Supplementary information

Epitaxial Growth of Dual-Color-Emitting Organic Heterostructures via Binary Solvent Synergism Driven Sequential Crystallization

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Materials. Tris-(8-hydroxyquinoline)aluminium (Alq3) and 9,10-bis(phenylethynyl)anthracene (BPEA) were purchased from Alfa Aesar, and used without further treatment. The solvents (A.R.) were purchased from Beijing Chemical Agent, Inc.

Synthesis of organic heterostructures. The heterostructures were prepared by using an anti-solvent diffusion method combined solvent evaporation-induced self-assembly method. In a typical experiment, a stock solution containing both Alq3 (CAlq3 = 2, 4 or 8 mM) and BPEA (CBPEA = 4 mM) in CHCl3 was pre-prepared first (CAlq3:CBPEA = 1:2, 1:1 or 2:1). Subsequently, 20 μl of the mixture was dropped onto the precleaned Si substrate in a beaker containing 0.5 ml anti-solvent. (The Si substrate was previously cleaned by sequential ultrasonic rinses in acetone, ethanol and deionized water.) To avoid direct contact between the mixture and anti-solvent, the Si substrate was placed on top of an inner pillar taller than the level of anti-solvent. The beaker was sealed with cap to prevent the solvent from volatilizing quickly due to its relatively low boiling point. As the solvent and anti-solvent volatilized off thoroughly, organic heterostructures have been successfully prepared. The solvent evaporation time varied from 2 h to 12 h, according to the type of anti-solvent.

Synthesis of Alq3 spindle structures and BPEA microwires. In a typical synthesis, 20 μL of BPEA solution (4 mM) in good solvent of CHCl3, was dropped onto the precleaned Si substrate in a beaker containing 0.5 ml isopropanol. To avoid direct contact between the BPEA solution and anti-solvent, the Si substrate was placed on top of an inner pillar taller than the level of anti-solvent. The beaker was sealed with cap to prevent the solvent from volatilizing quickly. As the solvent and anti-solvent volatilized off thoroughly, BPEA microwires were obtained. Alq3 spindle structures were also synthesized following the similar experiment procedure.

characterization. The morphology and crystal structure of the as-prepared samples were examined by scanning electron microscopy (SEM, Hitachi S-4800) and X-ray diffraction (XRD, Bruker AXS D8 diffractometer), respectively. The fluorescence microscopy images
were taken with a fluorescence microscope by exciting the samples with the UV band (330-380 nm) of a mercury lamp. The PL spectra were measured with fluorescent spectroscopy (HITACHI F-4500). PL lifetime images were taken with an FLIM setup by scanning the samples with a 375 nm picosecond pulse laser. The FLIM setup (PicoQuant) was made up of picosecond pulsed diode laser (PDL 800-D), fiber coupling unit (FCU II), laser scanning microscope (Olympus FV-1000), four channel detector router (PHR 800), and photomultiplier detector assembly (PMA Series). All spectroscopic and optical measurements were carried out at room temperature.
**Figure S1.** Room-temperature PL intensities of individual Alq₃ (black) and BPEA (red) solution at different concentrations (C), recorded upon excitation at 365 nm.

The intensities of Alq₃ PL (black) increases almost linearly as a function of C until $C_{Alq3} = 0.5 \times 10^{-5}$ M, then drops with increasing the value of $C_{Alq3}$ probably due to the concentration quenching effect. This indicates the formation of Alq₃ aggregation with a threshold concentration of $0.5 \times 10^{-5}$ M. The threshold concentration of BPEA aggregation is determined to be $0.6 \times 10^{-5}$ M (red), which is almost the same as that for the formation of Alq₃ aggregation.
Figure S2. PL images of dispersed BPEA and Alq$_3$ nanostructures prepared when ethyl acetate (a), benzene (b) and tetrahydrofuran (c) were used as the anti-solvent, respectively. Scale bar is 250 μm.

When non-alcohol solvents were used as the anti-solvent, BPEA and Alq$_3$ almost simultaneously nucleated and crystallized into dispersed BPEA and Alq$_3$ nanostructures. (Ethyl acetate (a), benzene (b) and tetrahydrofuran (c))
Figure S3. The high-magnification SEM image of single BH. Scale bar is 10 μm.

The high-magnification SEM image indicates each trunk microwire has rectangular cross section.
Figure S4. LCFM images of a typical single BH collected from the (a, d) green-light and (b, e) red-light regions and (c, f) the dual-color superposed LCFM images. (BH prepared when isobutanol was used as anti-solvent at 0°C (a-c) or n-butanol was used as anti-solvent at 25°C (d-f)) The scale bars correspond to 30 μm.

LCFM images of a single BH clearly revealed the spatial distribution of Alq3 and BPEA. In particular, monitored in the red-light region (560-620 nm) showed only a red-emitting segment on the BH body (Figure S3b, e). And monitored in the green-light region (500-560 nm) showed clear emission of bright green light from the microstructure on the microwires surface, whereas the red-emitting body was nearly nonemissive (Figure S3a, d). The superposed LCFM image (Figure S3c, f) also confirms the growth of Alq3 on the BPEA microwires surface.
Figure S5. PL images of BHs prepared when isoamyl alcohol was used as the anti-solvent. Scale bar is 150 µm.
Figure S6. (a, b) XRD patterns of various BHs formed by using different anti-solvents.

