## Supporting Information

## Prolonging the Lifetime of Quasi-2D Perovskite Blue LEDs via DMAcPA Doping for Defect Passivation

Yu-Chuan Huang<sup>1</sup>,<sup>†</sup>, Chien-Cheng Li<sup>1</sup>,<sup>†</sup>, Tzu-Yu Huang<sup>1</sup>, Yu-Hsuan Lai<sup>1</sup>, Xin-Kai Gao<sup>1</sup>, Jia-Xin Li<sup>2</sup>, Chang-Hua Liu<sup>2</sup>, Hao-Chung Kuo<sup>3</sup>, Ray-Hua Horng<sup>1</sup>, Chih-Shan Tan<sup>1\*</sup>

<sup>1</sup> Institute of Electronics, National Yang Ming Chiao Tung University, Hsinchu 30010, Taiwan

<sup>2</sup> Institute of Photonics Technologies, National Tsing Hua University, Hsinchu 30013, Taiwan

<sup>3</sup> Department of Photonics and Graduate Institute of Electro-Optical Engineering, College of

Electrical and Computer Engineering, National Yang Ming Chiao Tung University, Hsinchu 30010, Taiwan

† These authors contributed equally to this work.



Figure S1. PL spectra of perovskite thin films with and without DMAcPA modified.



**Figure S2.** Binding energy spectra of pristine and doped (target) films: Normalized spectra of (a) Cs 3d, (b) Pb 4f and P 2p, and (c) Br 3d and Cl 2p.



**Figure S3.** FTIR spectra of pristine and doped (target) films, compared with bare DMAcPA on glass.



Figure S4. Tauc plot of PeLEDs: (a) PEDOT:PSS, and (b) TPBi.



**Figure S5.** UPS spectra of PeLEDs: (a) PEDOT:PSS, (b) Pristine, (c) Doped (target), and (d) TPBi. A He I source with a photon energy of 21.22 eV was used for excitation in the UPS measurements. The calculated valence band energies were 5.22 eV for PEDOT:PSS, 6.17 eV for pristine, 6.07 eV for doped (target), and 6.20 eV for TPBi.



Figure S6. CIE coordinates of pristine and doped (target) PeLEDs at maximum luminance.



**Figure S7.** Temperature-dependent PL measurements and optical characteristics: (a) A schematic diagram of the optical setup for temperature-dependent PL measurements. (b) and (c) Exciton binding energy ( $E_b$ ) for pristine and doped (target) films obtained from the relationship between PL intensity and 1/T. (d) and (e) Huang-Rhys factor (S) for pristine and doped (target) films obtained by fitting the profile of FWHM vs. *T*.



Figure S8. Normalized transient photocurrent (TPC) spectra of pristine and doped (target) devices.



**Figure S9.** Extraction of the relative dielectric constant ( $\varepsilon_r$ ) from capacitance-frequency measurements, using the device structure of ITO/PEDOT/perovskite/Al.



**Figure S10.** (a), (b)Hole mobility measurement using the SCLC method, with the hole-only devices structure ITO/PEDOT/perovskite/MoO<sub>3</sub>/Ag. (c), (d) Electron mobility measurement using the SCLC method, with the electron-only devices structure ITO/TPBi/perovskite/TPBi/LiF/Al.



**Figure S11.** Sample surface images:(a), (b) Top-view scanning electron microscope (SEM) images of the pristine and doped (target) films. (c), (d) Atomic force microscope (AFM) images of the pristine and doped (target) films (scale bar: 1 µm).



**Figure S12.** CIE coordinates showing the largest performance gap between pristine and doped (target) devices.

**Table S1.** Time-resolved photoluminescence decay fitting parameters for pristine and doped (target) perovskite films.

	A <sub>1</sub>	<b>T</b> <sub>1</sub>	A <sub>2</sub>	<b>T</b> _2	A <sub>3</sub>	<b>т</b> <sub>3</sub>	T avg
Pristine	1.11293	0.23608	0.03209	2.50201	0.01124	0.0594	0.765
Target	0.56036	0.17565	0.405	0.88862	0.06544	4.21157	2.041

The average exciton lifetime  $(\tau_{avg})$  was calculated using the following equation:

$$\tau_{avg} = \frac{A_1 \tau_1^2 + A_2 \tau_2^2 + A_3 \tau_3^2}{A_1 \tau_1 + A_2 \tau_2 + A_3 \tau_3}$$