



National Taiwan University
Chemical Engineering

2024.02.20

Optoelectronic Polymer Laboratory

Advanced Polymer and Nanotechnology Laboratory

Advisor:

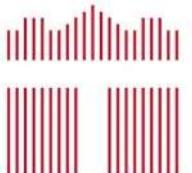
Prof. Wen-Chang Chen

Prof. Chi-Ching Kuo

Presenter :

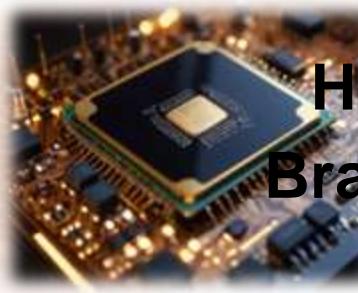
Wei-Cheng Chen

(W.-C. Chen)

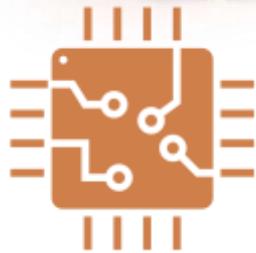


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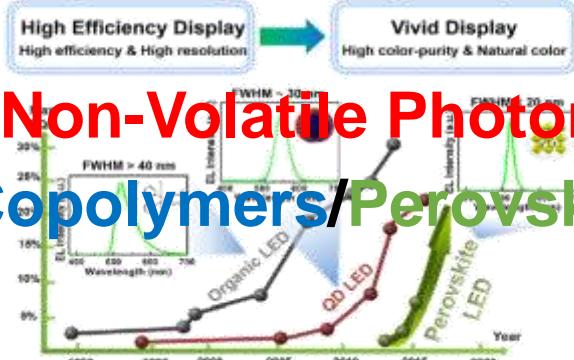
Introduction & Motivation



High Performance Non-Volatile Photomemory Utilizing Branched Triblock Copolymers/Peroovskite Quantum Dots



- ✗ Data latency
 - ✗ Intralayer consumption
 - ✗ Only electric operation



- ✓ High potential
 - ✓ High color purity
 - ✓ Solution process

**emory Utilizing
te Quantum Dots**

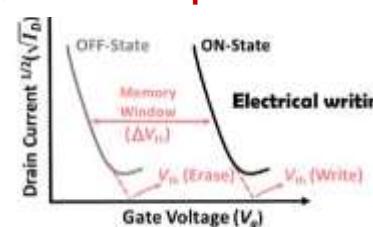
→ Foldable → Stretchable

Photomemory

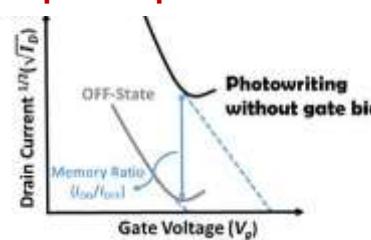


- ✓ Low latency
 - ✓ Low energy consumption
 - ✓ Arbitrary wavelength

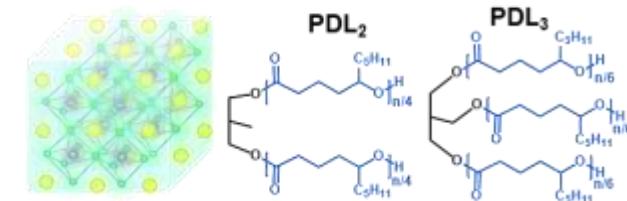
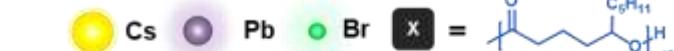
Electric operation



Optical operation



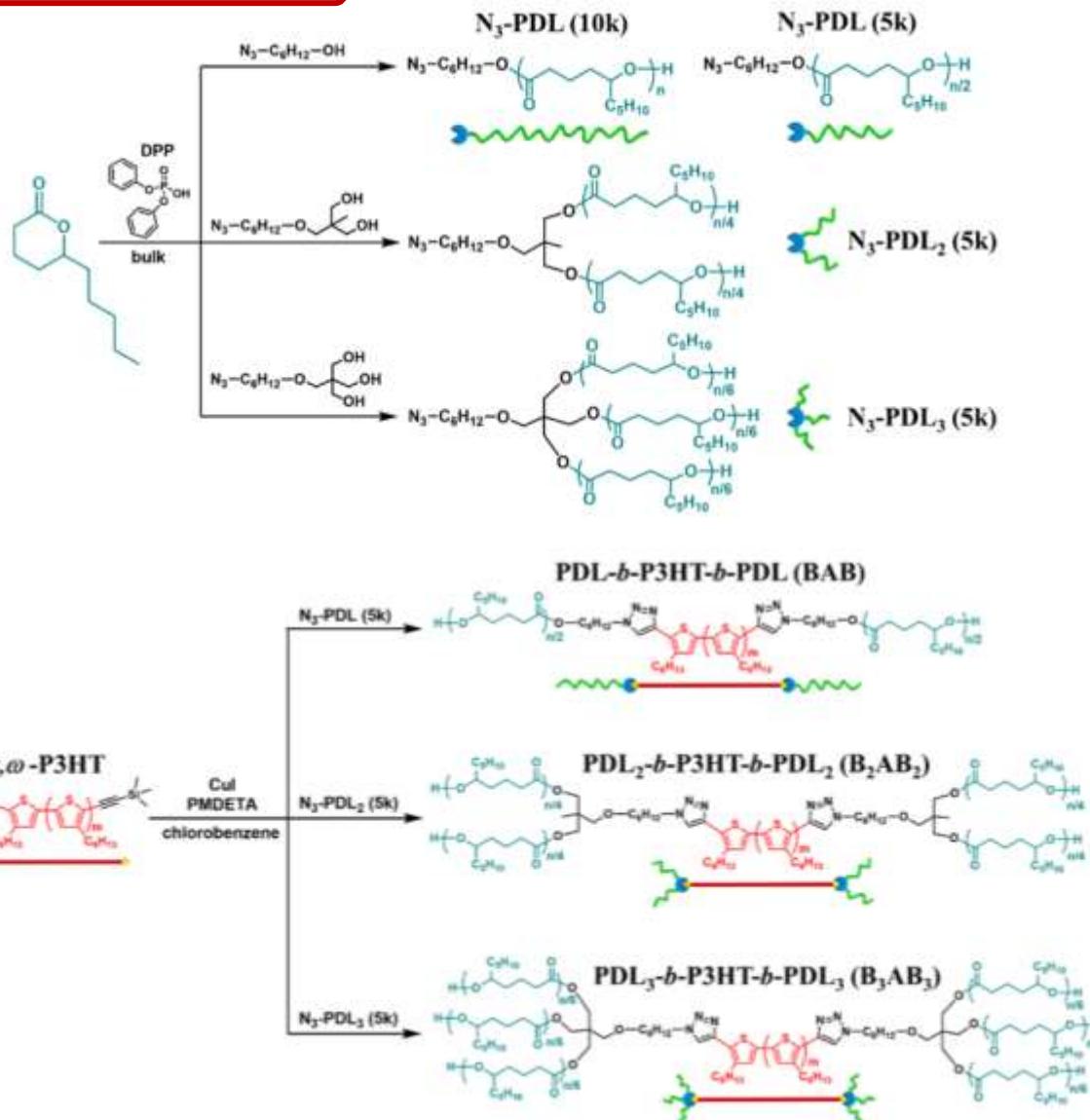
- ✗ Self-aggregation
 - ✗ Ion dissociation
 - ✗ Solution stability



