solar cells,^{6–8} lasers⁹ and photocatalysis.^{10–12} Currently, the most common synthesis methods for PNCs are the hot-injection method¹³ and the ligand-assisted reprecipitation (LARP) method.^{14,15} In the LARP method, PNCs are obtained through the regulation of volume and concentration of solvents and precursors. Precursors are firstly dissolved in good solvents, such as dimethyl sulfoxide (DMSO) and dimethylformamide (DMF),

and then transferred into poor solvents (antisolvents), such as toluene and *n*-octane, to form a supersaturated solution and precipitate the crystals under the assistance of ligands, such as oleic acid (OA) and oleylamine (OAm). This method is promising due to its ease of operation, low equipment requirement and mild operational temperatures. However, there are two disadvantages of the LARP method. On the one hand, a large amount of toxic organic solvents is inevitably used, which significantly limits their applications. On the other hand, the proton exchange between OA and OAm may induce the ligands to be easily lost from the surface of PNCs during the purification process or storage period, resulting in poor stability of PNCs.

Recently, great efforts have been devoted to developing effective ways to solve the above problems. Li¹⁶ *et al.* developed a novel low-toxicity antisolvent synthesis (LTAS) process based on tetraethyl orthosilicate (TEOS) to obtain monodispersed and luminescent all-inorganic PNCs. Wang's¹⁷ group proposed a facile method to synthesize green CsPbBr₃ PNCs using the ionic liquid (IL) 1-butyl-3-methylimidazolium bromide ([Bmim]Br). Luo¹⁸ *et al.* recently reported the synthesis of blue-emission CsPbBr₃ PNCs with the assistance of hydrophobic ILs. However, ILs have some disadvantages, such as complex preparation processes, high cost and poor biocompatibility.^{19,20} In addition, Tan²¹ *et al.* proposed a novel strategy using octylphosphonic acid (OPA) as the capping

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ligand to solve the problem of ligands loss and improve the stability of CsPbX₃ PNCs. Mishra²² *et al.* reported bromopropane as a novel bromine precursor for the completely amine-free colloidal synthesis of CsPbBr₃ PNCs. Research on the direction of amine-free synthesis emerges in an endless stream.^{23–25}

Deep eutectic solvents (DESs), introduced by Abbott²⁶ *et al.* in 2003, have been regarded as a new class of green solvents. DESs are low-eutectic mixtures of compounds composed of a hydrogen bond donor (HBA) and a hydrogen bond acceptor (HBD).²⁷ They possess not only similar favorable properties to ILs, but also the advantage of being easy to prepare, cost-effective, and biodegradable.^{28,29} Among them, natural deep eutectic solvents (NADESSs), composed of natural compounds such as carboxylic acids, sugars, alcohols, amines and amino acids, have attracted more attention because they fully meet the green chemistry principles.³⁰ As the most promising alternative to toxic organic solvents, NADESSs have been applied successfully in the fields of catalysis,^{31,32} extraction,^{33,34} electrochemistry^{35,36} and organic synthesis.^{37,38} Furthermore, the –COOH, –OH, and –NH₂ groups in these natural compounds can coordinate with metal ions, which provides feasibility for using NADESSs as ligands. To the best of our knowledge, NADESSs used simultaneously as solvents and surface ligands in the synthesis of PNCs have not been reported so far.

Herein, a facile, rapid and toxic organic solvent-free synthesis strategy for CsPbBr₃ PNCs was developed for the first time *via* the ligand-assisted reprecipitation (LARP) method using NADESSs as solvents and surface ligands. First, NADESSs, consisting of thymol as the HBA and L-lactic acid (Thy:LacA) and *n*-octanol (Thy:Oct) as the HBD, were used as good solvents, and a NADES consisting of thymol as the HBA and decanoic acid (Thy:DecA) as the HBD was used as antisolvent for the green synthesis of NADES-CsPbBr₃ PNCs. Moreover, the –COOH and –OH groups in Thy:Oct and Thy:DecA can chelate with Pb²⁺ and provide a strong interaction with CsPbBr₃ PNCs. Thereby, Thy:Oct and Thy:DecA can also act as surface ligands to significantly enhance the optical properties and stability of CsPbBr₃ PNCs. This facile, rapid and green synthesis method of PNCs will provide a new avenue for their sustainable industrial production in various fields.

Experimental

Reagents and solutions

DL-Menthol (C₁₀H₂₀O, 98%), thymol (C₁₀H₁₄O, 98%), L-lactic acid (C₃H₆O₃, 90%), *n*-octanoic acid (C₈H₁₆O₂, 99%), decanoic acid (C₁₀H₂₀O₂, 99%), lauric acid (C₁₂H₂₄O₂, 98%), oleic acid (C₁₈H₃₄O₂, AR), ethanol (C₂H₆O, 99%), *n*-butanol (C₄H₁₀O, 99%), *n*-hexanol (C₆H₁₄O, 99%), *n*-octanol (C₈H₁₈O, 99%), and cesium bromide (CsBr, 99.5%) were obtained from Aladdin Industrial Co. Ltd (Shanghai, China). *n*-Decanol (C₁₀H₂₂O, 98%), lead(II) bromide (PbBr₂, 99%), and oleylamine (C₁₈H₃₇N, 80–90%) were purchased from Macklin Biochemical

Technology Co. Ltd. (Shanghai, China). All reagents and solutions were used as received without further purification.

