

# Graphene-based Polymer Bulk Heterojunction Solar Cells

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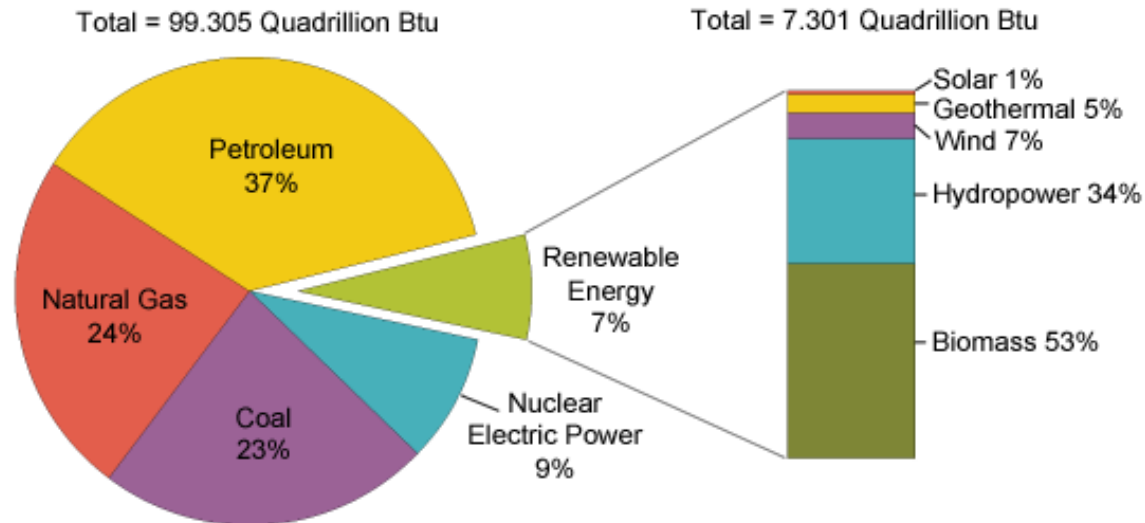
University of Cincinnati

APS March Meeting 2012, Boston

# Solar Energy In Development

- Renewable
- Potential for High coverage
- Low emission

The Role of Renewable Energy in the Nation's Energy Supply, 2008



Note: Sum of components may not equal 100% due to independent rounding.  
Source: Energy Information Administration, *Renewable Energy Consumption and Electricity Preliminary Statistics 2008*,  
Table 1: U.S. Energy Consumption by Energy Source, 2004-2008 (July 2009).

# A comparison

## Inorganic solar cells

- From 1941
- High processing cost
- Thickness in microns
- Not flexible
- 25.0% for Si cells\*