Our Research (2/9)

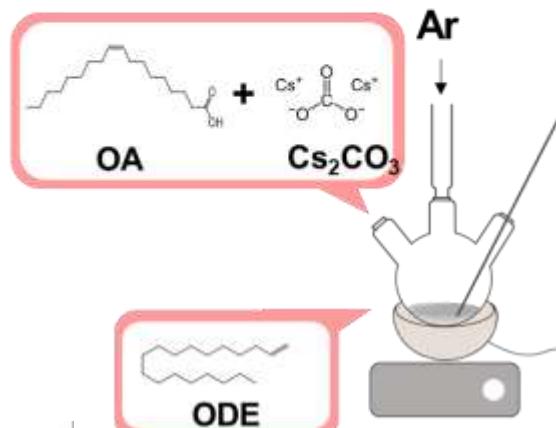
Synthesis of Copolymers and Perovskite

Synthesis

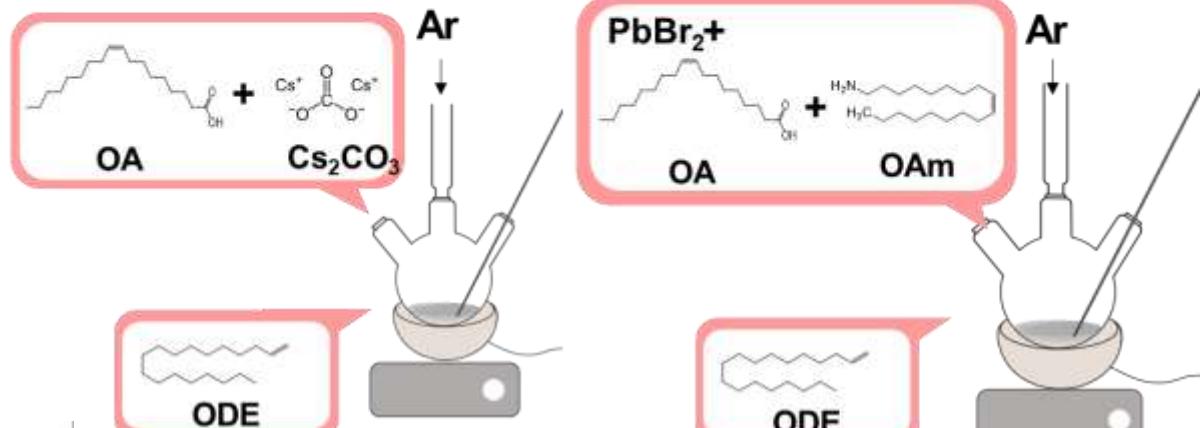


Hot Injection Method

Preparation of Cs-oleate.



Preparation of Pb precursor.



ODE
Magnetic Stirrer



Hot injection 180°C 5s



Purification with EA

Toluene: Ethyl acetate (EA) = 1:2

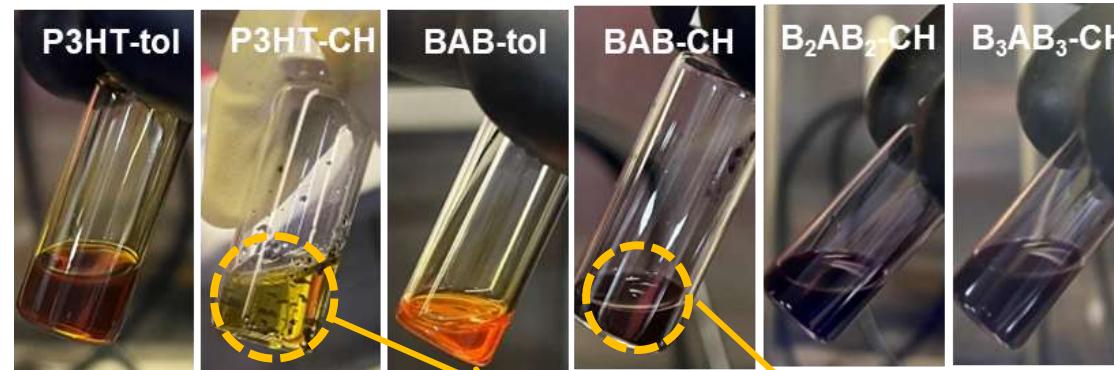


Cooling to room temp.