Apparatus

Fourier transform infrared (FTIR) spectra (NICOLET iS 10) and ¹H NMR (Avance 500 MHz; Bruker) were applied to determine the formation of NADESSs. The phase of the NADES-CsPbBr₃ PNCs was identified *via* an X-ray diffractometer (XRD, Rigaku Ultima IV). Ultraviolet–visible (UV-vis) absorption spectra were tested using a UV-vis spectrophotometer (Specord plus 200). Photoluminescence (PL) emission spectra were recorded using a spectrometer (Techcomp FL970 Plus). NADES-CsPbBr₃ PNCs were characterized using luminescence decay curves and photoluminescence quantum yields (PLQY; FLS980, Edinburgh) under excitation at 365 nm. Transmission electron microscopy (TEM) images and high-resolution TEM images were taken on a JEM-2100.

Preparation of NADESSs

In this study, different NADESSs were prepared by the following steps. Firstly, the [HBA:HBD] of all NADESSs was mixed at a molar ratio of 1:1, and using DL-menthol and thymol as the HBA, combined with eight different HBDs comprising L-lactic acid, *n*-octanoic acid, decanoic acid, lauric acid, ethanol, *n*-butanol, *n*-hexanol and *n*-octanol. Secondly, the mixture was heated and stirred at 80 °C and 700 rpm for 90 minutes until a clear homogeneous liquid was obtained, which indicated the formation of NADESSs. The as-obtained NADESSs were as follows: DL-menthol:L-lactic acid (Man:LacA), DL-menthol:*n*-octanoic acid (Man:OctA), DL-menthol:decanoic acid (Man:DecA), DL-menthol:lauric acid (Man:LauA), thymol:*n*-octanoic acid (Thy:OctA), thymol:lauric acid (Thy:LauA), thymol:ethanol (Thy:Eth), thymol:*n*-butanol (Thy:But), thymol:*n*-hexanol (Thy:Hex), Thy:LacA, Thy:Oct and Thy:DecA, as shown in Table S1.†

Synthesis of NADES-CsPbBr₃ PNCs

The synthesis processes of NADESSs and NADES-CsPbBr₃ PNCs are illustrated in Fig. 1. and the NADESSs used are listed in Table S2.† In a typical procedure, under vigorous stirring at 50 °C, 0.02 mmol of PbBr₂, 100 μL of OA and 75 μL of OAm were dissolved in 1 mL of Thy:Oct, and 0.3 mmol of CsBr were dissolved in 1 mL of Thy:Lac to form Pb- and Cs-precursor solutions. Then, 600 μL of Pb-precursor and 119 μL of Cs-precursor solution (molar ratio of Cs:Pb = 3:1) was quickly added to 3 mL of Thy:DecA at the same time under vigorous stirring, and bright blue-green PL emission was observed immediately. After 5 hours, the CsPbBr₃ colloidal solution was centrifuged at 10 000 rpm for 10 min to obtain NADES-CsPbBr₃ PNCs. Finally, the product was washed and stored in *n*-hexane.

Fabrication of the white LED device

First, the as-synthesized NADES-CsPbBr₃ PNCs powders were blended with silicone gel A and B (A:B = 1:4) and mixed by stirring to obtain a fine dispersion of NADES-CsPbBr₃ PNCs. Then the mixture was vacuumed to discharge air bubbles. A

