Abstract: Hybrid electroluminescence (EL) devices with mixed emission layers (EMLs) were developed to achieve balanced peaks from red quantum dots (QDs) and blue phosphorescent small molecules. The EML was prepared by mixing QDs and small molecules in an organic solvent, and then adding polystyrene to disrupt the aggregation of the host material. These unique bichromatic devices exhibited two distinct EL peaks with similar intensities over 10 V, and the voltage dependent EL spectra were investigated systematically. These hybrid EL devices showed a maximum luminance of 1057.7 cd/m² and a current efficiency of 2.45 cd/A. These results indicate that the unique mixed EMLs have potential for use as white devices using fewer fabrication steps.

Keywords: colloidal quantum dots; organic small molecules; electroluminescence

1. Introduction

Colloidal quantum dots (QD) have an adjustable emission wavelength, through size-control, with high color purity and their outer organic ligands make the fabrication process cost effective by the solution process [1–4]. For this reason, QDs are promising semiconductor emitters for the future display industry, and the performance of quantum dot light-emitting diodes (QLEDs) is now comparable with that of organic light-emitting diodes (OLEDs) because of the advantages of a narrow emission line and enhanced electroluminescence (EL) characteristics [5–7].

Compared to monochromatic devices, studies on white-emitting QLEDs have been limited owing to the structural complexity of the device and the difficulty in simultaneous multi-emission. The stacked tandem structure is considered the way to achieve white emission as in the case of white-emitting OLEDs. However, its complicated fabrication requires further attention because of its in-between buffer layers and multiple emission layers (EMLs) [8,9]. Other attempts using mixed EMLs using more than two QDs have been made, but the inevitable energy loss to adjacent QDs is difficult to control [10,11]. Studies about organic and inorganic hybridization for white EL have been limited. White light-emitting diodes with a hybrid polymer and QDs have been reported, but the luminance was less than 500 cd/m² [12,13]. Recently, our group reported highly efficient white QLEDs with blue and green mixed QDs and red organic phosphorescent molecules. They had a maximum luminance of 20,453 cd/m² and an external quantum efficiency (EQE) of 9.19% [14]. However, the fabrication process for two EMLs consists of two different deposition methods—spin coating and thermal evaporation.

Although vacuum deposition is the most commonly employed process for the fabrication of EL devices, the equipment is very expensive, and the efficiency of the deposited material is as low as approximately 10%. These problems can be critical in large panels. Recently, the solution process in OLEDs has received considerable attention as a result of the low manufacturing cost, and
high utilization efficiency of the deposited material by minimizing the use of materials, and their performance is comparable with that of conventional, vacuum-processed devices [15]. Shinar et al. successfully fabricated a green OLED by spin coating one of the solution processes and achieved an efficiency of 69 cd/A [16].

Herein, we report hybrid EL devices with solution-processed EMLs using red QDs and blue phosphorescent molecules. Because blue remains a weak point in QDs, we chose QDs for red emission and small molecules for blue emission. Firstly, small-molecule host materials and blue dopant materials were dispersed in an organic solvent, and then a unique solution-processed EML was prepared by mixing it with red QDs. As most small-molecule materials exhibit poor film-forming properties owing to molecular aggregation, a small amount of polystyrene (PS) was added to the mixed solution to suppress this. Hybrid EL devices with the addition of small amounts of PS showed better optoelectrical performance with stable film morphology.

2. Materials and Methods

2.1. Synthesis of Red QDs

Red emitting CdSeS/ZnS QDs were prepared by the synthesis procedure in the recent publication of Yang’s group [11]. Briefly, a mixture of 1.1 mmol of Cd acetate and 5 mL of oleic acid (OA) was degassed and heated to 150 °C, followed by the addition of 20 mL of 1-octadecene (ODE) and further heating to 305 °C. At that temperature, an anionic stock solution, prepared by dissolving 0.27 mmol of Se and 0.03 mmol of S in 0.3 mL of trioctylphosphine (TOP), was quickly injected. And then the reaction was maintained for 90 s for CdSeS core growth, followed by a dropwise addition of 1.7 mmol of 1-octanethiol. For the ZnS shelling, a Zn solution with 2.86 mmol of Zn acetate dihydrate dissolved in 4 mL of OA and 1 mL of ODE was first injected and a S solution with 6.75 mmol of S dissolved in 3.5 mL of TOP was then injected. The absorption and photoluminescence (PL) spectra and X-ray diffraction (XRD) patterns of red CdSeS/ZnS QDs are shown in Figure 1 [17].