Self-assembly of Copolymers

Solvent selectivity



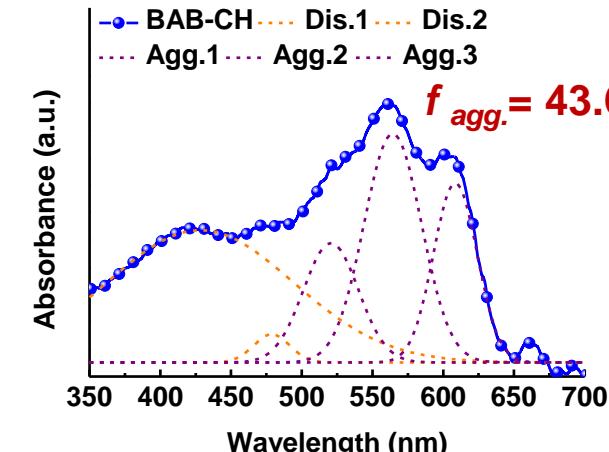
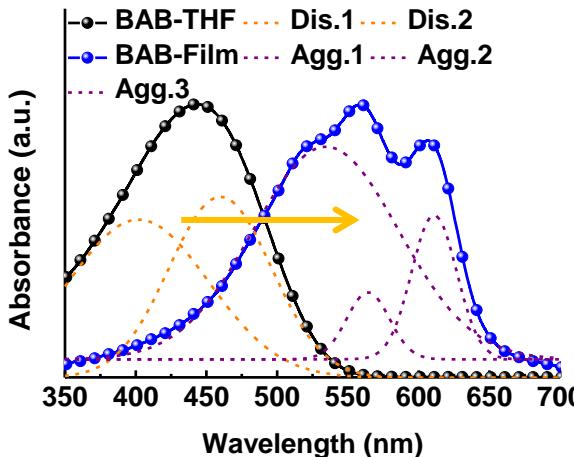
Tol = Toluene
CH = Cyclohexane

Insoluble at room temperature

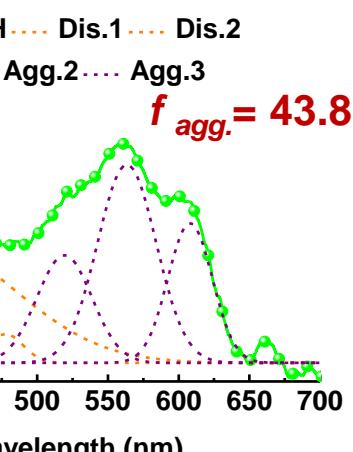
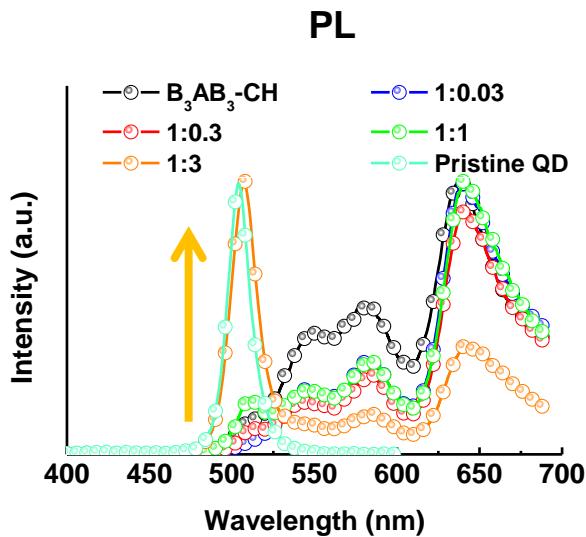
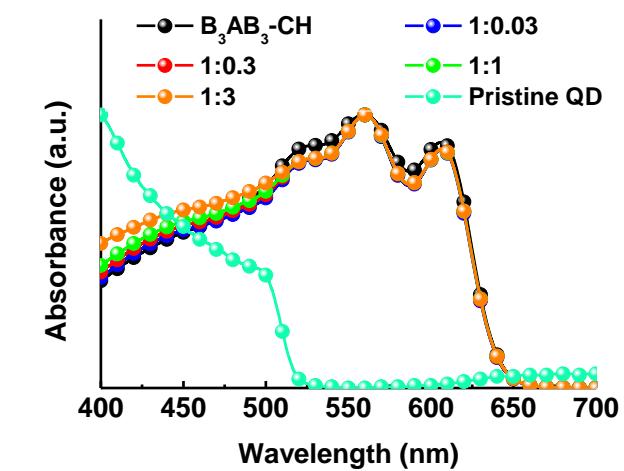
Phase separation

$$f_{\text{aggregation}} = \frac{\frac{A_{\text{aggregate}}}{F}}{A_{\text{disorser}} + \frac{A_{\text{aggregate}}}{F}}$$