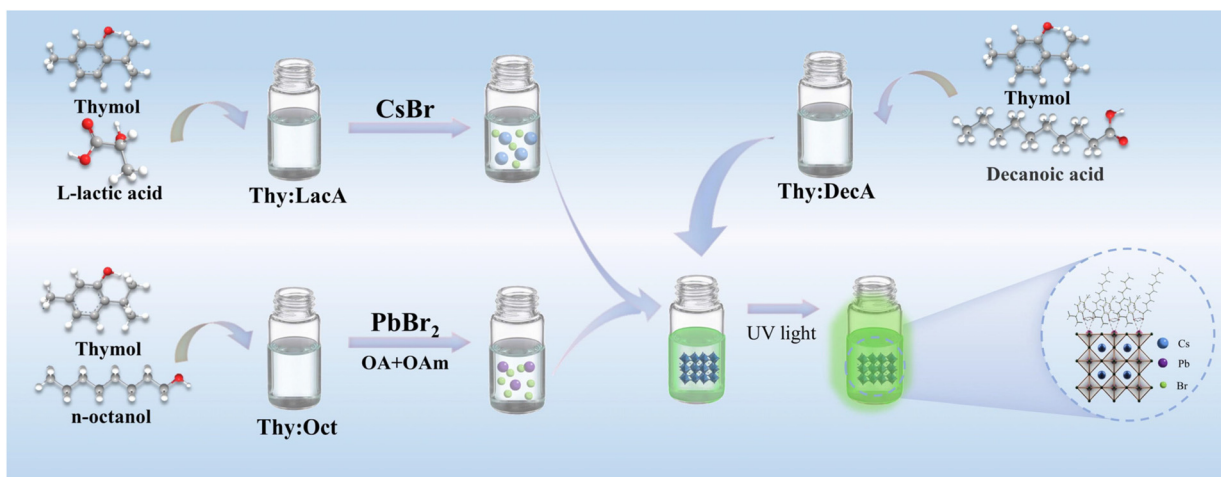


Fig. 1 Schematic illustration of synthetic procedure for NADES-CsPbBr₃ PNCs.

WLED device was constructed by combining a 480 nm GaN LED blue chip with the green-emitting NADES-CsPbBr₃ PNC glass and commercial red-emitting KSF phosphor.

Results and discussion

Characterization of NADESs

NADESs were formed by the hydrogen bond between HBA and HBD. To prove the formation of Thy:Oct, the FTIR and ¹H NMR characterizations of Thy:Oct and their constituents were performed, as shown in Fig. 2. From the FTIR spectra of Thy:Oct and its constituents (Fig. 2a), the broad absorption bands associated with the stretching vibrations of O–H groups were observed in the range 3100–3400 cm⁻¹. Compared with the O–H stretching bands of thymol and octanol respectively located at 3231.40 cm⁻¹ and 3317.54 cm⁻¹, the O–H stretching band of Thy:Oct, located at 3342.30 cm⁻¹, was shifted to a higher wavenumber. This result could be regarded as the contribution of the hydrogen bond formation between HBA and HBD, which indicated the formation of Thy:Oct. The ¹H NMR spectra of Thy:Oct and its constituents are presented in Fig. 2b. The characteristic H resonance peaks of thymol and

octanol can be observed in spectrum of Thy:Oct. Furthermore, the peaks of the –OH of thymol from $\delta = 4.78$ to $\delta = 6.79$ and the –OH group of octanol from $\delta = 3.74$ to $\delta = 2.88$ can be observed. The chemical shift value changes because the formation of hydrogen bonds changes the electron density around the hydrogen nucleus, verifying the formation of Thy:Oct. Similarly, as shown in Fig. S1 and S2,[†] the formation of Thy:LacA and Thy:DecA can also be proved by FT-IR and ¹H NMR.^{39,40}

The polarity of the NADESs can be evaluated based on the solvatochromic method. Nile red solvatochromic dyes comprise one of the most widely used probes that has been extensively employed to measure the polarity of NADESs.⁴¹ The solvatochromic parameter E_{NR} can be calculated using the following equation:

$$E_{NR} = 28591/\lambda_{max}$$

where E_{NR} is in kcal mol⁻¹ and λ_{max} is the wavelength of the maximum absorption of the probe in the NADESs. High E_{NR} values correspond to lower polarity and low E_{NR} values correspond to higher polarity of NADESs. The absorbance spectra of Nile red dissolved in Thy:LacA, Thy:Oct and Thy:DecA is shown in Fig. S3.[†] The λ_{max} values were located at 592 nm, 566 nm and 563 nm respectively. The polarity values of Thy:LacA, Thy:Oct and Thy:DecA as calculated by E_{NR} scales were 48.3 kcal mol⁻¹, 50.5 kcal mol⁻¹ and 50.7 kcal mol⁻¹ respectively.



Fig. 2 (a) The FT-IR spectra of thymol, *n*-octanol and Thy:Oct; (b) ¹H NMR spectra of thymol, *n*-octanol and Thy:Oct.

Characterization of NADES-CsPbBr₃ PNCs

The phase of NDES-CsPbBr₃ NCs were conducted *via* X-ray diffraction (XRD) (Fig. 3a). The diffraction peaks of NADES-CsPbBr₃ PNCs and typical CsPbBr₃ PNCs were well coincident with a monoclinic phase of CsPbBr₃ (PDF card no. 00-018-0364). The diffraction peaks located at $2\theta = 15.1^\circ$, 21.5° , 30.7° , 34.2° , 37.6° and 43.7° can be assigned to (001), (110), (200), (210), (211) and (202) crystal planes, respectively. Furthermore, compared with the typical LARP-synthesized CsPbBr₃ PNCs, NADES-CsPbBr₃ PNCs exhibited higher and sharper peaks indicating NADES-CsPbBr₃ PNCs have better



