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# High efficiency blue light-emitting devices based on quantum dots with core-shell structure design and surface modification

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Blue quantum dot (QD) light emitting diode (QLED) developments are far lagging behind the red and green ones as it becomes difficult to balance charge injection and photo stability than the latter. Here, we introduced a combination of a low band energy shell with better surfactants, which largely meet both abovementioned requirements. Our simulation pinpoints that it is the exposed Se on the QD surface, which causes non-radiative relaxations. By adding tributyl phosphine (TBP), which is a good ligand to Se, we recover photoluminescence quantum yield (PLQY) from less than 8.0% up to above 85.0%. The corresponding external quantum efficiency (EQE) of QLEDs increases from 3.1% to 10.1%. This demonstrates that the low bandgap shell with effective surfactant passivation is a promising strategy to enhance QLED performance.

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## Introduction

Electrically driven light emission of quantum dot light emitting diodes (QLEDs) has attracted considerable interest in recent years due to its promising potentials as a novel display technology.<sup>1–3</sup> It benefits from the merits of foldability, wide color gamut and low manufacturing cost due to its flexible structure, narrow full width at half maximum of quantum dots (QDs), and ink-jet printing process, respectively.<sup>6</sup> Therefore, QLEDs foresee a bright future among the competitive innovative display technologies.

By combining efforts of QD synthetic optimizations and device fabrication improvements during last two decades, the QLED performance gradually enhanced.<sup>1,7,8</sup> During the last few years, with increasing involvement from major display manufacturers, QLED embraced an accelerated development, largely for the red and green ones.<sup>2,3</sup> Blue QLEDs, on the other hand, have far lagged behind mainly due to the synergetic effects of the poor charge injection balance and high applied voltage.<sup>9</sup> While we are preparing this manuscript, Samsung reports ZnSe<sub>x</sub>Te<sub>1–x</sub>-based blue quantum dots with the assistance of the ZnCl<sub>2</sub> surface treatment that achieved high external quantum efficiency (EQE) and extensive operational stability.<sup>10</sup> ZnCl<sub>2</sub> is used to passivate the anionic dangling bonds on the surface in

order to mitigate surface non-radiative traps and also facilitate the charge injection.

With the state-of-art device structure using ZnO nanoparticles as electron transfer layer (ETL) and TFB as hole transfer layer (HTL), energy levels for both electron and hole injections are fixed. In order to balance the charge transfer, one way is to insert a charge block layer. Although it can effectively improve device EQE and operation stability, an additional thin electron blocking layer suffers from reproducibility issues due to the poor control of the film uniformity and thickness.<sup>1</sup> Another way is to manipulate the energy levels of QD core shell structures so as to accommodate the alignment offsets from ETL and HTL. Cao *et al.* utilized a narrow bandgap ZnSe shell to effectively mitigate the hole injection barrier of red QDs.<sup>3</sup> With this approach, it reports an EQE of 15.1% and the T95 device operation lifetime of more than 2300 h with an initial brightness of 1000 cd m<sup>–2</sup>, the longest up to date. Besides red, ZnSe shell has also been successfully utilized in previous studies to facilitate carrier injections for blue QLEDs, further demonstrating its generality.<sup>4,5</sup> Inspired by this shell design, here, we tackle the blue QLED charge problem using a similar approach with necessary surface modifications in order to preserve the photostability.

In previous studies, very often ZnS is used as the outer shell to strongly confine excitonic wavefunctions in blue cores aiming at maintaining high photo stability.<sup>11</sup> However, the large bandgap of the shell impedes both electron and hole from passing into the core, hence compromising device performance. On the other hand, the tentative use of ZnSe as the shell favors the charge injection, but severely deteriorates QD photoluminescence due to the weak confinement. One way to solve

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this issue is to eliminate surface traps with selectively picked surfactants.<sup>6</sup>