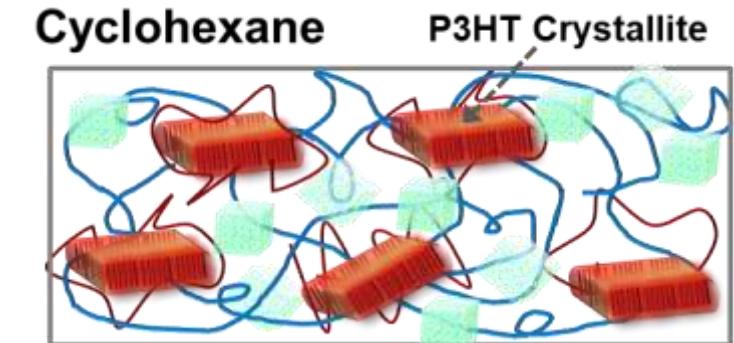
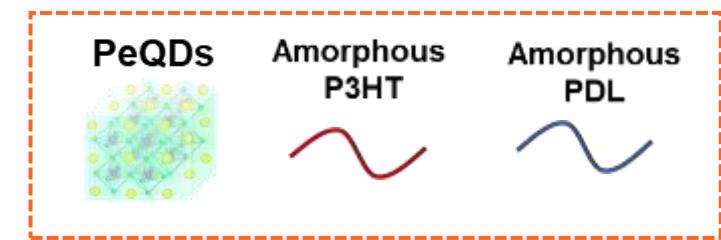
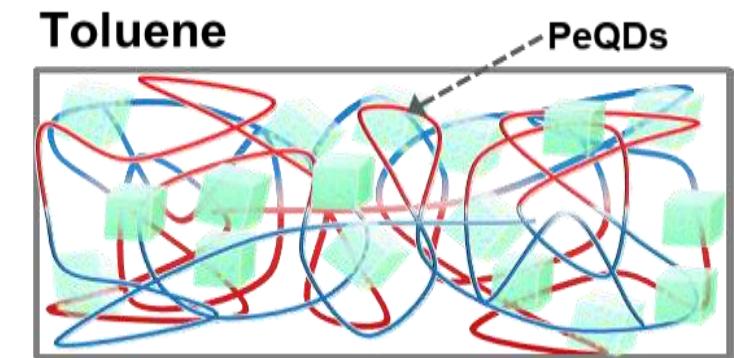
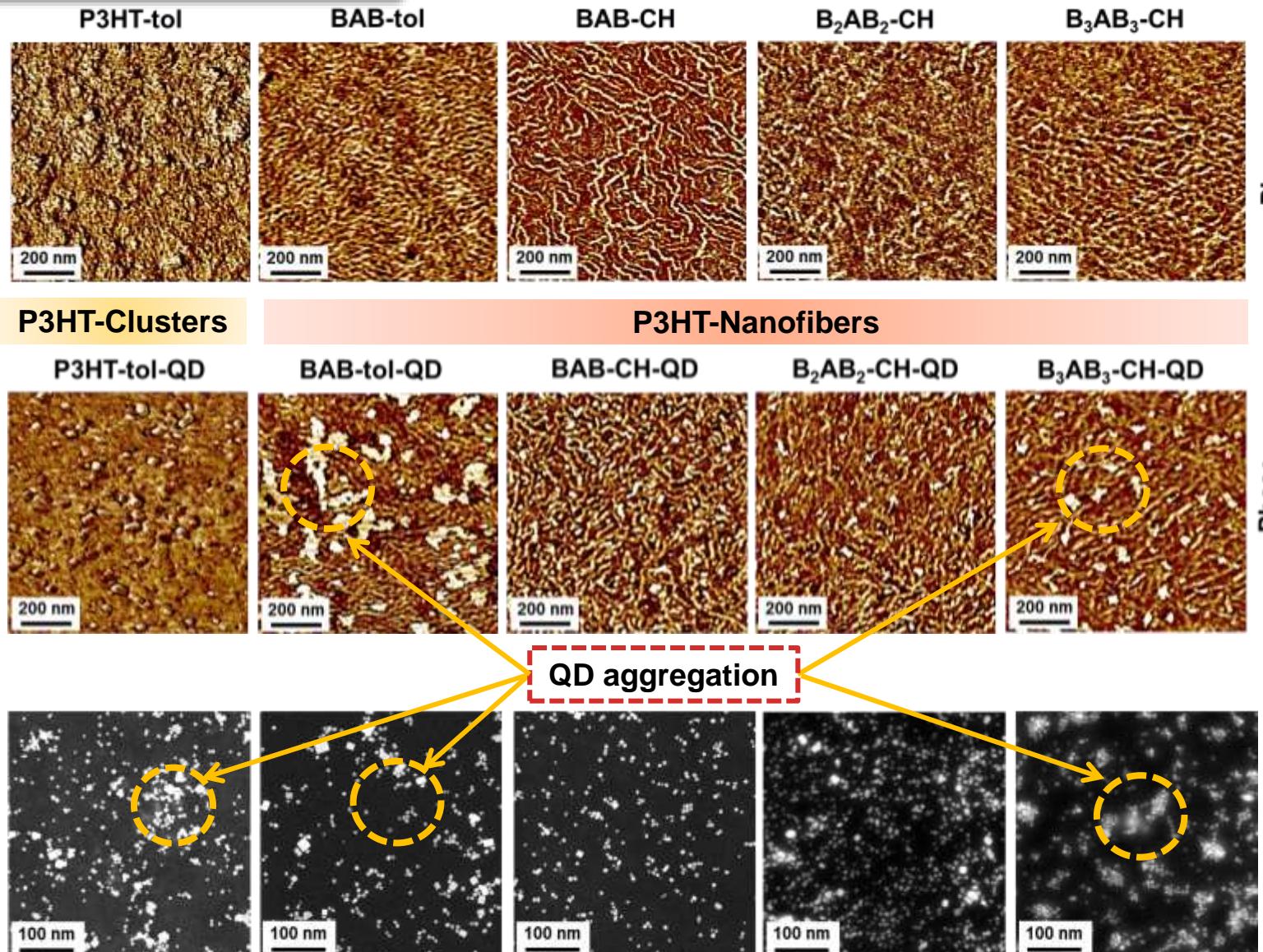
Fraction of aggregation