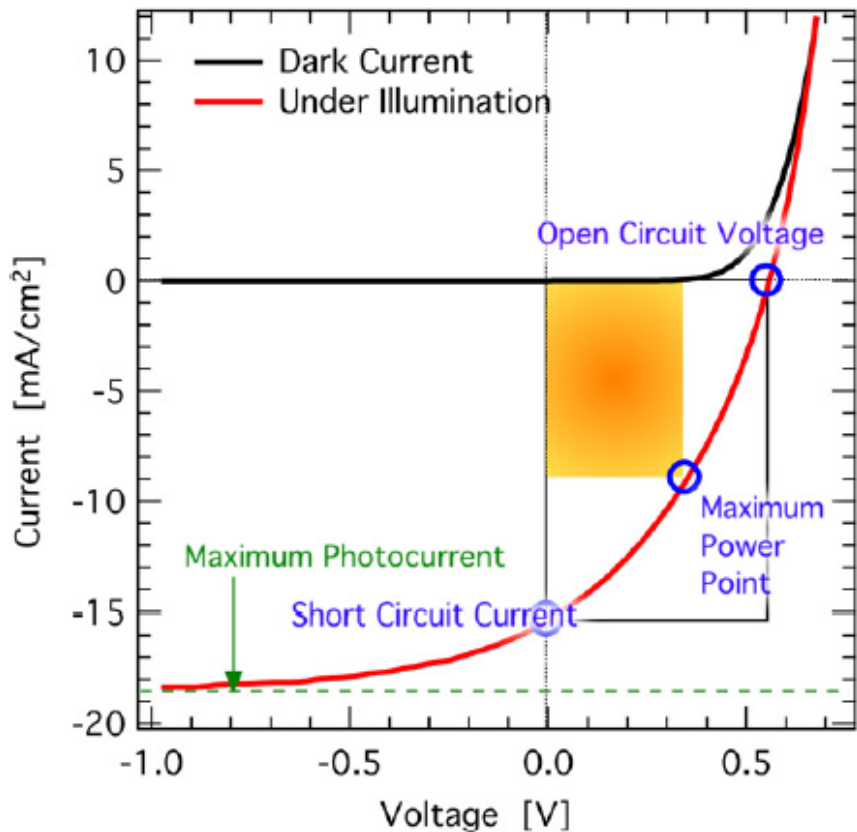
## Organic solar cells

- From 1954
- Solution processible
- 100~300 nm thick
- Flexible
- 6.1% for polymer BHJ cells\*\*

\* Green, Progress in Photovoltaics, 2009. **17**(3): p. 183-189.

\*\* Park *et al.*, Nat. Photonics, 2009. **3**(5): p. 297-U5.

# Solar Cell Parameters

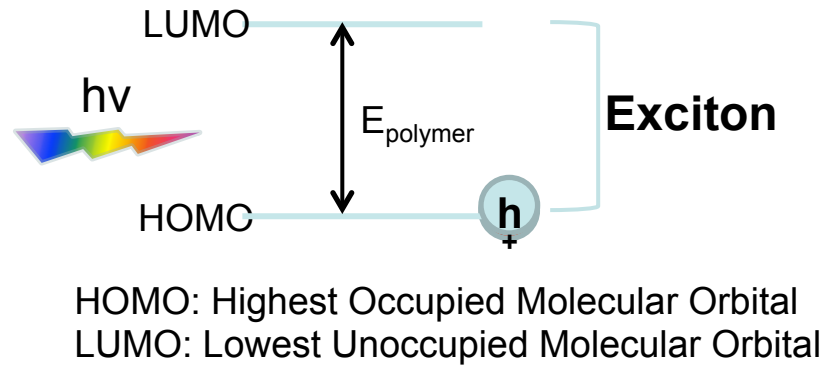
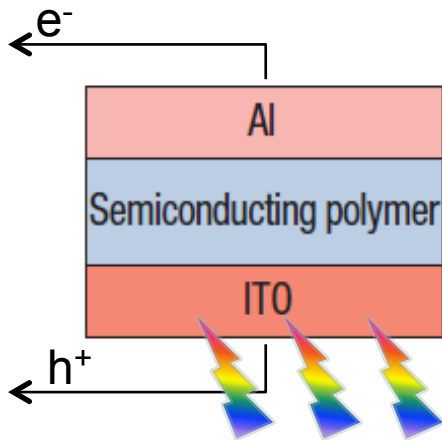


- $J_{sc}$ : Short-circuit current density
- $V_{oc}$ : Open-circuit voltage
- $P_{max}$ : Maximum output power
- **FF**: Fill factor
- **Power conversion efficiency (POC)**