Fig. 3 (a) The XRD patterns of CsPbBr₃ PNCs synthesized by two methods; (b and c) TEM images of NADES-CsPbBr₃ PNCs. Inset in (c) shows the corresponding HRTEM image; (d) size distribution of NADES-CsPbBr₃ PNCs; (e) absorption and PL spectra of NADES-CsPbBr₃ PNCs. Inset in (e) shows the photograph of NADES-CsPbBr₃ PNCs under daylight (left) and under UV light (right); (f) the decay curve of NADES-CsPbBr₃ PNCs.

crystalline quality. The transmission electron microscopy (TEM) graph of NADES-CsPbBr₃ PNCs indicated that NADES-CsPbBr₃ PNCs possess a cubic shape and is well dispersed (Fig. 3b and c). Moreover, the high-resolution TEM image (inset of Fig. 3c) revealed a highly ordered crystalline lattice and the lattice spacing was calculated to be 0.29 nm, which corresponds to the (200) lattice space of monoclinic CsPbBr₃ PNCs. The size ranged from 7 to 18 nm and the average widths were around 11.8 nm as illustrated in Fig. 3d.

Optical properties of the as-synthesized NADES-CsPbBr₃ PNCs were also studied. The UV-vis absorption and PL emission spectra of NADES-CsPbBr₃ PNCs are presented in Fig. 3e. The UV-vis absorption spectra showed an obvious shoulder peak at about 504 nm. Meanwhile, a single strong PL peak appeared at around 511 nm with a narrow FWHM of ~18.8 nm. As shown in Fig. S4,† the as-synthesized NADES-CsPbBr₃ NCs exhibited a high PLQY value of ~96.8%. The inset of Fig. 3e shows a NADES-CsPbBr₃ PNC yellow-green solution under daylight, and bright green light emission under UV light. Furthermore, the time-resolved PL measurements were carried out to verify the exciton recombination dynamics, as shown in Fig. 3f. The PL decay could be well fitted by a biexponential function,¹⁵

$$A(t) = A_1 \exp\left(\frac{-t}{\tau_1}\right) + A_2 \exp\left(\frac{-t}{\tau_2}\right)$$

where A , A_1 , and A_2 are constants and τ_1 and τ_2 are two biexponential lifetimes with amplitudes A_1 and A_2 , respectively. The average lifetime (τ_{ave}) can be calculated as

$$\tau_{\text{ave}} = \frac{A_1 \tau_1^2 + A_2 \tau_2^2}{A_1 \tau_1 + A_2 \tau_2}$$

The obtained NADES-CsPbBr₃ PNCs exhibit two lifetimes (τ): τ_1 of 5.78 ns accounting for 67.6%, and τ_2 of 19.00 ns accounting for 32.4%, respectively. The short decay time (τ_1) originated from excitonic radiative recombination, while the

long decay time originated from a longer charge trapping/detrapping process.⁴² The longer τ_2 of NADES-CsPbBr₃ PNCs is consistent with its higher surface defects, while the smaller proportion of τ_2 indicates the low defect density of the synthesized NADES-CsPbBr₃ PNCs. However, this may need further spectroscopic study.

The selection of NADESs

The synthesis of CsPbBr₃ PNCs by the supersaturation recrystallization method was achieved through dissolving the precursors in a good solvent, and then transferring them into a poor solvent (antisolvents) to form a supersaturated solution and precipitate crystals. Therefore, the solubility of precursors in the related solvents plays an important role in this method. The solubility of CsBr and PbBr₂ in 13 kinds of NADESs was investigated and is summarized in Table S3.† As can be seen, CsBr could dissolve in Thy:LacA, and PbBr₂ could dissolve in four NADESs, comprising Thy:Oct, Thy:But, Thy:Hex and Thy:Oct. To further obtain the best dissolve solvents of PbBr₂, four different schemes (Table 1) were designed to synthesize NADES-CsPbBr₃ PNCs and the PL spectra of the products were investigated (Fig. 4a). As shown in Fig. 4a, the NADES-CsPbBr₃ PNCs synthesized *via* the TLTO scheme exhibited the best optical properties. Hence, Thy:LacA and Thy:Oct were used as the dissolving solvents for CsBr and PbBr₂, respectively. Furthermore, the optimal NADES to be used as an antisolvent was also evaluated. It can be seen from Fig. 4b and Table S4† that the NADES-CsPbBr₃ PNC synthesized using Thy:DecA as an antisolvent showed the strongest PL intensity. Therefore, Thy:DecA was selected as the antisolvent in this experiment.