Optimizing QD ratio



Morphology and QD distribution

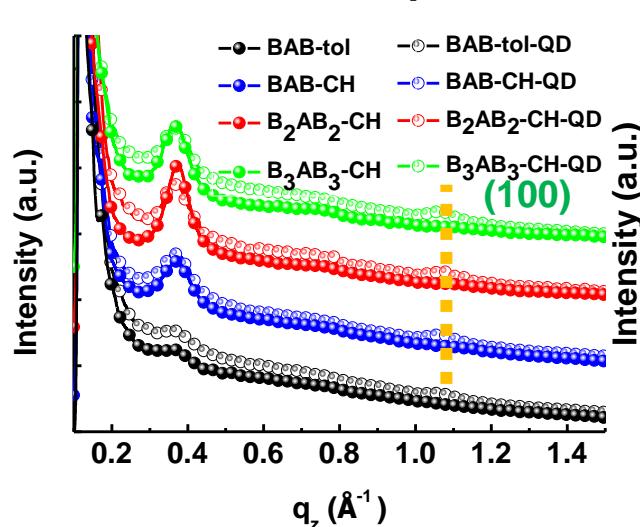


Dispersed region ↑ QD Aggregation ↑

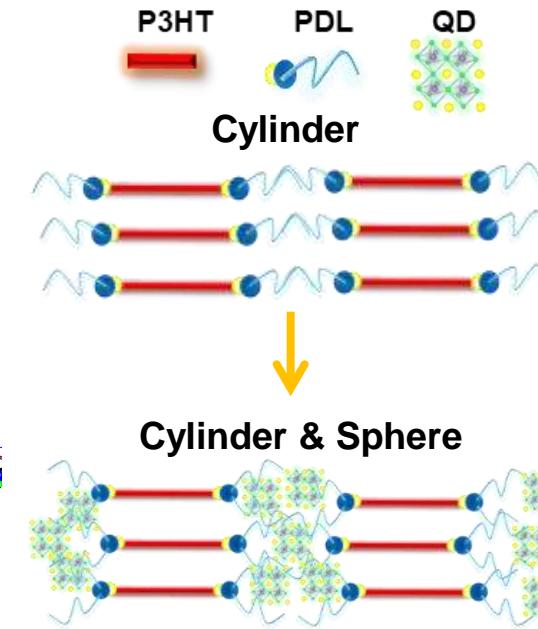
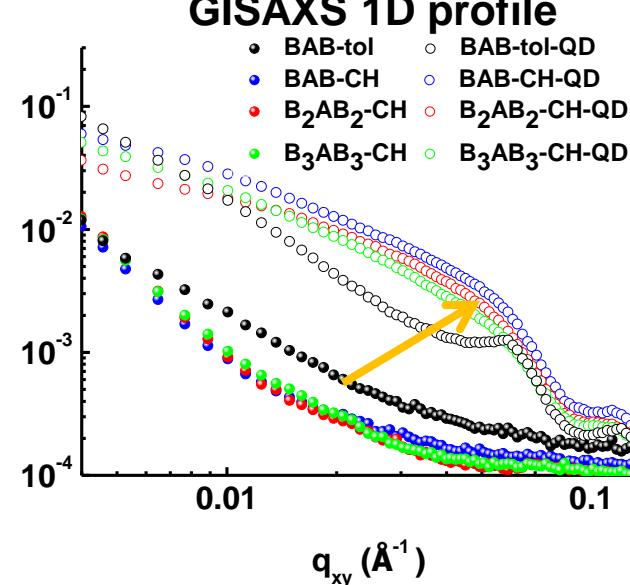
Our Research (5/9)

GIWAXS and SAXS for Crystallites

GIWAXS 1D profile



GISAXS 1D profile



Parameter

D = Domain size

Xc = peak center of GIWAXS

K = 0.9 λ = 0.102164 nm

R_g = Radius of Gyration

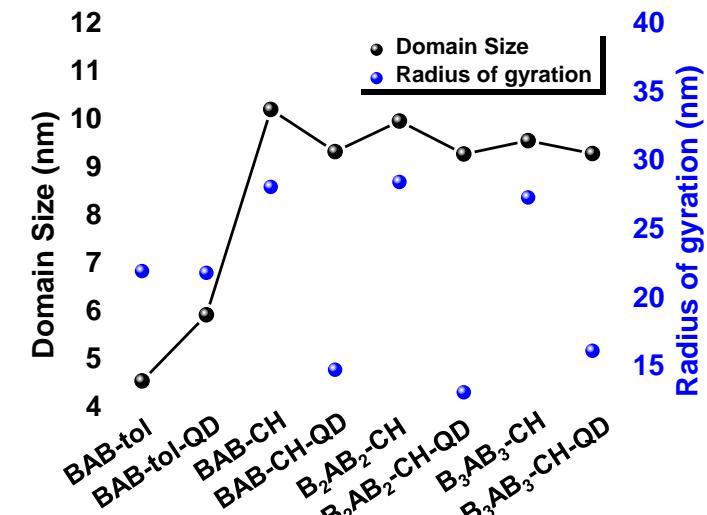
$$\text{Domain size} = \frac{K\lambda}{FWHM \times \cos \theta}$$