$$\eta = P_{max} / P_{in}$$

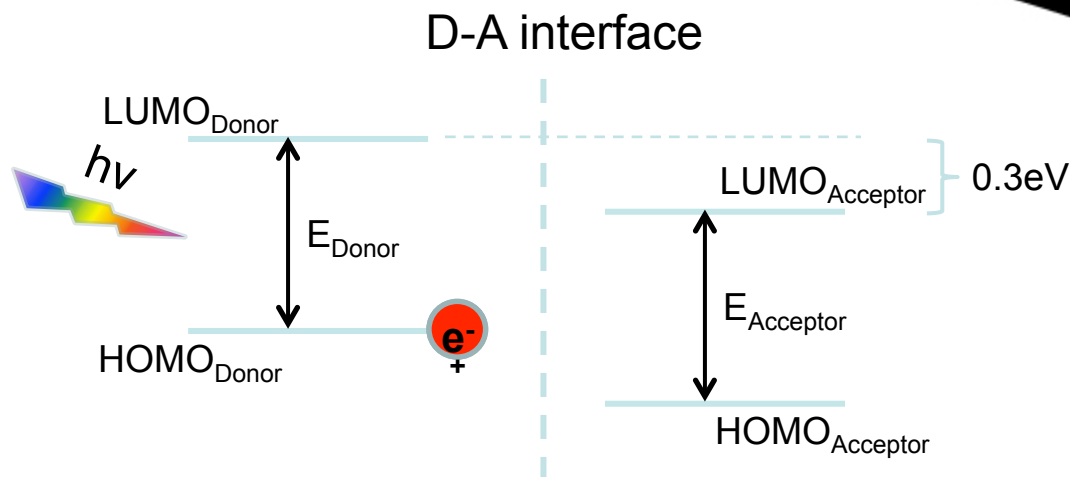
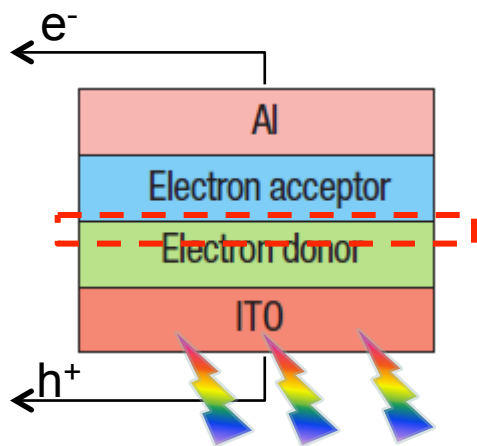
# Polymer solar cells

## Single-layer device



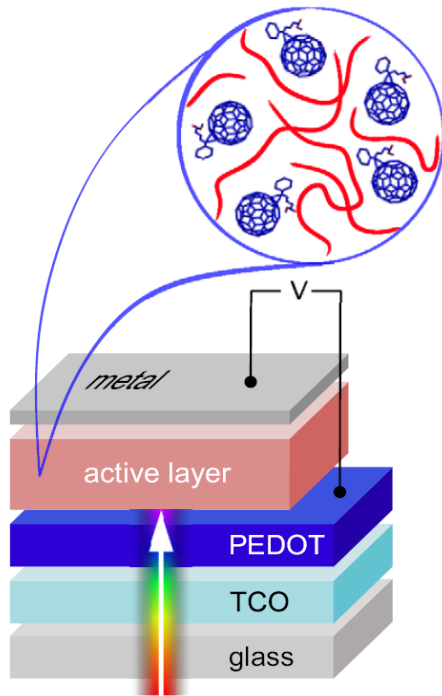
- $\sim 0.3$  eV energy is needed to dissociate excitons
- An external voltage is required
- Recombination of free charge carriers

# Bilayer Device



- D-A interface facilitates exciton dissociation
- Electron transfer from donor (semiconducting polymer) to acceptor
- Exciton dissociation is energetically favorable
- Exciton diffusion length (~10 nm)
- D-A interfacial area is limited by device geometry

# Bulk Heterojunction(BHJ) Solar Cells

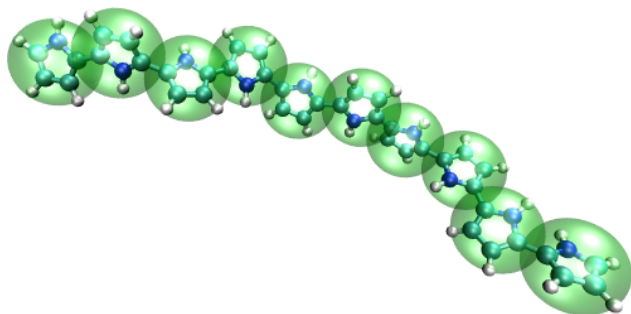


- Nanoscale penetrating network
- D-A interface close to where exciton is generated
- Much increased D-A interfacial area
- Over 6% PCE for P3HT:PCBM BHJs\*

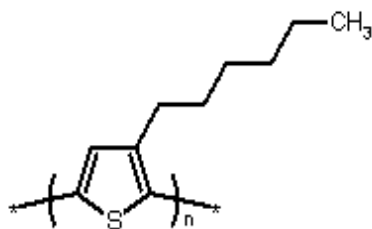
(Picture source: Deibel and Dyakonov, Reports on Progress in Physics, 2010. 73(9): p. 1-39)

\*Peet et al., Nature Materials, 2007. 6(7) : p. 497-500.