Effect of the concentration of Cs- and Pb-precursor solutions

As the structural disintegration of PNCs is likely to occur in a good solvent, it is necessary to use highly concentrated precursor solutions to reduce the amount of good solvent required in the LARP method. To investigate the influence of the concentration of Cs- and Pb-precursor solutions on the luminescence intensity of NADES-CsPbBr₃ PNCs in this method, a series of experiments were studied using various concentrations of Cs- and Pb-precursor solutions (Table S5 and S6†). Fig. 4c and d showed that the luminescence intensity of NADES-CsPbBr₃ PNCs was increased with the increase in the concentration of Cs- and Pb-precursor solutions, and the highest intensity was

Table 1 Different combination schemes used to synthesize NADES-CsPbBr₃ PNCs

Name	CsBr	PbBr ₂	Antisolvent
TLTE	Thy:LacA (0.3 mol·L ⁻¹)	Thy:Eth (0.02 mol·L ⁻¹)	Thy:DecA (3 mL)
TLTB	Thy:LacA (0.3 mol·L ⁻¹)	Thy:But (0.02 mol·L ⁻¹)	Thy:DecA (3 mL)
TLTH	Thy:LacA (0.3 mol·L ⁻¹)	Thy:Hex (0.02 mol·L ⁻¹)	Thy:DecA (3 mL)
TLTO	Thy:LacA (0.3 mol·L ⁻¹)	Thy:Oct (0.02 mol·L ⁻¹)	Thy:DecA (3 mL)



Fig. 4 (a) PL spectra of NADES-CsPbBr₃ PNCs synthesized by different schemes; (b) PL spectra of NADES-CsPbBr₃ PNCs synthesized by different antisolvents; (c) PL spectra of NADES-CsPbBr₃ PNCs synthesized by different concentration of Cs-precursor; (d) PL spectra of NADES-CsPbBr₃ PNCs synthesized by different concentrations of Pb-precursor.

obtained at up to 0.3 mol L⁻¹ Cs-precursor solutions and 0.02 mol L⁻¹ Pb-precursor solutions, respectively.

Optimum molar ratios of Pb- and Cs-precursor

The molar ratios of Pb- and Cs-precursor ($n_{\text{Cs}}/n_{\text{Pb}}$) significantly affect the phase transformation of PNCs. Fig. 5 presents the XRD patterns of PNCs with different $n_{\text{Cs}}/n_{\text{Pb}}$ ratios (1 : 3, 1 : 1 and 3 : 1). The phase of the PNCs gradually changed from CsPb₂Br₅ to CsPbBr₃ PNCs with the increase in $n_{\text{Cs}}/n_{\text{Pb}}$. Different from the pure phase of CsPbBr₃ PNCs formed in the typical method, mixed phases of CsPb₂Br₅ and CsPbBr₃ PNCs were obtained with $n_{\text{Cs}}/n_{\text{Pb}} = 1 : 1$ in this method. This was probably caused by the strong coordination between Cs⁺ and Thy:LacA. When $n_{\text{Cs}}/n_{\text{Pb}}$ reached 3 : 1, the pure phase of CsPbBr₃ PNCs was obtained, for which the XRD pattern was assigned to three dimensional CsPbBr₃ (PDF card no.00-018-0364). Hence, $n_{\text{Cs}}/n_{\text{Pb}} = 3 : 1$ was selected in our experiment.

Stability of NADES-CsPbBr₃ PNCs

To avoid the reduction of the applied device performance by the residual organic ligands, purification processes of PNCs are necessary. Unfortunately, in the typical LARP-synthesized CsPbBr₃ PNCs, the shift of the UV-vis peak and the PL quenching phenomenon were observed because of severe aggregation of PNCs due to insufficient surface ligands after only two washing cycles (Fig. S5†). According to previous reports,^{43,44} the OA/Oam ligands were easily washed off because of their proton exchange and weak coordination ability with PNCs. In contrast, NADES-CsPbBr₃ PNCs exhibited much higher stability



Fig. 5 XRD patterns of PNCs synthesized with different molar ratios of CsBr and PbBr₂.

ity during purification processes. NADESs have a stronger coordination ability with PNCs and act as better surface ligands, which prevents the aggregation of PNCs. As shown in Fig. 6a and b, the UV-vis absorption and PL spectra of NADES-CsPbBr₃ PNCs barely changed after five cycles of the same washing processes.

The storage, light, thermal and water stability of NADES-CsPbBr₃ PNCs were further determined. The storage stability of the NADES-CsPbBr₃ PNCs was evaluated in ambient conditions and the results are shown in Fig. 6c. It can be seen that the PL emission intensity could preserve 82.9% retention after 70 days. As shown in Fig. 6d, the typical LARP-synthesized CsPbBr₃ PNCs only kept 0.6% of the initial PL intensity under irradiation with a UV lamp after 9 h, while 47.6% of the initial PL intensity of NADES-CsPbBr₃ PNCs was still reserved. In addition, as exhibited in Fig. 6e, the typical LARP-synthesized CsPbBr₃ PNCs kept 0.7% of the initial PL intensity after nine heating cycles at 85 °C, while the PL intensity of NADES-CsPbBr₃ PNCs could still be maintained at 43.8%. Finally, the water stability of CsPbBr₃ PNCs was tested by 2 mL of hexane solution put on the top of 1 mL of water (Fig. 6f). For the typical LARP-synthesized CsPbBr₃ PNCs, green fluorescence disappeared completely after 40 h, while the NADES-CsPbBr₃ PNCs maintained bright fluorescence. All these results demonstrate that, compared to the typical LARP-synthesized CsPbBr₃ PNCs, the stability of NADES-CsPbBr₃ PNCs was significantly improved.