$$q = \frac{2\pi}{d} = \frac{4\pi \sin \theta}{\lambda}$$

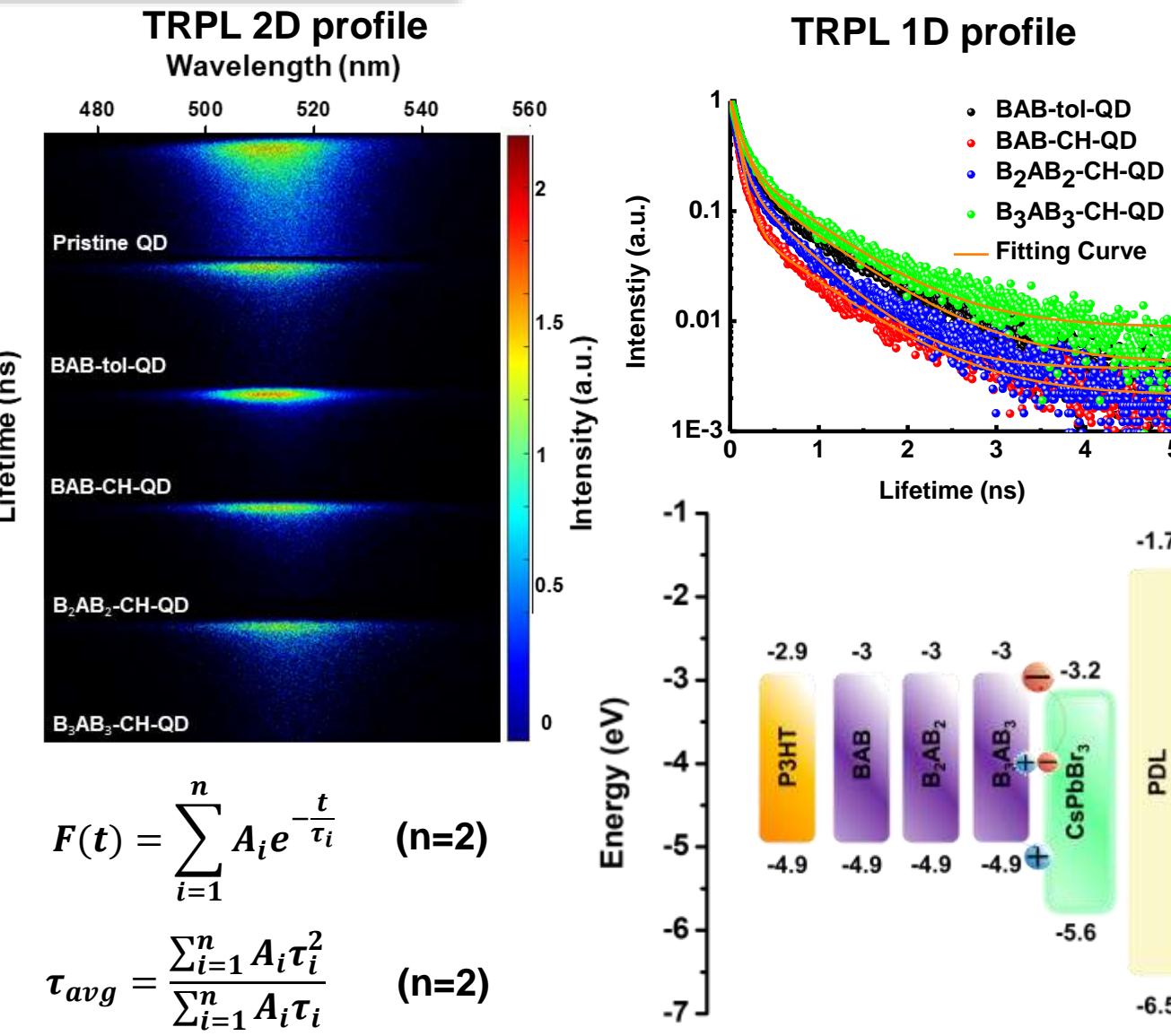
| | D (nm) | Xc (q, Å⁻¹) | d-spacing (Å) | Rg |
|---------------------------------------|--------|-------------|---------------|-------|
| BAB-CH | 10.28 | 0.368 | 17.1 | 28.15 |
| BAB-CH-QD | 9.42 | 0.366 | 17.2 | 14.82 |
| B ₂ AB ₂ -CH | 9.99 | 0.369 | 17.0 | 28.52 |
| B ₂ AB ₂ -CH-QD | 9.30 | 0.367 | 17.1 | 13.17 |
| B ₃ AB ₃ -CH | 9.62 | 0.372 | 16.9 | 27.39 |
| B ₃ AB ₃ -CH-QD | 9.31 | 0.366 | 17.2 | 16.20 |
| BAB-tol | 4.57 | 0.356 | 17.7 | 21.89 |
| BAB-tol-QD | 5.95 | 0.363 | 17.3 | 22.01 |

Guinier Plot:

$$\ln(I(q)) = \ln(I_0) - \frac{R_q^2}{3} \times q^2$$



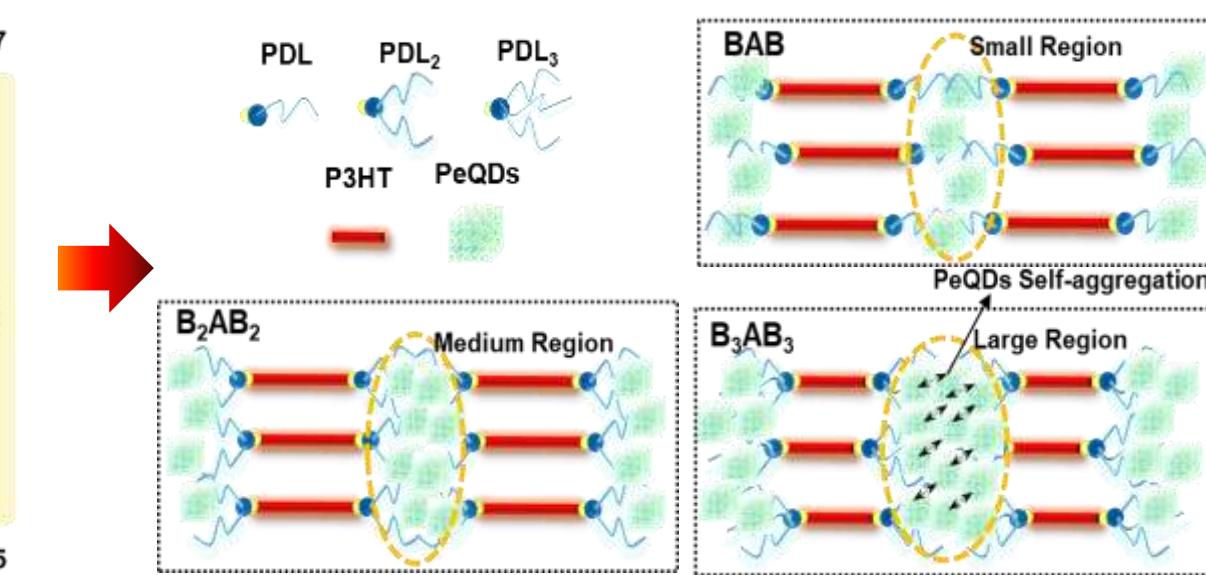
Time-resolved PL and Charge Transfer Efficiency



