Supporting Information

Highly efficient and stable hybrid quantum-dot light-emitting field-effect transistors

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Figure S1. a The output curves of the TFTs with the channel layer of InScO and InScO/ZnO; b The TEM image of the TFT with the channel layer of InScO/ZnO.



Figure S2. a The schematic structure of the bottom-emitting QLED. **b** The schematic structure of the equivalent top-emitting QLED.



Figure S3. The gate leakage current (I_G) recorded simultaneously during the transfer characteristic (I_{DS} VS V_{GS}) measurements of the QD-HLET.



Figure S4. Microscopic images of red QD-HLETs with a constant V_{DS} of 30 V and different V_{GS} of -50 V, -30 V, -10 V, 10 V, and 30 V.



Figure S5. a Current density–luminance–voltage (*J-L-V*) curves of the equivalent topemitting red QLED. **b** Current efficiency and EQE as functions of current density for the equivalent top-emitting red QLED with a structure of Ag(200 nm)/ ITO(10 nm)/ZnO(38 nm)/QDs(25 nm)/TCTA(50 nm)/MoO_x(8 nm)/Au(20 nm). **c** The operating lifetime curves of the equivalent top-emitting red QLED at a constant current density of 71 mA cm⁻², corresponding to an initial luminance of 6300 cd m⁻². **d** Current density–luminance–voltage (*J-L-V*) curves, **e** Current efficiency and EQE as functions of current density, **f** operational lifetime curves (at a constant current density of 71 mA cm⁻², corresponding to an initial luminance of 6450 cd m⁻²) of the equivalent bottom-emitting red QLED with a structure of / ITO(150 nm)/ZnO(38 nm)/QDs(25 nm)/TCTA(50 nm)/MoO_x(8 nm)/Al(120 nm).



Figure S6. a The electronical and optical output characteristics of the red QD-HLET with a hole transport layer of TCTA.



Figure S7. The UPS **a** and transmittance **b** of InScO and ZnO-nanoparticle. The $(\alpha hv)^2$ versus hv curves of InScO **c** and ZnO **d**. The thickness of InScO and ZnO-nanoparticle is 3.4 nm and 38 nm, respectively.

The energy band levels of InScO and ZnO-nanoparticle films can be derived from the ultraviolet photoelectron spectroscopy (UPS) and UV-visible absorption spectroscopy measurements, as shown in Figure S7a, c and d, respectively. The optical bandgap (E_{G}^{OPT}) is calculated using Tauc analysis:¹

$$\alpha h \upsilon = C (h \upsilon - E_G^{OPT})^b \tag{1}$$

where α is the optical absorbance of the material, *hv* is the incident photon energy, C is a constant, and b is an exponent that depends on the nature of the semiconductor bandgap. Because InScO is a direct bandgap semiconductor (See Figure S8), b=1/2 is used. Figure S7c and d show the corresponding $(\alpha hv)^2$ versus hv curves. The extracted

optical bandgaps are 3.91 and 3.47 eV InScO (3.4 nm) and ZnO-nanoparticle (38 nm), respectively. Considering Burstein-Moss shift,^{2,3} which is usually attributed to occupation of the conduction band (inducing optical transitions at energies higher than the minimum-energy fundamental electronic gap), the fundamental electronic bandgap (E_G^F) is smaller than the optical bandgap, and the magnitude of the Burstein-Moss shift (ΔE_G^{BM}), under free-electron theory, is described as

$$\Delta E_G^{BM} = E_G^{OPT} - E_G^F = \frac{\mathbf{h}^2}{2m^*} (3\pi^2 n_e)^{2/3}$$
⁽²⁾

where, m* is the effective mass of the electrons, and n_e is the electron density $(9.2 \times 10^{19} \text{ cm}^{-3} \text{ for InScO} \text{ and } 2.0 \times 10^{14} \text{ cm}^{-3} \text{ for ZnO-nanoparticle})$. m* for InScO is calculated from first-principles calculations to be ~0.2m_e (me is the Electron rest mass), as shown in Figure S8. From Eq. (2), Δ EGBM is calculated to be 0.37 and 0.01 eV (corresponding to EGF of 3.54 and 3.46 eV) for InScO and ZnO-nanoparticle, respectively. The valance band maximum (VBM) is derived from the UPS spectra to be 7.58 and 7.47 eV for InScO and ZnO-nanoparticle, respectively. Then, the conduction band minimum (CBM) is calculated to be 4.04 and 4.01 eV for InScO and ZnO-nanoparticle, respectively.