# Donor

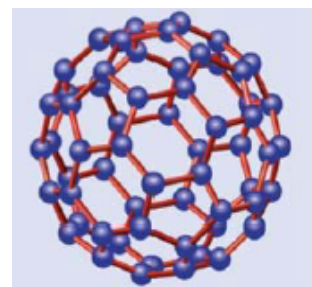


Conjugated polymer

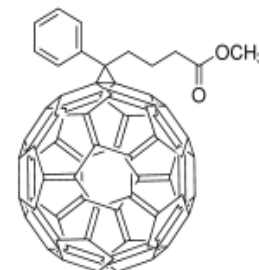


P3HT

# Acceptor



Fullerene(C<sub>60</sub>)



PCBM



# Factors Affecting BHJ Performance

- **Choice of donor and acceptor materials:** band gap and miscibility
- **Choice of solvent:** polymer chain packing
- **Donor-acceptor ratio:** domain size
- **Annealing conditions:** reorganize polymer chains, crystallization
- **Other post-production treatments:** DC voltage during annealing for ordered structure \*

Morphology

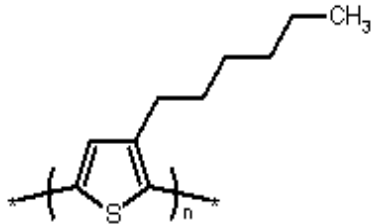


Performance

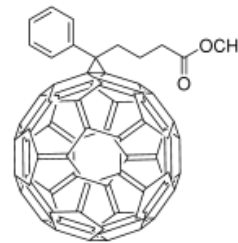
# BHJ features

Polymer:Fullerene BHJ device

- High interfacial area for exciton dissociation
- Bicontinuous network for charge transport
- 50:50 w/w P3HT:PCBM for optimum performance
- Increase P3HT ratio to capture more solar energy