XRD patterns of various BHs formed by using different anti-solvents show that the BHs agree well with those of the corresponding α-phase crystals of BPEA and Alq3. Moreover, it is worth noting that the addition of Alq3 did not destroy the original crystal structure of the seed microwires.
Figure S7. PL spectra of BHs formed using different anti-solvents. The excitation wavelength is 365 nm.

To investigate the effect of Alq3 growth on the optical properties of the seed microwires, PL measurements were carried out. The PL spectrum of the various BHs consists of a peak at 560 nm and 601 nm due to the BPEA component and another peak at 512 nm due to the Alq3 component. However, the PL spectrum of the BHs is not a simple mixture or sum of the individual PL spectra of the component because the emission from BPEA is so relative to that of Alq3. The relatively intense BPEA PL (560 nm) indicates efficient direct energy transfer from Alq3 to BPEA the heterojunction interface because of the good overlap of the excitation spectrum of BPEA with the emission spectrum of Alq3.
Figure S8. PL images of BHs prepared when n-Propanol was used as the anti-solvent. Scale bar is 250 μm.

When n-Propanol was used as the anti-solvent, the heterojunctions obtained is similar to that of isopropanol or isobutanol as an anti-solvent, but it has a poorer degree of aggregation than the BHs described above.
Figure S9. PL images of BHs prepared when methanol (a) and ethanol (b) were used as the anti-solvents. Scale bars are 250 μm.

When methanol or ethanol was used as anti-solvents, only the presence of BPEA microwires was observed in the sample because the lower chain length alcohols were detrimental to the crystallization of Alq₃.
Figure S10. (a-c) PL images of BHs prepared at three different molar ratios of Alq$_3$ to BPEA: (a) 1:2, (b) 1:1 and (c) 2:1. All scale bars are 150 μm.

In order to fine-tune the BHs, the number of Alq$_3$ microstructures needs to be controlled by systematically adjusting the molar ratio of Alq$_3$ to BPEA. If $C_{\text{BPEA}}$ was fixed at 4 mM, when the $C_{\text{Alq3}}$ increased from 2 to 4 and to 8 mM, the amount of Alq$_3$ microstructures deposited on the surfaces of the BPEA microwires can be effectively tuned from several to several dozen.
Figure S11. (a, b) PL images of BHs obtained at low temperature (0°C). All scale bars are 150 μm.
Figure S12. Calculated heat of formation for the CHCl3-ROH mixtures.

The first-principles calculations have been performed within the framework of electronic density functional theory (DFT), as implemented in the Vienne Ab initio Simulation Package (VASP). The exchange and correlation interaction was described by the generalized gradient approximation (GGA) with the Perdewe-Wang (PW91) parameterization. The interactions between ions and valence electrons were modeled by the projector-augmented wave (PAW) method. A plain wave cutoff energy for 500 eV has been used. Brillouin zone integrations were modeled by using Gama point. The heats of formation for mixing CHCl3 and C3H7OH (isopropanol) with different ratio can be obtained from the following formula:

$$E_{\text{from}}^{\text{CHCl}_3-\text{C}_3\text{H}_7\text{OH}} = \left[E_{\text{total}}^{\text{CHCl}_3-\text{C}_3\text{H}_7\text{OH}} - (aE_{\text{CHCl}_3} + bE_{\text{C}_3\text{H}_7\text{OH}})\right]/(a + b)$$

Negative heat of formation for mixing CHCl3 and C3H7OH show a strong interaction when mixing alcohols anti-solvent with CHCl3 solvent, indicating that there exist a stable CHCl3-ROH complex in the mixing solvent.
**Figure S13.** Solubility of BPEA and Alq₃ molecules in different ratios of C₃H₇OH-CHCl₃ mixtures.
Figure S14. (a) FLIM images of Alq₃ spindle structures; Scale bar is 30 μm. (b) Corresponding PL decay curves (collection of PL decay time in selected areas), instrumental response functions (scattering from the excitation laser), fitted lifetime curves and residuals of the selected areas.

In the absence of BPEA energy acceptor, the PL lifetime of Alq₃ was 12.06 ns, revealing the intrinsic exciton lifetime of Alq₃ without extrinsic PL quenching.
Figure S15. (a-c) FLIM images of BHs with different Alq3 microstructure numbers when n-Butanol was used as anti-solvent; Scale bars are 30 μm. (d-f) Corresponding PL decay curves (collection of PL decay time in selected areas), instrumental response functions (scattering from the excitation laser), fitted lifetime curves and residuals of the selected areas.

The sharp shortening of the Alq3 lifetime in the heterostructures prove the FRET process from Alq3 to the BPEA acceptor molecules.
Figure S16. (a) FLIM images of BHs when 1-Pentanol was used as anti-solvent; Scale bars are 30 μm. (b) Corresponding PL decay curves (collection of PL decay time in selected areas), instrumental response functions (scattering from the excitation laser), fitted lifetime curves and residuals of the selected areas.

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