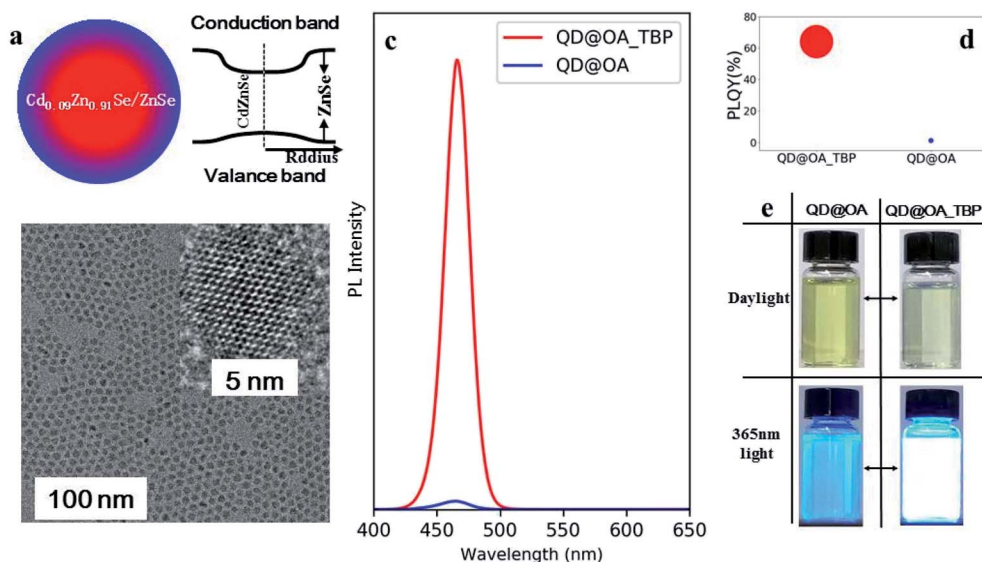
Here, we demonstrate the use of both oleic acid (OA) and alkyl phosphine to passivate cations and anions on the surface, respectively. Blue-colored  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}$  core shells can not only preserve high photoluminescence but also achieve excellent EQE. The pristine OA surfactants mainly passivate dangling bonds on zinc. By addition of alkyl phosphine post synthesis, it can effectively bind on the hole trap-intending selenium and therefore improve photoluminescence from less than 8.0% up to above 85.0%. Furthermore, the corresponding EQE of electrical-excited device increased from 3.1% up to 10.1%. It provides a new insight that the low bandgap shell with pin-pointing surfactant passivation is an effective strategy to enhance the performance QLED.

## Results and discussions

The  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}$  core shells are synthesized following a three-step process. First, ZnSe nuclei is formed by the swift injection of Se into a redundant Zn ion solution, then Cd ions are continuously added to form the  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}$  core before it is completed with the further slow addition of Se to form a ZnSe shell. TEM shows the size of QDs is approximately 6–7 nm with roughly spherical morphology and excellent crystallinity (Fig. 1b). Uniformly alloyed core and smooth interface between the core and shell are guaranteed using both high annealing temperature and sufficiently long cooking duration (Fig. 1a left). Both theoretical<sup>12</sup> and experimental<sup>13</sup> evidence demonstrate that smoothly connected energy levels in QDs can both mitigate Auger recombination and facilitate charge injections (Fig. 1a right). Our synthetic strategy is designed to exclusively accommodate this requirement. The protective ZnS

layer is intentionally eliminated in this case to ease both electron and hole injections.<sup>3</sup> Unsurprisingly, as a result of it, the QDs suffer from decrease in the photoluminescence quantum yield (PLQY) after purification (Fig. 1c blue curve and 1d blue spot). After standard three time washings, the PLQY drops to below 8.0%. This can be elucidated by the unavoidable wave-function extensions to the surface. Our calculation shows that the dangling bonds of Se on the QD surface tend to be hole traps, which in turn create non-radiative relaxation channels. The as-synthesized QDs exhibit decent photoluminescence due to the usage of both oleic acid and trioctyl phosphine (TOP) during synthesis, with former binding to Zn ions and latter to Se. However, after purification, the free ligands were cleaned away as well as the weakly bound surfactants such as TOP, leaving Se exposed. To solve this issue, we reintroduced phosphine ligands during the post purification process to enhance PLQY. Phosphines including TOP and tributyl phosphine (TBP) are good ligands to Se, which are routinely used as precursors in the QD synthesis.<sup>14</sup> By adding a trace amount of phosphine into the QD solution, the PLQY quickly bounced back to 64.0% and with further optimization of the phosphine amount, it can reach above 85.0% (Fig. 1c red curve and 1d red spot). Also, this little amount of ligand can boost photoluminescence but not enough to significantly affect the electrical property of the following light emitting devices, therefore perfectly solving the trap problem. The emission peak is at 465.8 nm with the full width at half maximum (FWHM) of ~21 nm, which is the same as that of the pristine QDs.