Non-radiative recombination

$$CTE = \frac{\tau_{QD} - \tau_{copolymers\&QD}}{\tau_{QD}} \times 100\%$$

| Range: 475~540 nm | A_1 | τ_1 | A_2 | τ_2 | τ_{avg} (ns) | CTE (%) |
|----------------------|------------------------------------------------------------------------------------------|----------|-------|----------|------------------------------------------------------------------------------------------|-------------------------------------------------------------------------------------------|
| Pristine QD | 0.55 | 0.25 | 0.45 | 1.57 | 1.36 | |
| BAB-tol-QD | 0.81 | 0.12 | 0.19 | 0.75 | 0.50 | 63.58 |
| BAB-CH-QD | 0.93  | 0.08 | 0.07 | 0.73 | 0.33  | 75.64  |
| B_2AB_2 -CH-QD | 0.83 | 0.08 | 0.17 | 0.55 | 0.35 | 74.32 |
| B_3AB_3 -CH-QD | 0.76 | 0.11 | 0.24 | 0.71 | 0.52 | 61.96 |

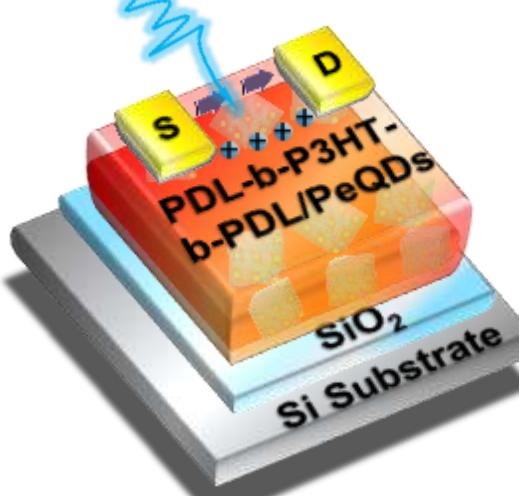


Self-aggregation of PeQDs decreases CTE.

Our Research (7/9)