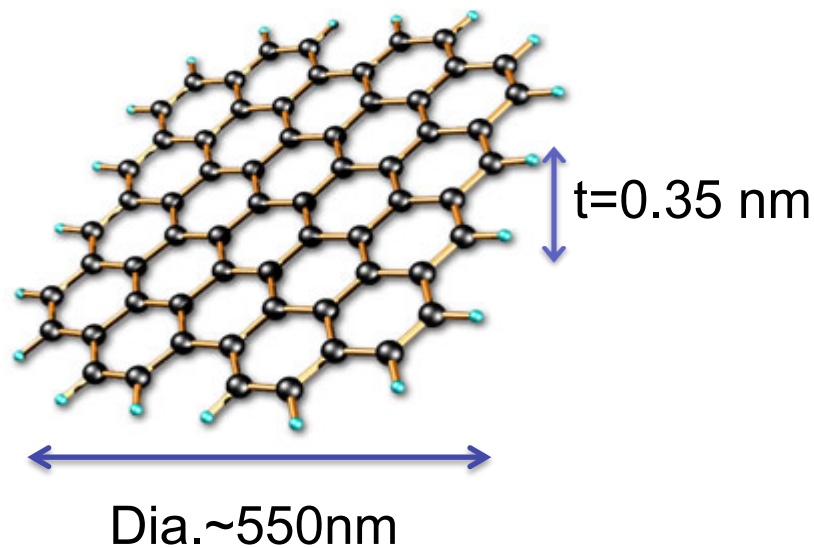
P3HT



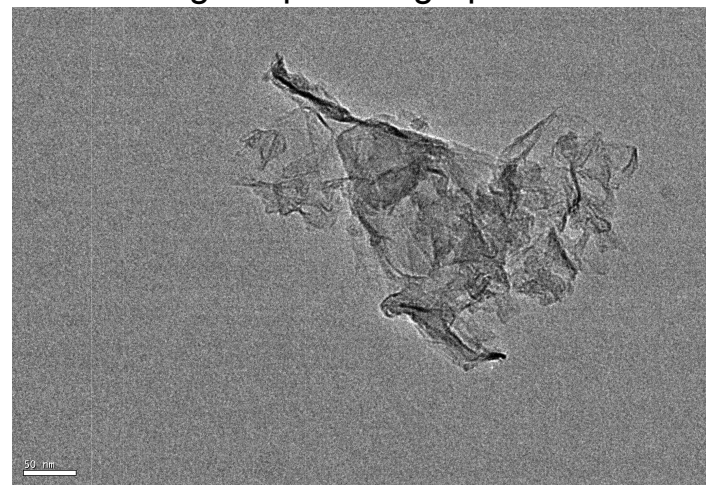
PCBM

# Pristine Graphene

- OPVs with chemically modified graphenes were reported\*
- Excellent conductivity and high aspect ratio
- Percolation paths at very low fraction