Formation mechanism of NADES-CsPbBr₃ PNCs

To further understand the formation mechanism of NADES-CsPbBr₃ PNCs, the role of NADESs in this method is discussed. On the one hand, NADESs were used as solvents in the LARP method. To measure the dissolving capacity of three NADESs for CsBr and PbBr₂, their polarity was evaluated through the Nile red polarity scale. As a result, CsBr can directly dissolved in Thy:LacA of high polarity, and PbBr₂ can



Fig. 6 (a) The UV-vis patterns of NADES-CsPbBr₃ NC by different centrifugation times; (b) PL spectra of NADES-CsPbBr₃ PNCs by different centrifugation times; (c) time stability of NADES-CsPbBr₃ PNCs; (d) light stability of CsPbBr₃ PNCs; (e) thermal stability of CsPbBr₃ PNCs; (f) water stability of CsPbBr₃ PNCs. Above is NADES-CsPbBr₃ PNCs; below is typical LARP-synthesized CsPbBr₃ PNCs. Upper layer, hexane; bottom layer, water.

dissolve in Thy : Oct of lower polarity with the help of the coordination of OA/OAm. Meanwhile, Thy : DecA of the lowest polarity can be used as an antisolvent, leading to the mixture becoming a supersaturated solution and precipitating crystals to form NADES-CsPbBr₃ PNCs. On the other hand, NADESs were also used as surface ligands of CsPbBr₃ PNCs. For the synthesis of PNCs, introducing some organic molecules is necessary, which can strongly interact with ions and control crystal growth kinetics to reduce the defect density.⁴⁵ To verify the feasibility of using NADESs as surface ligands, the crystal growth kinetics and ¹H NMR characterization of NADES-CsPbBr₃ PNCs were performed. It can be seen from Fig. S6,† that the fluorescence of NADES-CsPbBr₃ PNCs changed from blue to green and the intensity gradually increased with the reaction time and reached the highest value at 5 h. The main reason for the difference from the rapid crystallization of typical synthetic PNCs is that the -COOH and -OH groups of NADESs can strongly coordinate with the Pb of CsPbBr₃ PNCs. From the ¹H NMR spectra of NADES-CsPbBr₃ PNCs OA and OAm (Fig. 7), it can be found that there is no carbon-carbon double bond characteristic hydrogen peak ($\delta = 5.37$) of OA and OAm in NADES-CsPbBr₃ PNCs, which indicates that not oleylamine oleate but NADESs can act as surface ligands to effectively passivate PNCs.

Light-emitting devices

Finally, the obtained NADES-CsPbBr₃ NCs with a bright pure green emission were applied for fabricating a WLED lamp based on a commercially available 460 nm GaN LED blue chip (the inset in Fig. 8a). Fig. 8a shows the typical electroluminescence (EL) spectra of the WLED at a driving current of 20 mA. To further investigate the potential applications of NADES-CsPbBr₃ NCs, we measured the luminous efficiency of the WLED driven by different currents, as shown in Fig. S7.†



Fig. 7 ¹H NMR spectra of NADES-CsPbBr₃ PNCs, OA and OAm.

The highest power efficiency reached 55.61 Lm W⁻¹ under a 20 mA current. The color coordinates of the sample were labeled as a red dot with a corresponding color coordinate of (0.3164, 0.3242) in the Commission Internationale de l'Eclairage (CIE) chromaticity diagram (Fig. 8b). Therefore, the obtained NADES-CsPbBr₃ PNCs reveal their potential commercial application in the light field.

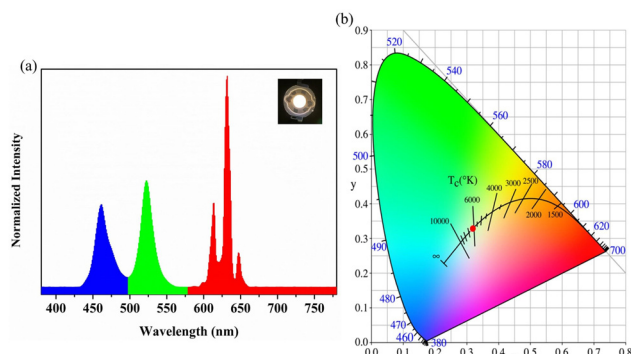


Fig. 8 (a) PL emission spectrum. The inset shows the photograph of device in open states; (b) CIE coordinates of NADES-CsPbBr₃ PNCs in CIE space.