![Figure 1. (a) Absorption and photoluminescence spectra and (b) X-ray diffraction (XRD) patterns of CdSeS/ZnS QDs.](image)

2.2. Fabrication of QLEDs

A patterned indium–tin oxide (ITO) glass was cleaned by sonication using isopropyl alcohol and deionized water. The surface of the pre-cleaned ITO glass was subjected to an ultraviolet (UV) plasma treatment for 15 min. For a device with an inverted structure, the electron transport layer (ETL) of ZnO nanoparticles (NPs) in ethanol was spin-coated on the cleaned ITO glass. The organic molecule of 1,3-Bis(N-carbazolyl)benzene (mCP) was used as the host material and Bis[2-(4,6-difluorophenyl)pyridinato-C2,N](picolinato) iridium(III) (Flrpic) was used as the dopant for the mixed EML. Then, mCP:Flrpic was mixed in a 10:1 weight ratio and dissolved in toluene to be mixed again with red CdSeS/ZnS CQDs. The QD solution and organic solution were mixed in 1:3 and

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1:5 volume ratios, respectively. The polystyrene was dissolved separately in chlorobenzene to prepare a solution of 20 mg/mL concentration. Afterwards, a 10 wt.% polystyrene solution was added. To deposit EML, the solution of mixed QDs, organic phosphors, and polystyrene was spin-coated on the ZnO layer at a spin rate of 2000 rpm for 5 s. After the solution process, 4,4′-Bis(N-carbazolyl)-1,1′-biphenyl (CBP) was directly deposited onto the EML to a thickness of 40 nm, using vacuum evaporation. Then, the cathode (MoO$_3$ and Al) was deposited by thermal evaporation to thicknesses of 10 nm and 100 nm, respectively. Finally, the devices were encapsulated by an encapsulation glass using an UV sealant.

2.3. Characterization

The current density–voltage–luminance (J–V–L) characteristics of the fabricated devices were measured using a spectroradiometer (Minolta CS 2000, Tokyo, Japan) with a Keithley 2400 source meter under ambient conditions. From these J–V–L measurements, the changes in the devices’ luminance, current efficiency and electroluminescence spectrum as a function of the applied voltage were studied systematically. The surface roughness was measured by atomic force microscopy (AFM, PSIA XE-100, Neuchatel, Switzerland) was used for non-contact mode measurement.

3. Results and Discussion

Figure 2 illustrates a schematic diagram of hybrid EL devices, which consists of ITO/ZnO NPs (50 nm)/red QDs + mCP:Flrpic (mixed EML) (30 nm)/CBP (40 nm)/MoO$_3$ (10 nm)/Al. The inverted device structure was chosen for efficient injection of electrons and holes into the EML. ZnO NPs are representative inorganic materials as an ETL for QLEDs, because their conduction band minimum is well-matched with the work function of the ITO electrode and they are robust against various organic solvents [18]. In this study, a unique mixed EML was designed for simultaneous red and blue emissions from the QDs and Flrpic, respectively. It is well known that phosphorescent OLED devices have a host-dopant system to efficiently transfer exciton energy, and mCP was selected as the host for the blue emitting dopant, Flrpic. However, mCP is mainly used in conventional, vacuum-processed OLEDs owing to the crystallization during solution processing caused by the low glass transition temperature and symmetric molecular structure [15]. To diminish this aggregation, the electrically isolating PS was dissolved separately and then added to the mixed EML.

![Figure 2. Schematic of hybrid electroluminescence (EL) devices with mixed emission layers (EMLs) of red quantum dots (QDs) and blue small molecules.](image-url)

Figure 3 shows the EL spectra of hybrid EL devices with mixed EMLs. Two distinct peaks (blue from Flrpic and red from QDs) are simultaneously observed at an applied voltage of 10 V. Different mixing ratios of QD and organic solution in the EML were compared, and similar EL intensities from red QDs and blue Flrpic were obtained using a mixing ratio of 1:5. As QDs with organic ligands are
more stable than FIrpic in the organic solvent and the emission from FIrpic needs an exciton energy transfer from mCP, it is reasonable to conclude that more FIrpic is needed for balanced EL peaks.

Figure 3. EL spectra of (a) the monochromatic EL device with only 1,3-Bis(N-carbazolyl)benzene (mCP):Bis[2-(4,6-difluorophenyl)pyridinato-C2,N](picolinato) iridium(III) (FIrpic) and (b) the bichromatic hybrid EL device with mixed EMLs at 10 V.