In a typical synthesis,  $\text{Zn}(\text{Ac})_2$  (12 mmol), oleic acid (20 ml) and ODE (10 ml) are filled into a 50 ml three-necked round bottom flask. The mixture is stirred and heated to 100 °C before purging with  $\text{N}_2$  for around 30 min to dissolve the salt and remove acetic acid. After it totally dissolves, the solution



**Fig. 1** Colloidal quantum dots. (a) The schemes of  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}$  core shells and its energy levels. (b) TEM images with HRTEM in the inset, the QD are roughly spherical and size is around 6–7 nm. (c) Luminescence peaks comparison between oleic acid (OA) and tributyl phosphine (TBP) (red curve) and OA only (blue curve). (d) The PLQY of OA\_TBP and OA only capped QDs, the sizes of the spots are proportionate to their PLQY values. (e) The images of QD@OA and QD@OA\_TBP under daylight and blue light excitations.



becomes either clear or slightly yellowish in color. Then, after the solution is heated up to 300 °C, the TOPSe solution (3.6 ml with a concentration of 0.67 M) is swiftly injected. 40 s later,  $\text{Cd}(\text{OA})_2$  (1.5 ml with a concentration of 0.2 M) is continuously injected at a rate of  $75 \mu\text{l min}^{-1}$  for 20 min. After the addition, the solution is heated for another 40 min for annealing. It is then followed by the slow addition of TOPSe (1.34 ml with a concentration of 0.67 M) at a rate of  $45 \mu\text{l min}^{-1}$  for 30 min. After the addition, the solution is naturally cooled to room temperature and subjected to washing process.

Due to the high surface to volume ratio of quantum dots, the trap states caused by the dangling bonds of surface atoms cannot be ignored. These mid-gap states can act as non-radiative recombination centers.<sup>15,16</sup> To demonstrate from a theoretical perspective, we performed simulations *via* density functional theory (DFT). The surface of quantum dots was simulated to understand the trap states caused by the dangling bonds of the surface atoms. According to our quantum dot structure, the (100) wurtzite ZnSe is used as the surface model. The density of states (DOS) of passivated and un-passivated surfaces is compared. As shown in Fig. 2, without passivation, a significant amount of defect states are presented within the bandgap. These defect states contribute to the Auger recombination, thus decreasing the quantum efficiency. After passivating the Zn and Se dangling bonds on the surface, 84% of the defect states in the mid-gap disappear as shown by the red line in Fig. 2. Therefore, choosing the appropriate ligand to passivate the QD surface is critical.

In previous studies, different ligands have been used to passivate the surface of QDs in order to achieve a better device efficiency.<sup>6,17–19</sup> In our case, we demonstrate that the OA and alkyl phosphine are complementary with each other to bind both Zn and Se on the QD surface. In particular, alkyl phosphine forms a coordination bond with Se and is regularly used as the Se precursor for the metal chalcogenide QD synthesis. To demonstrate this claim, phosphine ligands are introduced into the purified blue QD solution, and the PLQY of the colloidal