Conclusions

In summary, the work focuses on designing a toxic organic solvent-free approach to the synthesis of CsPbBr₃ PNCs based on NADESS, for meeting the environment-friendly requirements to achieve the preparation of high-performance LED chips. The as-prepared NADES-CsPbBr₃ PNCs exhibit a high PLQY of ~96.8%, a narrow FWHM of ~18.8 nm and excellent stability. NADESS can not only use function as solvents, but also serve as a surface ligand to effectively reduce defects for the green synthesis of highly stable, bright-luminous NADES-CsPbBr₃ PNCs. Besides, a WLED lamp was fabricated to verify its practical application. Compared to the traditional LARP method, our method is simple and eco-friendly, which is desirable for the large-scale production of perovskite, and opens a new synthesis pathway to green exploitation of perovskite materials.

Author contributions

Heng Lu: conceptualization, methodology, investigation, writing – original draft, formal analysis. Xiaohong Tan: methodology and resources. Guobin Huang: data curation, writing – review and editing. Shaoru Wu: investigation. Yanmei Zhou: visualization and supervision. Junying Zhang: investigation. Qiaowen Zheng: investigation and editing. Tianju Chen: visualization. Feiming Li: editing. Zhixiong Cai: editing. Jingbin Zeng: methodology, investigation and editing. Maosheng Zhang: supervision, project administration, funding acquisition.

Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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References

- G. Huang, Y. Zhou, F. Li, X. Tan, Z. Cai, D. Luo, T. Chen and M. Zhang, *Sens. Actuators, B*, 2021, **347**, 130618.
- J. Song, T. Fang, J. Li, L. Xu, F. Zhang, B. Han, Q. Shan and H. Zeng, *Adv. Mater.*, 2018, **30**, 1805409.
- G. Xing, N. Mathews, S. S. Lim, N. Yantara, X. Liu, D. Sabba, M. Grätzel, S. Mhaisalkar and T. C. Sum, *Nat. Mater.*, 2014, **13**, 476–480.
- Y. Zhou, S. Wu, G. Huang, J. Zeng, B. E. Meteku, X. Tan, H. Lu, F. Li, Z. Cai, X. Wang and M. Zhang, *J. Alloys Compd.*, 2022, **918**, 165565.
- K. Lin, J. Xing, L. N. Quan, F. P. G. de Arquer, X. Gong, J. Lu, L. Xie, W. Zhao, D. Zhang, C. Yan, W. Li, X. Liu, Y. Lu, J. Kirman, E. H. Sargent, Q. Xiong and Z. Wei, *Nature*, 2018, **562**, 245–248.
- J. Ali, Y. Li, P. Gao, T. Hao, J. Song, Q. Zhang, L. Zhu, J. Wang, W. Feng, H. Hu and F. Liu, *Nanoscale*, 2020, **12**, 5719–5745.
- K. Fu, C. T. Nelson, M. C. Scott, A. Minor, N. Mathews and L. H. Wong, *Nanoscale*, 2016, **8**, 4181–4193.
- S. Ghimire and C. Klinke, *Nanoscale*, 2021, **13**, 12394–12422.
- L. Lei, Q. Dong, K. Gundogdu and F. So, *Adv. Funct. Mater.*, 2021, **31**, 2010144.
- M. Palabathuni, S. Akhil, R. Singh and N. Mishra, *ACS Appl. Nano Mater.*, 2022, **5**, 10097–10117.
- S. Akhil, V. G. V. Dutt, R. Singh and N. Mishra, *J. Phys. Chem. C*, 2021, **125**, 22133–22141.
- S. Akhil, V. G. V. Dutt and N. Mishra, *Nanoscale Adv.*, 2021, **3**, 2547–2553.
- L. Protesescu, S. Yakunin, M. I. Bodnarchuk, F. Krieg, R. Caputo, C. H. Hendon, R. X. Yang, A. Walsh and M. V. Kovalenko, *Nano Lett.*, 2015, **15**, 3692–3696.
- X. Li, Y. Wu, S. Zhang, B. Cai, Y. Gu, J. Song and H. Zeng, *Adv. Funct. Mater.*, 2016, **26**, 2435–2445.
- F. Zhang, H. Zhong, C. Chen, X.-G. Wu, X. Hu, H. Huang, J. Han, B. Zou and Y. Dong, *ACS Nano*, 2015, **9**, 4533–4542.
- W. Li, W. Deng, X. Fan, F. Chun, M. Xie, C. Luo, S. Yang, H. Osman, C. Liu, X. Zheng and W. Yang, *Ceram. Int.*, 2018, **44**, 18123–18128.
- T. Chen, Y. Xu, Z. Xie, W. Jiang, L. Wang and W. Jiang, *Nanoscale*, 2020, **12**, 9569–9580.
- H. Luo, Y. Huang, H. Liu, B. Zhang and J. Song, *Chem. Eng. J.*, 2022, **430**, 132790.