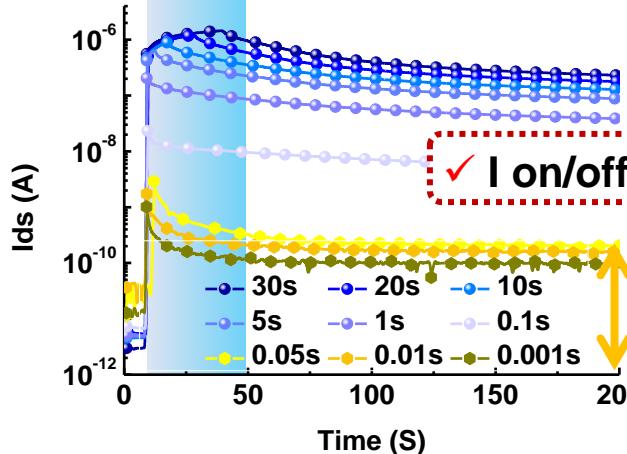
Memory Device Performance

Light stimulus

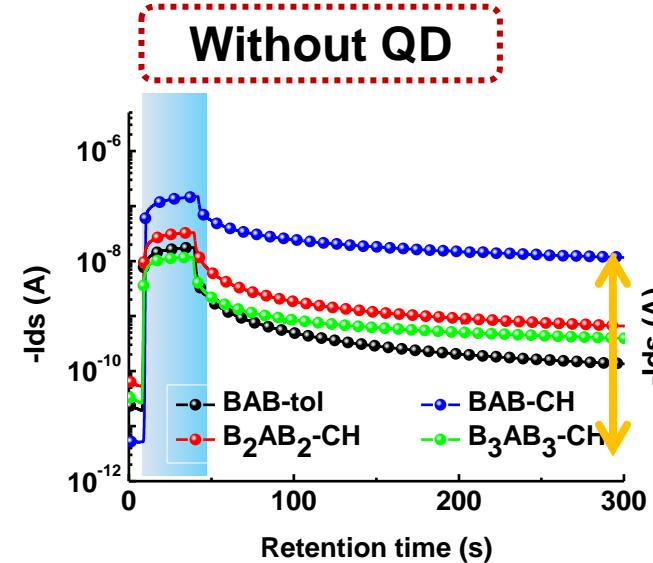


Transient Photocurrent

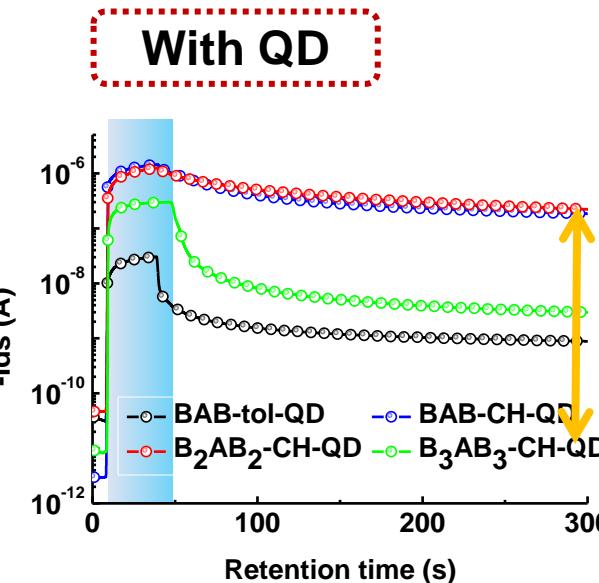
BAB-CH-QD



Without QD

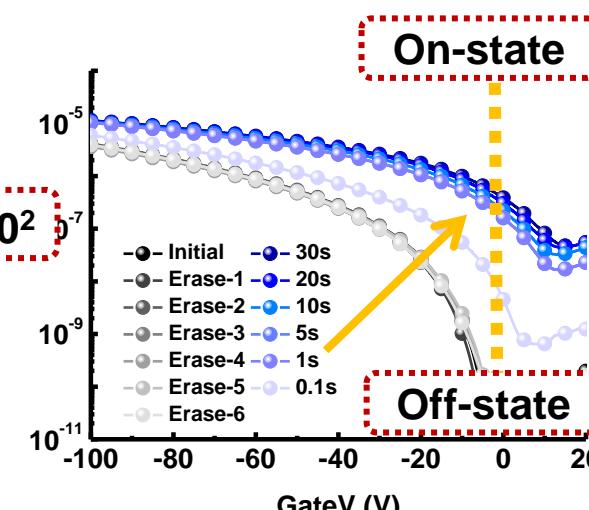


With QD

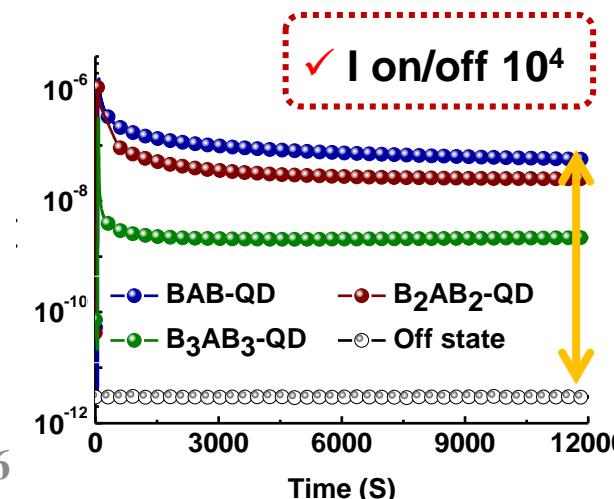


✓ The larger domain size of P3HT crystallites reveals better electron trapping.

Transfer Curve



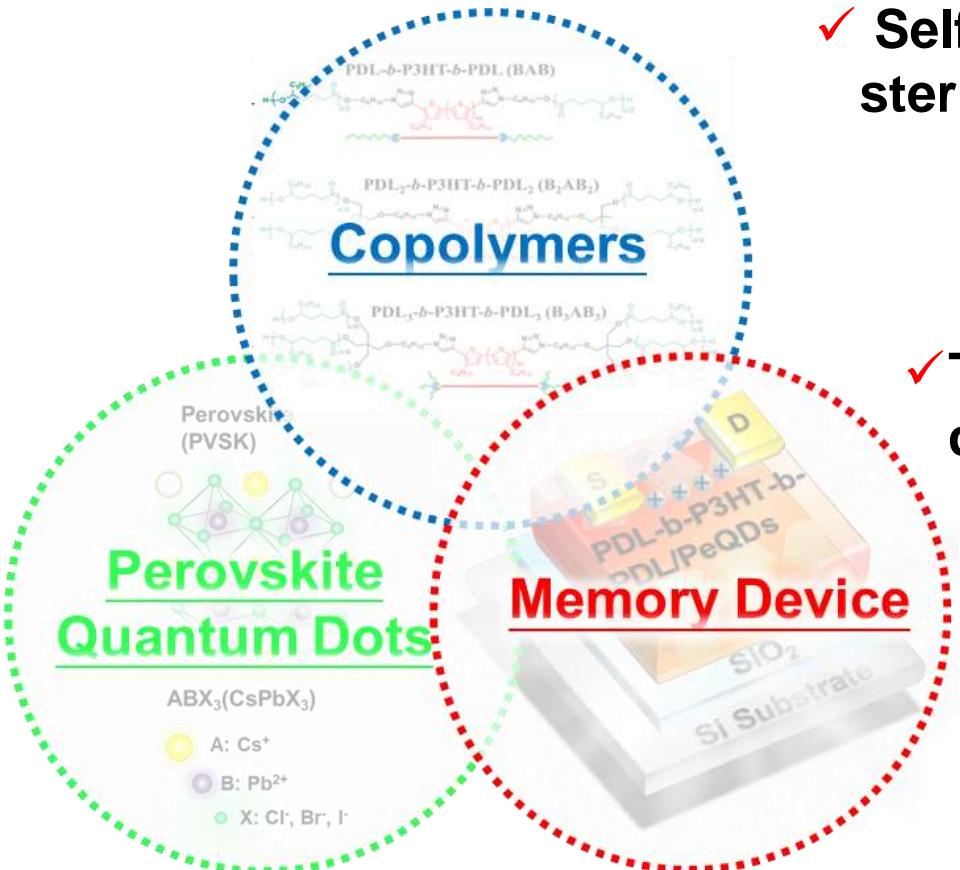
Retention 10⁴ s



✓ The better distribution of QD in copolymers presents better photo-sensitivity.

✓ The closer distance between QD and copolymers crystallites presents better electron trapping.

Conclusion



- ✓ Self-aggregation of PeQDs is improved by an appropriate steric hindrance and self-assembly of triblock copolymers.
- ✓ The good distribution of PeQDs and larger P3HT crystallite domain size present higher photosensitivity.
- ✓ A superior charge transfer efficiency (75.6%) is demonstrated by optimizing the interfaces of QD and P3HT crystallites.

Our Research (9/9)

Acknowledgments

Optoelectronic Polymer Laboratory



National Taiwan University



Advanced Polymer and Nanotechnology Laboratory



National Taipei
University of Technology



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