quantum dots before and after adding the ligands is investigated. As shown in Fig. 3a, as the phosphine ligand concentration increases, the PLQY of the  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}@OA$  blue QDs is significantly improved due to the effective passivation of phosphine ligands on under-coordinated Se anion sites,<sup>20,21</sup> corresponding well with the simulation results above, showing the major trap states of the  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}@OA$  blue QDs are the Se anion defect sites. Moreover, phosphine ligands with the alkyl chain of different length affect the PLQY recovery. With TOP that has three octyl branches, the PLQY of the QDs rise to ~74% at the optimal TOP ligand's quantity, while TBP that has three butyl branches increase the PLQY of the blue QDs to almost 90% under the same condition. The smaller molecular size of the TBP ligands and the corresponding higher surface ligand coverage on the blue QDs is the reason to cause this difference and lead to a better surface passivation. Moreover, other types of ligands with different function groups, such as pristine oleic acid (OA) and octanethiol (OT), had also been tested with the  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}$  blue QDs, and the optimal PLQY of the blue QDs after surface passivation with each type of ligands has been summarized in Fig. 3b. It can be seen that the surface ligands with carboxylate or thiol functional groups do not recover the PLQY of the  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}@OA$  blue QDs as effective as alkyl phosphines. It is well known that the carboxylate and thiol ligands prefer surface cation sites passivation, but only have limited effect on anions.<sup>22,23</sup> Therefore, the surface ligands with the carboxylate group or thiol groups do not effectively passivate with the Se defect sites, and the major trap states of the  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}@OA$  blue QDs as shown in the simulation results, resulting in an unsatisfied PLQY recovery of the blue QDs.

To illustrate the influence of ligands on the device performance,<sup>23</sup> the blue QLED devices ( $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}@OA$  and  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}@OA\_TBP$  devices) were prepared. The QLED device configuration is as follows: glass/ITO (50 nm)/poly(3,4-ethylenedioxythiophene) polystyrene sulfonate (PEDOT: PSS 35 nm)/poly(9,9-dioctylfluorene)-*co*-(*N*-(4-butylphenyl)diphenylamine) (TFB 25 nm)/QDs (20 nm)/ZnO (35 nm)/Al (100 nm) (Fig. 4a). For the organic-inorganic hybrid QLED devices, all functional layers were spin-coated except the anode and cathode.<sup>24</sup> A glass substrate with patterned 50 nm indium tin oxide (ITO, sheet resistance is about  $50 \Omega \square^{-1}$ ) anode were cleaned by ultrasonic treatment in a tergitol/deionized (DI)-water solution, DI-water, and isopropanol, respectively, followed by treatment with under ultraviolet (UV)-ozone for 15 min. A PEDOT: PSS layer as the hole injection layer (HIL) was spin-coated onto the pre-cleaned ITO substrates at 5000 rpm and then baked at 150 °C under air condition. TFB dispersed in chlorobenzene was spin-coated at 2000 rpm and annealed at 150 °C under a  $\text{N}_2$  atmosphere. After that, one QD and ZnO nanoparticle layer were sequentially deposited from the octane and ethanol solution in the glove-box, respectively, *via* the spin-coating method and annealed in the same environment at 80 °C. The Al cathode was thermally deposited in a vacuum chamber under the chamber pressure less than  $10^{-6}$  torr through a shadow mask to obtain active areas of  $2 \times 2 \text{ mm}^2$ . Finally, the blue QLED devices were encapsulated using a UV-

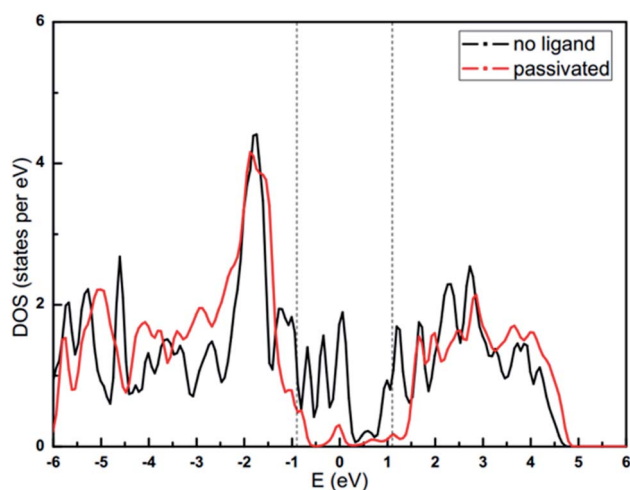
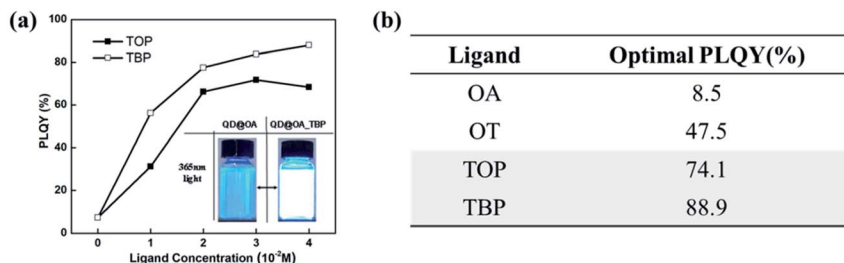