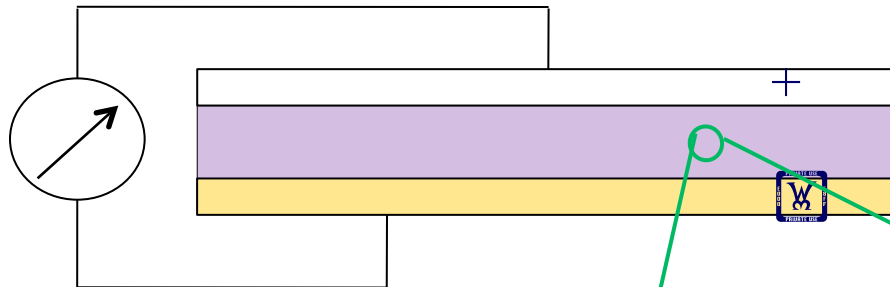





TEM image of pristine graphene flake

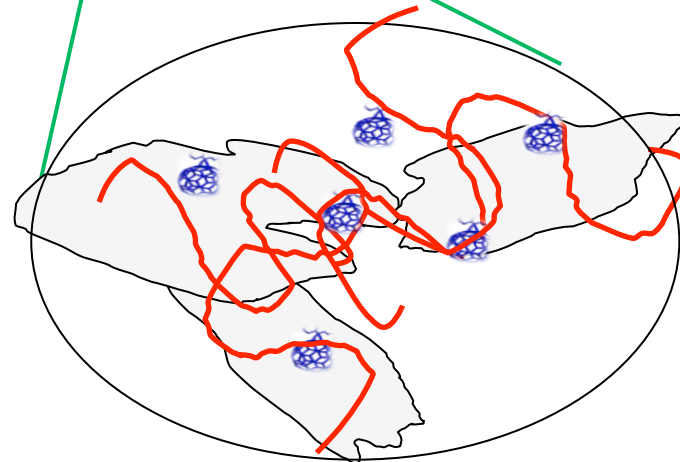


Scale bar=50nm

# The Active layer

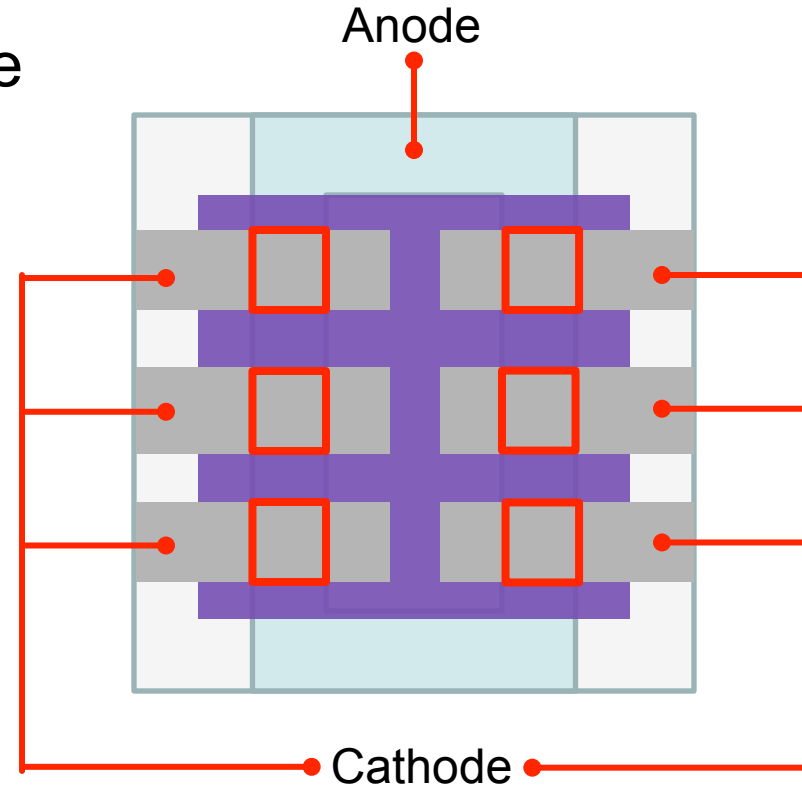