- 19 M. T. Clough, K. Geyer, P. A. Hunt, S. Son, U. Vagt and T. Welton, *Green Chem.*, 2015, **17**, 231–243.
- 20 L. Yang, H. Wang, Y.-G. Zu, C. Zhao, L. Zhang, X. Chen and Z. Zhang, *Chem. Eng. J.*, 2011, **172**, 705–712.
- 21 Y. Tan, Y. Zou, L. Wu, Q. Huang, D. Yang, M. Chen, M. Ban, C. Wu, T. Wu, S. Bai, T. Song, Q. Zhang and B. Sun, *ACS Appl. Mater. Interfaces*, 2018, **10**, 3784–3792.
- 22 S. Akhil, V. G. V. Dutt and N. Mishra, *Nanoscale*, 2021, **13**, 13142–13151.
- 23 S. Akhil, V. G. V. Dutt, R. Singh and N. Mishra, *J. Phys. Chem. C*, 2022, **126**, 10742–10751.
- 24 S. Akhil, V. G. V. Dutt and N. Mishra, *ChemNanoMat*, 2021, **7**, 342–353.
- 25 S. Akhil, V. G. V. Dutt and N. Mishra, *Chem. – Eur. J.*, 2020, **26**, 17195–17202.
- 26 A. P. Abbott, G. Capper, D. L. Davies, R. K. Rasheed and V. Tambyrajah, *Chem. Commun.*, 2003, **1**, 70–71.
- 27 Y. Fan, H. Wu, D. Cai, T. Yang and L. Yang, *Sep. Purif. Technol.*, 2020, **250**, 117211.
- 28 A. P. Abbott, D. Boothby, G. Capper, D. L. Davies and R. K. Rasheed, *J. Am. Chem. Soc.*, 2004, **126**, 9142–9147.
- 29 A. Paiva, R. Craveiro, I. Aroso, M. Martins, R. L. Reis and A. R. C. Duarte, *ACS Sustainable Chem. Eng.*, 2014, **2**, 1063–1071.
- 30 Y. Dai, J. van Spronsen, G. J. Witkamp, R. Verpoorte and Y. H. Choi, *Anal. Chim. Acta*, 2013, **766**, 61–68.
- 31 Y. Ma, Y. Li, S. Ali, P. Li, W. Zhang, M. C. R. Rauch, S. J. P. Willot, D. Ribitsch, Y. H. Choi, M. Alcalde, F. Hollmann and Y. Wang, *ChemCatChem*, 2020, **12**, 989–994.
- 32 T.-X. Yang, L.-Q. Zhao, J. Wang, G.-L. Song, H.-M. Liu, H. Cheng and Z. Yang, *ACS Sustainable Chem. Eng.*, 2017, **5**, 5713–5722.
- 33 W. Liu, K. Zhang, J. Chen and J. Yu, *J. Mol. Liq.*, 2018, **260**, 173–179.
- 34 Á. Santana-Mayor, A. V. Herrera-Herrera, R. Rodríguez-Ramos, B. Socas-Rodríguez and M. Á. Rodríguez-Delgado, *ACS Sustainable Chem. Eng.*, 2021, **9**, 2161–2170.
- 35 D. Fuchs, B. C. Bayer, T. Gupta, G. L. Szabo, R. A. Wilhelm, D. Eder, J. C. Meyer, S. Steiner and B. Gollas, *ACS Appl. Mater. Interfaces*, 2020, **12**, 40937–40948.
- 36 F. J. V. Gomez, M. Espino, M. de los Angeles Fernandez, J. Raba and M. F. Silva, *Anal. Chim. Acta*, 2016, **936**, 91–96.
- 37 Q. Hu, H. Yu, S. Gong, Q. Han and W. Wu, *J. Mater. Chem. C*, 2022, **10**, 6002–6008.
- 38 M. Wang, F. Qiao and H. Yan, *Green Chem.*, 2021, **23**, 5179–5188.
- 39 J. Yue, Z. Zhu, J. Yi, H. Li, B. Chen and J. Rao, *Food Chem.*, 2022, **376**, 131943.
- 40 K. Zhu, Q. Wei, H. Li and X. Ren, *ACS Sustainable Chem. Eng.*, 2022, **10**, 2125–2135.
- 41 A. K. Dwamena and D. E. Raynie, *J. Chem. Eng. Data*, 2020, **65**, 640–646.
- 42 M. T. Hoang, N. D. Pham, Y. Yang, V. T. Tiong, C. Zhang, K. Gui, H. Chen, J. Chang, J. Wang, D. Golberg, J. Bell and H. Wang, *Green Chem.*, 2020, **22**, 3433–3440.
- 43 Y. Liu, D. Li, L. Zhang, Y. Chen, C. Geng, S. Shi, Z. Zhang, W. Bi and S. Xu, *Chem. Mater.*, 2020, **32**, 1904–1913.
- 44 Q. Zhong, M. Cao, Y. Xu, P. Li, Y. Zhang, H. Hu, D. Yang, Y. Xu, L. Wang, Y. Li, X. Zhang and Q. Zhang, *Nano Lett.*, 2019, **19**, 4151–4157.
- 45 Y. Liu, X. Zheng, Y. Fang, Y. Zhou, Z. Ni, X. Xiao, S. Chen and J. Huang, *Nat. Commun.*, 2021, **12**, 1686.