Figure S8. The calculated energy band structures and effective mass for InScO. The first-principles calculations were performed using the Vienna ab initio simulation package (VASP), where the projected augmented wave (PAW) method was adopted with the exchange-correlation functional in the form of Perdew-Burke-Ernzerhof (PBE). The atomic positions and cell parameters of candidates were fully relaxed until the energy differences were less than 0.001 eV. The cut-off energy of the plane wave was set to be 480 eV. The k meshes are distributed in reciprocal space with an average interval of 0.5Å^{-1} . With the structural recognition, we have excluded the duplicate configurations for the high-throughput screening with the first-principles calculations. We have adopted a supercell of In₂O₃ in bixbyite phase containing 32 In and 48 O atoms. Three Sc atoms have been doped into the supercell to replace In atoms, where 118 unique structures have been found for the first-principles

calculations.



Figure S9. a The carrier distribution when the device is in off-state. b The carrier

distribution when the device is in on-state.



Figure S10. The electronical and optical output characteristics of the red QD-HLET at $V_{GS} = -20$ V and -30 V.



Figure S11. The operational lifetime of red QD-HLET without ZnO nanoparticle layer tested at constant current density of 71 mA cm⁻².



Figure S12. The device structure **a** wide-angle interference **b** and multiple-beam interference **c** of the QD-HLETs.



Figure S13. a The electrical and optical transfer characteristics curves, **b** current efficiency versus current density curves and **c**, EQE versus current density curves of red QD-HLET with 50 nm thick ZnO-nanoparticle (V_{DS} was held at different voltage of 10, 20, and 30 V, and V_{GS} swept from -50 to 30 V at each V_{DS}). **d** The electrical and optical output characteristics curves, **e** current efficiency versus current density curves and **f** corresponding EQE versus current density curves of Devices D (V_{GS} was held at different voltages between -20 to 30 V in step of 10 V, and V_{DS} swept from 0 to 30 V at each V_{GS}); the insets of (**e**, **f**) were the corresponding amplified curves.



Figure S14. a The electrical and optical transfer characteristics curves, **b** current efficiency versus current density curves and **c**, EQE versus current density curves of red QD-HLET with 26 nm thick ZnO-nanoparticle (V_{DS} was held at different voltage of 10, 20, and 30 V, and V_{GS} swept from -50 to 30 V at each V_{DS}). **d** The electrical and optical output characteristics curves, **e** current efficiency versus current density curves and **f** corresponding EQE versus current density curves of Devices E (V_{GS} was held at different voltages between -20 to 30 V in step of 10 V, and V_{DS} swept from 0 to 30 V at each V_{GS}); the insets of (**e**, **f**) were the corresponding amplified curves.

Device types	Mobility	EQE _{max}	L_{\max}	Current	T_{50} at 100	Reference
	$(cm^2 V^{-1} s^{-1})$	(%)	(cd/m^2)	on/off ratio	cd m ⁻² (h)	
Red-QDs	e 3.1	22.8	145000	~10 ⁵	153000	This work
Red-QDs	_e 0.8	8.7	13400	105	N/A	4
Green-QDs	_e 25	11	8000	N/A	N/A	5
Perovskite	_e 20	0.2	18	106	N/A	6
Alq ₃ :DCJTB	N/A	6.5	2190	N/A	N/A	7
NT4N	e 2.3×10 ⁻¹	N/A	1.7	N/A	N/A	8
	_h 4.7×10 ⁻³					
Green-yellow	e 5	0.8	29000	106	N/A	9
TADF	_h 0.11	3.76	1890	104	N/A	10
Super-yellow	e 1.2×10 ⁻¹	1.9	2100	1.7×10 ⁵	N/A	11
	_h 3×10 ⁻³					
F8BT	N/A	8.2	5500	<102	N/A	12
Alq ₃ :DCM	e 5×10 ⁻¹	5.5	N/A	<10 ²	N/A	13
	_h 3×10 ⁻⁵					

Table S1. Comparison of electrical and optical properties of LETs in this work with others reported elsewhere.

N/A, not available; $_{\rm e}$, electron mobility; $_{\rm h}$, hole mobility

Notes and References

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