Fig. 2 The DOS of the (100) ZnSe surface with and without ligand passivation. The figure is reprinted from ref. 6.

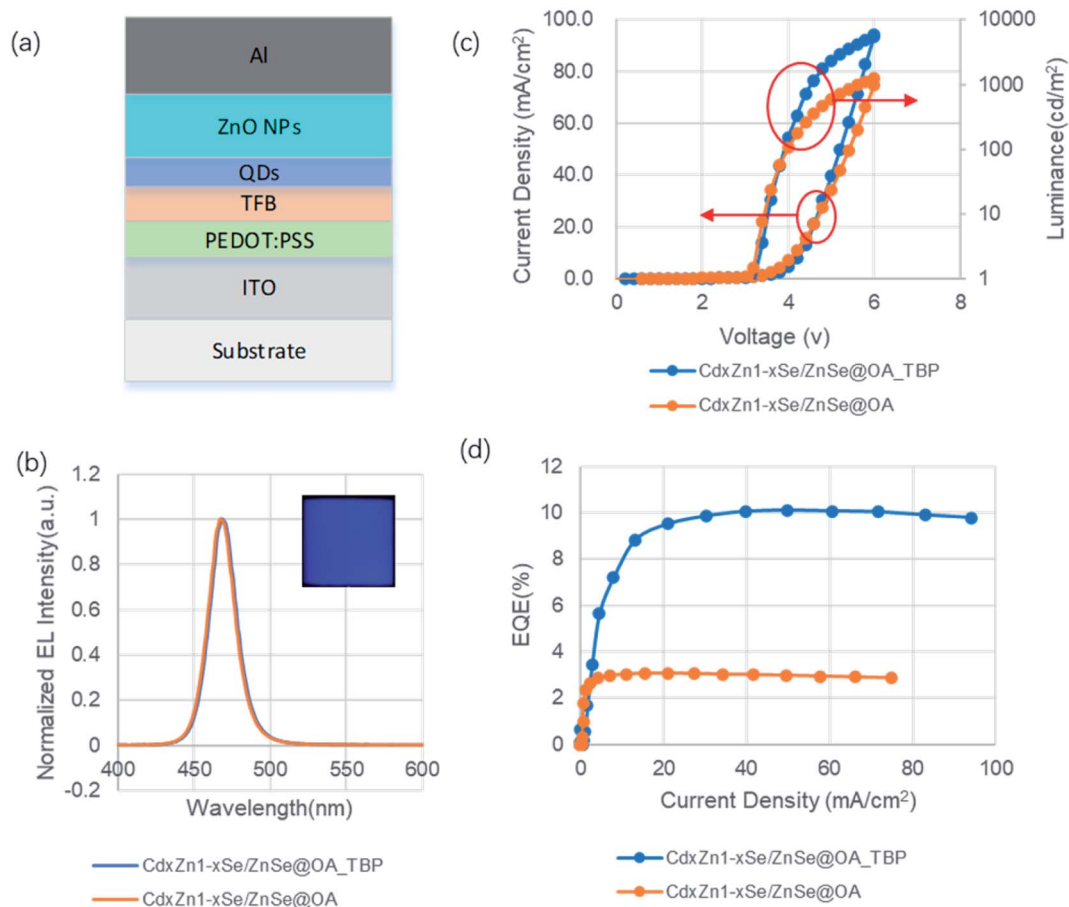




**Fig. 3** The effect of phosphine ligands on PLQY of the  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}@OA$  blue QDs. (a)  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}@OA$  blue QD PLQY after adding TOP or TBP with different concentrations of phosphine ligands, respectively. The inset photograph taken under illumination at 365 nm, from left to right: before and after adding TBP ligands into the purified blue QDs. (b) The optimal photoluminescence quantum yield of the  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}@OA$  blue QDs after surface passivation with different coordination groups of surface ligands.