-  P3HT (~90.99%)
-  PCBM (~9%)
-  Graphene (~0.01%)



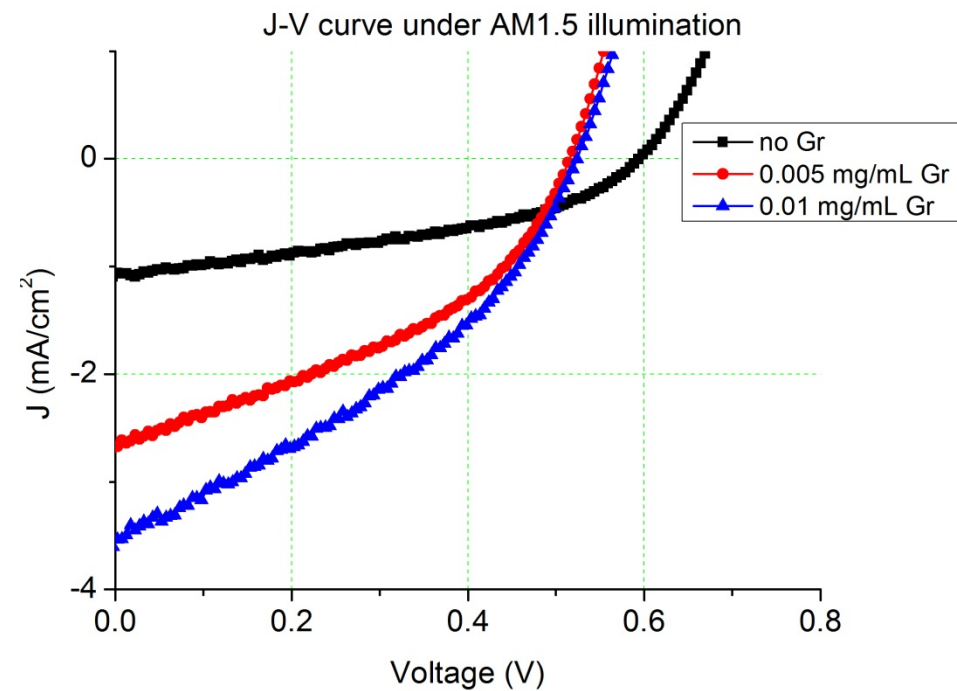
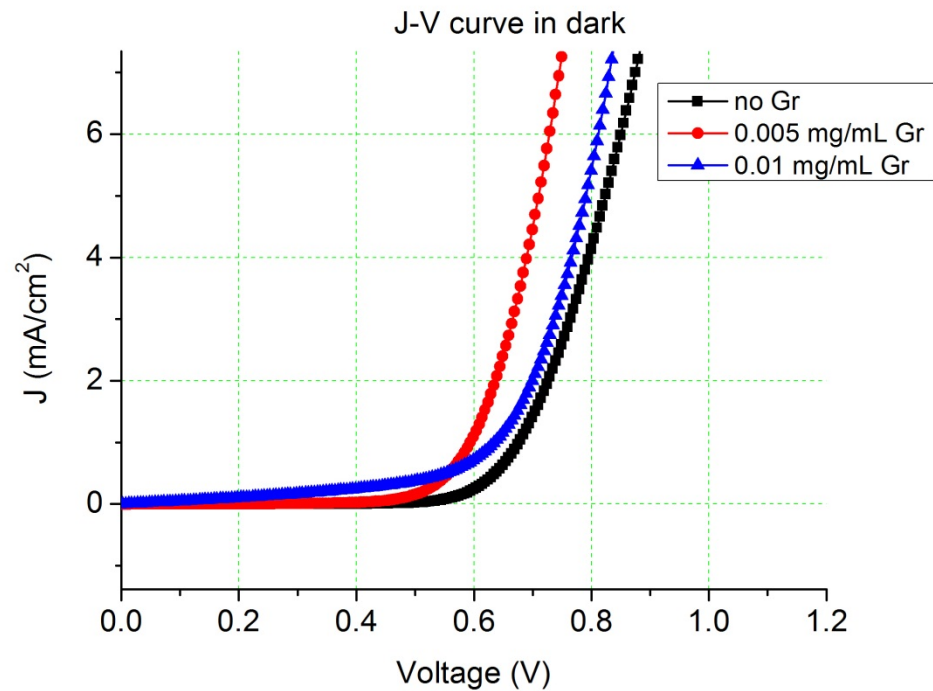
# Device Fabrication

