



## The effect of doping iodine on organic light-emitting diode

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### Abstract

The performances of organic light-emitting diodes (OLEDs) with the configuration Al/Alq<sub>3</sub> (Aluminum Tris-(8-hydroxyquinoline))/TPD(N,N'-diphenyl-N,N'-bis-(3-methylphenyl)-1,1'-bipheny-4,4'-diamine)/ITO have been significantly improved by doping iodine (I<sub>2</sub>) on both Alq<sub>3</sub> and TPD layers. The luminance is promoted from 2800 cd/m<sup>2</sup> without doping to 8000 cd/m<sup>2</sup> with I<sub>2</sub> doping under bias 10 V. Additionally, the driving voltage (@100 cd/m<sup>2</sup>) was reduced from 7.5 V without doping to 5.2 V with I<sub>2</sub> doping. We attribute the promotions to the reduction of the electron and hole injection energy barrier at Al/Alq<sub>3</sub> and TPD/ITO interfaces and the expansion of trap energy states beneath the LUMO of Alq<sub>3</sub> generated by I<sub>2</sub> doping. The mechanism is illustrated comprehensively with a schematic energy diagram model and nicely supported with photoluminescence (PL), electroluminescence (EL) spectra and other experimental results.

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### 1. Introduction

Organic light emitting diodes (OLED) have been widely applied on cell phone display, liquid-crystal-display and television for the advantages of low power dissipation, wider vision angle, high

luminescence, shorter response time and simplified fabrication [1]. In the past, the technologies to improve the performances, such as electroluminescence efficiency and driving voltage, have been concentrated on lowering barrier height on metal contact [1,2], using high luminescence organic materials [1,3–7] and buffer layer or inter-layers [8–12]. Even these technologies could enhance the performance of OLED, however they also complicate the fabrication process or need some special doping materials [13]. Therefore, there are still

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needed the simpler methods or more common doping materials.

In general, for the bi-layer OLED as shown in the inset of Fig. 1, Al/Alq<sub>3</sub> (Aluminum Tris-(8-hydroxyquinoline))/TPD(N,N'-diphenyl-N,N'-bis-(3-methylphenyl)-1,1'-bipheny-4,4'-diamine)/ITO is adopted to study for its simple structure, where Alq<sub>3</sub> and TPD are electron transport layer (ETL) and hole transport layer (HTL), respectively [14]. To reduce the driving voltage, the common approach is to lower the barrier at Al/Alq<sub>3</sub> interface, for the barrier of Al/Alq<sub>3</sub> contact is larger than the barrier of TPD/ITO [15] and the current is controlled by electrons injection into Alq<sub>3</sub> [16]. In other word, lowering Al/Alq<sub>3</sub> barrier is more effective than lowering the TPD/ITO barrier [17–19]. On the other hand, it has been evidenced that the incorporation of iodine (I<sub>2</sub>) on organic material could increase the conductivity [20]. Hence, in this study, we lowered Al/Alq<sub>3</sub> barrier with doping iodine on ETL and improve the output luminance by doping iodine (I<sub>2</sub>) on both ETL and HTL layer simultaneously. The doped I<sub>2</sub> molecules on ETL layer generate traps states beneath the lowest unoccupied molecular orbital (LUMO) of Alq<sub>3</sub> [15,21], thus expanding the excited band of LUMO and lowering the Al/Alq<sub>3</sub> barrier for electron to inject. Compared to the reported methods [1–12], the technique possesses the advantage of low cost, convenience and be compatible to the current OLEDs fabrication process. Furthermore,

a schematic energy diagram model has been employed to illustrate the improving mechanism comprehensively.

## 2. Device design, fabrication and measurement

The bi-layer OLEDs samples with the configuration of Al/Alq<sub>3</sub>/TPD/ITO, as schematically described in the inset of Fig. 1, were prepared in the glass substrates pre-coated by indium tin oxide (ITO) with sheet resistance  $R_s \leq 10 \Omega/\text{square}$  and work function of 4.9 eV. After ultrasonic cleaning in H<sub>2</sub>O–H<sub>2</sub>O<sub>2</sub>–NH<sub>3</sub>OH solution, the substrates were taken into a stain steel chamber which then be evacuated to  $1 \times 10^{-5}$  Torr to deposit TPD as HTL. The thickness of TPD layers is 500 Å. Next, Alq<sub>3</sub> with thickness of 500 Å as ETL and Al with thickness of 1500 Å as cathode were evaporated sequentially in the chamber with  $1 \times 10^{-5}$  Torr. In this work, OLED samples were doped I<sub>2</sub> by evaporation of Alq<sub>3</sub> on ETL with various weight ratios I<sub>2</sub> powder (i.e. Alq<sub>3</sub>/I<sub>2</sub> = 1/1, 1/10) and on HTL by evaporation of TPD with weight ratio of 10/1 I<sub>2</sub> powder (i.e. TPD/I<sub>2</sub> = 1/10) respectively. On the other hand, HP4156 and TOPCON BMP were used to measure the electrical characteristics and output luminance, respectively. Additionally, the photoluminescence (PL) and electronluminescence (EL) were measured with Fluorolog-3 Fluorescence on device samples.

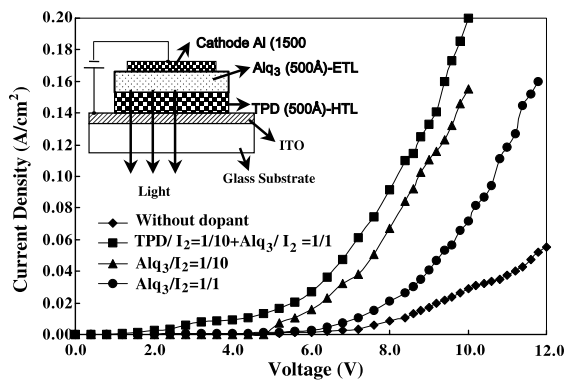


Fig. 1. The  $J/V$  curves of bi-layer OLEDs with Alq<sub>3</sub> or both Alq<sub>3</sub> and TPD simultaneously doped with doping different I<sub>2</sub> in weight ratio. The inset presents schematic structure diagram of OLEDs.

## 3. Results and discussion

Fig. 1 gives the measured current density–voltage ( $J/V$ ) curves with various weight ratios as parameter. With doping I<sub>2</sub> (i.e. Alq<sub>3</sub>/I<sub>2</sub> = 1/1), the current densities of OLED are improved in comparison to that without doping especially under large bias. Also, the improvement is enhanced with increase of dopant weight ratio, i.e., Alq<sub>3</sub>/I<sub>2</sub> = 1/10 is better than Alq<sub>3</sub>/I<sub>2</sub> = 1/1. For example, the current density with Alq<sub>3</sub>/I<sub>2</sub> = 1/10 doping increases to 67 mA/cm<sup>2</sup> and 155 mA/cm<sup>2</sup> from 9 mA/cm<sup>2</sup> and 29 mA/cm<sup>2</sup> for OLED without dopant under 8 V and 10 V biases, respectively. We attribute the increase of current density to the lowering of

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