epoxy resin and a cover glass in the  $\text{N}_2$  atmosphere. A digital source meter (Keithley 2400) was used to operate the devices and measure the  $J$ - $V$  characteristics. Electroluminescence spectra were recorded using an Ocean Optics USB 2000+ spectrometer. Light intensities of devices were measured with a Keithley 6485 Picoammeter together with a calibrated silicon detector (Edmund). EQEs were calculated from the luminance, current density, and EL spectra with the assumption of the Lambertian emission pattern of all devices.<sup>25,26</sup>

The normalized electroluminescence (EL) spectra of the devices based on  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}@OA$  and  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}@OA\_TBP$  are shown in Fig. 4b. To simplify the nomination, hereafter the former is named as QD@OA and the latter is QD@OA\_TBP. Both QLED devices exhibited the same narrow EL emission at a peak wavelength of 469 nm with a FWHM of 21 nm. While FWHM is preserved, the EL peak position redshifted 3.2 nm compared to that in the solution. This is mainly due to the increasing exciton wavefunction



**Fig. 4** Device structure and performance of the blue device: (a) bottom emission device structure, (b) normalized EL spectrum of QD@OA and QD@OA\_TBP, respectively, (c) the current density and luminance of QLEDs, and (d) EQE vs. current density.





communications of QD after they are closed packed on the solid film. Pure blue emission is observed in the inserted photograph of QLED fabricated using QD@OA\_TBP. Fig. 4c shows the current density ( $J$ )–voltage ( $V$ )–luminance ( $L$ ) characteristics of QD@OA and QD@OA\_TBP devices. Both devices show the same turn-on voltage ( $V_T$ ) of 3.2 V defined as the required operation voltage for achieving the luminance of  $1 \text{ cd m}^{-2}$ . The  $J$ – $V$ – $L$  characteristics are not that much different at a low  $J$  region ( $<4 \text{ V}$ ). However, the luminance of the QD@OA\_TBP device increases rapidly at the  $>4 \text{ V}$  region. The QD@OA\_TBP device shows about four times higher luminance ( $2322 \text{ cd m}^{-2}$ ) than that of the QD@OA device ( $590 \text{ cd m}^{-2}$ ). The external quantum efficiency ( $\eta_{\text{EQE}}$ ) of the QD@OA and QD@OA\_TBP devices are exhibited in Fig. 4d. The latter shows the maximum  $\eta_{\text{EQE}}$  value of 10.1%, which is about 3.3-fold higher than that of the former device. Similarly, high EQE was also reported in the literature *via* functional layer modifications.<sup>4,5</sup> To note, all the device fabrication processes are kept the same, and the only difference is to add trace amounts of TBP into the QD solution for the QD@OA\_TBP device. Compared to the OA only case which favours Zn, TBP complements the passivation of Se, which further mitigates the non-radiative relaxation centers. Besides, in contrast to TOP, which has large steric hindrance and high boiling point, the TBP molecule is small and volatile, which indicates a more effective passivation of QD and easy removal for the excessive ones during the QD film preparation process, both of which are beneficial for the device characteristic improvement.

## Conclusions

In summary, the present study demonstrates the blue QLEDs with high EQE and low threshold voltage. This is achieved through the synergistic effects of tuning the energy level of QD to match adjacent functional layers and effective ligand passivation to largely eliminate non-radiative traps on the surface. In particular, we introduce TBP to QD as additional surfactants, which complement the original OA to bind on the dangling bonds of Se on the surface. Compared to other long chain alkyl phosphines, TBP has the merits of low steric hindrance and high volatility, which favor more passivation to surface Se sites and ease for nonbinding ones to remove during device fabrication. TBP-modified QDs witness the PLQY recovery from less than 8.0% up to above 85.0%, and compared to the pristine QD@OA, the EQE of QD@OA\_TBP devices has 3.3-fold enhancement from 3.1% to 10.1% with a threshold voltage as low as 3.2 V. This demonstrates that a low bandgap shell design with effective surfactant passivation is a promising strategy to enhance the performance of QLED.