Figure 4 shows the optoelectronic properties of hybrid EL devices with the mixed EML (red QDs:blue FIrpic = 1:5) to verify the role of the PS additive. In Figure 4a, the hybrid EL device without PS shows a constant ohmic trend with increasing applied voltage. After adding PS in the mixed EML, the current density–voltage curve shifted to lower currents, because electrically isolating PS disrupted the charge carrier transport [19]. Therefore, it leads to an increment of both the turn-on voltage (from 3.5 to 4.5 V) and current efficiency (from 0.33 to 2.45 cd/A), respectively, as shown in Figure 4b,c. The peak luminance of 1057.7 cd/m² was achieved at 13 V (corresponding to a current density of 293 mA/cm²) and current efficiency of 2.45 cd/A were obtained at a current density of 0.13 mA/cm².

Figure 4. (a) Current density–voltage; (b) luminance–voltage; and (c) current efficiency–current density characteristics of monochromatic EL devices with mCP:FIrpic and hybrid EL devices with mixed EMLs and added PS.
Figure 5 shows the AFM images of the mixed EML (red QDs:blue FIrpic 1:5) to investigate the effect of added PS. Prior to fabrication of the whole QLEDs, the mixed EML was deposited on the ZnO/ITO glass to compare the surface roughness with and without PS addition. The surface roughness of the mixed EML decreased from 1.627 to 0.733 nm with the PS addition because of the effective suppression of the mCP aggregation [19].

![AFM images](image)

(a)

(b)

**Figure 5.** Atomic force microscopy (AFM) images of mixed EMLs on the ZnO/ITO glass: (a) without PS ($R_q = 1.627$ nm) and (b) with PS ($R_q = 0.733$ nm).

The spectral variation in the EL spectra and the shift in the Commission Internationale de l’Eclairage (CIE) coordinates of hybrid EL devices with the mixed EML (red QDs:blue FIrpic 1:5) at different voltages are shown in Figure 6. In Figure 6a, the EL intensity from blue FIrpic is approximately half that of the red QDs and the peak intensity was observed at 6 V. As shown in Figure 4a, an extremely high current density of hybrid EL devices without PS leads to unstable optoelectrical characteristics, such as early turn-on and sudden turn-off. Since the EL peak from red QDs is dominant and the intensity ratio between blue and red peaks is similar to increasing voltage, the CIE coordinate mainly stays in the red region with a small shift in Figure 6c. On the contrary, the hybrid EL devices with PS show balanced EL peaks above 10 V from both blue FIrpic and red QDs. The CIE coordinate moves to the white region with increasing voltage, as a result of the later blue emission of FIrpic. This is attributed to the large bandgap of FIrpic leading to effective emission in the higher voltage region.
Although the performance of the hybrid EL devices with the mixed EML does not match with the high-performance of recent QLEDs, these hybrid EL devices demonstrate a novel method to create white light over the traditional tandem structure. QDs are easy to apply to solution processes, however, efficient blue emission requires further work. Small molecules have been well-studied for EL applications, but the solution process is not the main process for practical manufacturing. While the hybridization of organic and inorganic materials for white emission can take advantage of each, the complicated interface between them results in low performance and unstable devices [12,13,20]. In this work, the present hybrid EL devices successfully showed two distinct emissions simultaneously from the mixed EML and the balanced EL peaks under higher applied voltage were achieved with the addition of PS. Since the limitation of charge injection into QDs and small molecules are different, more precise optimization is required for white emission with high color rendering index. Combining the advantages of QDs and small molecules in the solution based EML can reduce the fabrication cost and give the flexibility for future large-area white emitting EL devices.

4. Conclusions

Hybrid EL devices with mixed EMLs employing QDs and small molecules were successfully fabricated. The EL spectra showed a distinct blue emission from FIrpic and a distinct red emission from QDs simultaneously, and similar EL intensities were observed at a ratio of 1:5 (QDs:FIrpic). After
adding PS to mCP, the hybrid EL devices had a maximum luminance of 1057.7 cd/m², while emitting the balanced white light. Stable optoelectrical properties were obtained because electrically isolating PS effectively hinders the flow of charge. The mixed EML prepared in the solution process showed balanced peaks from inorganic QDs (red) and organic mCP:F1rpic (blue), which is a promising result for solution based white emitting EL devices.

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