## Experimental details

### Procedures for the blue QD preparations

**Stock solutions.** TOP-Se (1.0 M) is prepared by dissolving 5 mmol Se powder into a 5 ml TOP solvent.

$\text{Cd}(\text{OA})_2$  (0.2 M) is prepared by reacting CdO 1.0 mmol with OA 1 ml and ODE 4.0 ml at  $250^\circ\text{C}$ . The solution is bubbled with  $\text{N}_2$  at  $80^\circ\text{C}$  for 20 min before cooling down.

### Reaction procedures

(1)  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}$  core preparation.  $\text{Zn}(\text{ac})_2$  (5.0 mmol), OA (5 ml) and ODE (20 ml) are added into a 100 ml round bottom three-neck flask. The mixture is bubbled with  $\text{N}_2$  at  $130^\circ\text{C}$  for 1 h before heating up to  $310^\circ\text{C}$ . When the temperature is stable, the TOPSe (0.8 ml) and DPP (0.3 ml) mixture is swiftly injected into the solution, triggering the burst of ZnSe nuclei. The temperature drops down to  $300^\circ\text{C}$  and the solution continues to cook for 50 s before  $\text{Cd}(\text{OA})_2$  (0.4 ml) is rapidly injected into the solution. The  $\text{Zn}_{0.9}\text{Cd}_{0.1}\text{Se}$  core is furthermore cooked for 1 h at  $300^\circ\text{C}$ .

(2)  $\text{Cd}_{0.1}\text{Zn}_{0.9}\text{Se}/\text{ZnSe}$  shell coating. With the hot solution in section 1 at  $300^\circ\text{C}$ , TOPSe (0.5 ml) is slowly added for 13 min 20 s at constant rate using a syringe pump. With the excess  $\text{Zn}(\text{OA})_2$  in the solution, the ZnSe shell is gradually formed during this process. Then, the solution is cooled down to room temperature and subjected to cleaning by going through 3 times of precipitation with ethanol and dissolution with octane cycle. Then, the QD is stored in an octane solution for the following use.

### QD solution PLQY measurement

Blue QD solution PLQY is measured *via* the integrated sphere method using Edinburgh Instruments F55 SC-30. The instrument includes an F55 fluorescence plus SC-30 integrated sphere module. The procedures to perform the measurements are following:

The instrument is first to be turned on and warmed up for 30 min before the operation. Next step is to set the excitation and emission wavelengths both at 465 nm, and tune the both bandwidths to make sure signal rate reaching  $1 \times 10^6$  cps. The third is to set the emission range from 400 up to 650 nm with the step of 0.2 nm, dwell time of 0.2 s and the number of scan of 1. Also, the excitation wavelength was set at 360 nm. For the fourth step, the cuvette with the octane solvent was put into the center of the integrated sphere and the spectrum was recorded as the control. Then, the cuvette with the octane solution of QD with absorbance at the excitation wavelength around between 0.1 and 0.15 is to replace the previous control cuvette and perform the same scan. By subtracting the later with former, PLQY can be obtained.

### Device characterizations

The device testing system is composed of a high-precision digital source meter, high-performance optical spectrometer and optical fiber (inner diameter 50  $\mu\text{m}$ ), device testing probe and data acquisition system. The  $J$ – $V$  characteristics of quantum dot devices were measured under ambient condition using a digital source meter (Keithley 2400) measuring the sweeping voltages and currents. The maximum voltage was set up 6 V and the voltage variable step was set up 0.2 V during measuring. Simultaneously, light intensities were measured on a Keithley 6485 Picoammeter together and a calibrated silicon detector



(Edmund). Luminance was calibrated using a photometer (Spectra Scan PR655) with the assumption of the Lambertian emission pattern of all devices. Electroluminescence spectra were recorded using a high performance Ocean Optics USB 2000+ spectrometer.

## Conflicts of interest

There are no conflicts to declare.

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