- Patterned ITO as bottom electrode
- PEDOT:PSS by spin coating
- 10:1 P3HT:PCBM(w/w) with graphene by spin coating
- LiF and Aluminum
- Fabricated and annealed in  $N_2$

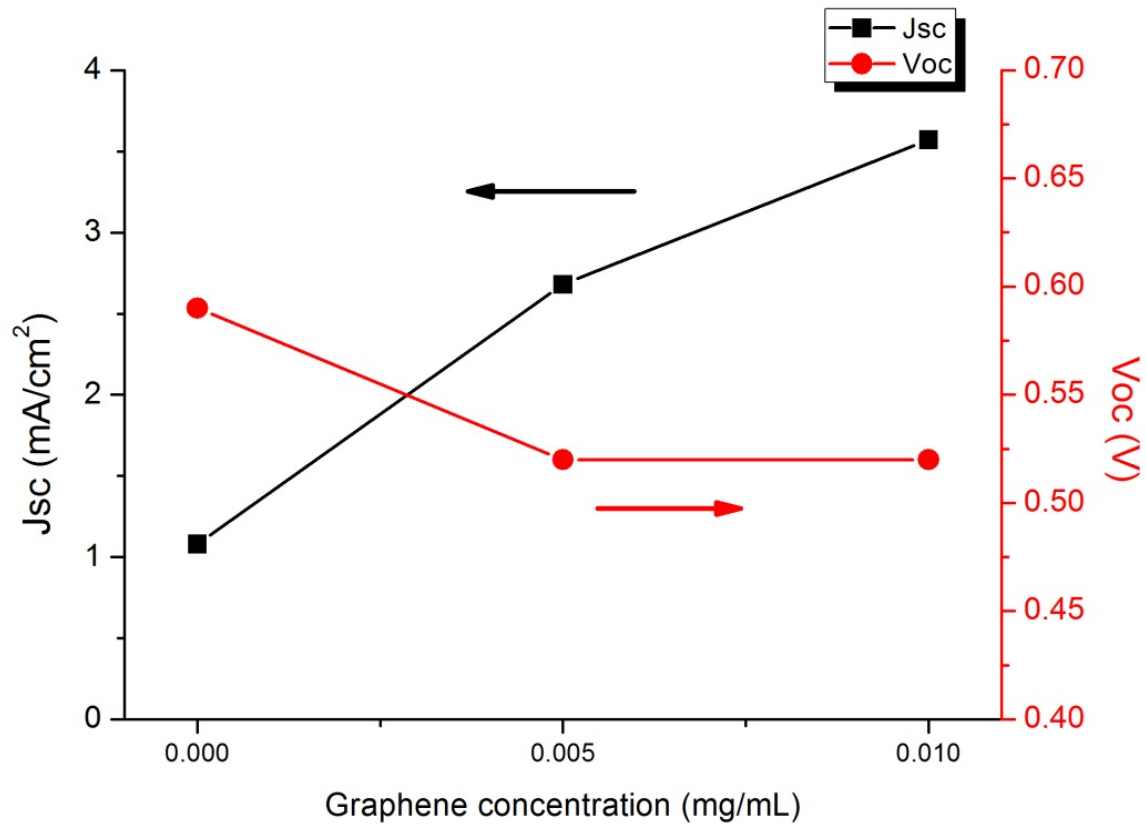


# Device Characterization

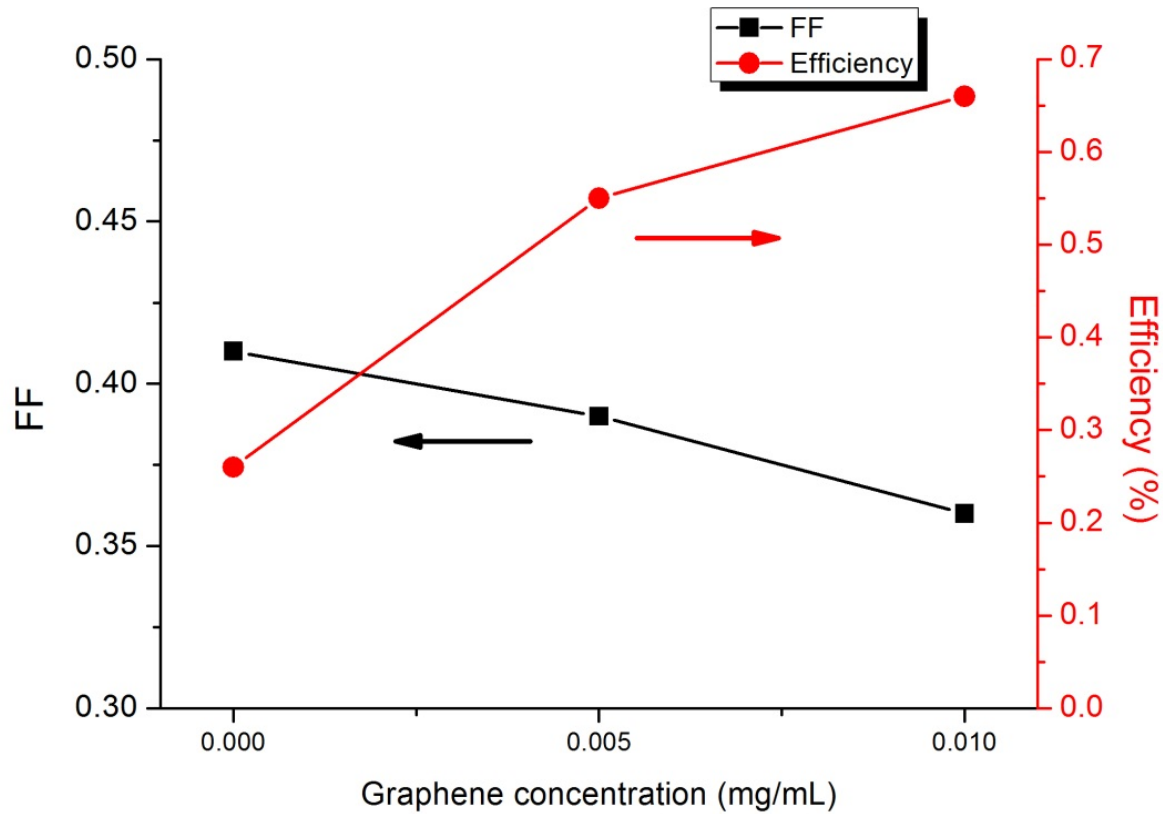
## ➤ J-V characteristics



## ➤ Cell performance summary



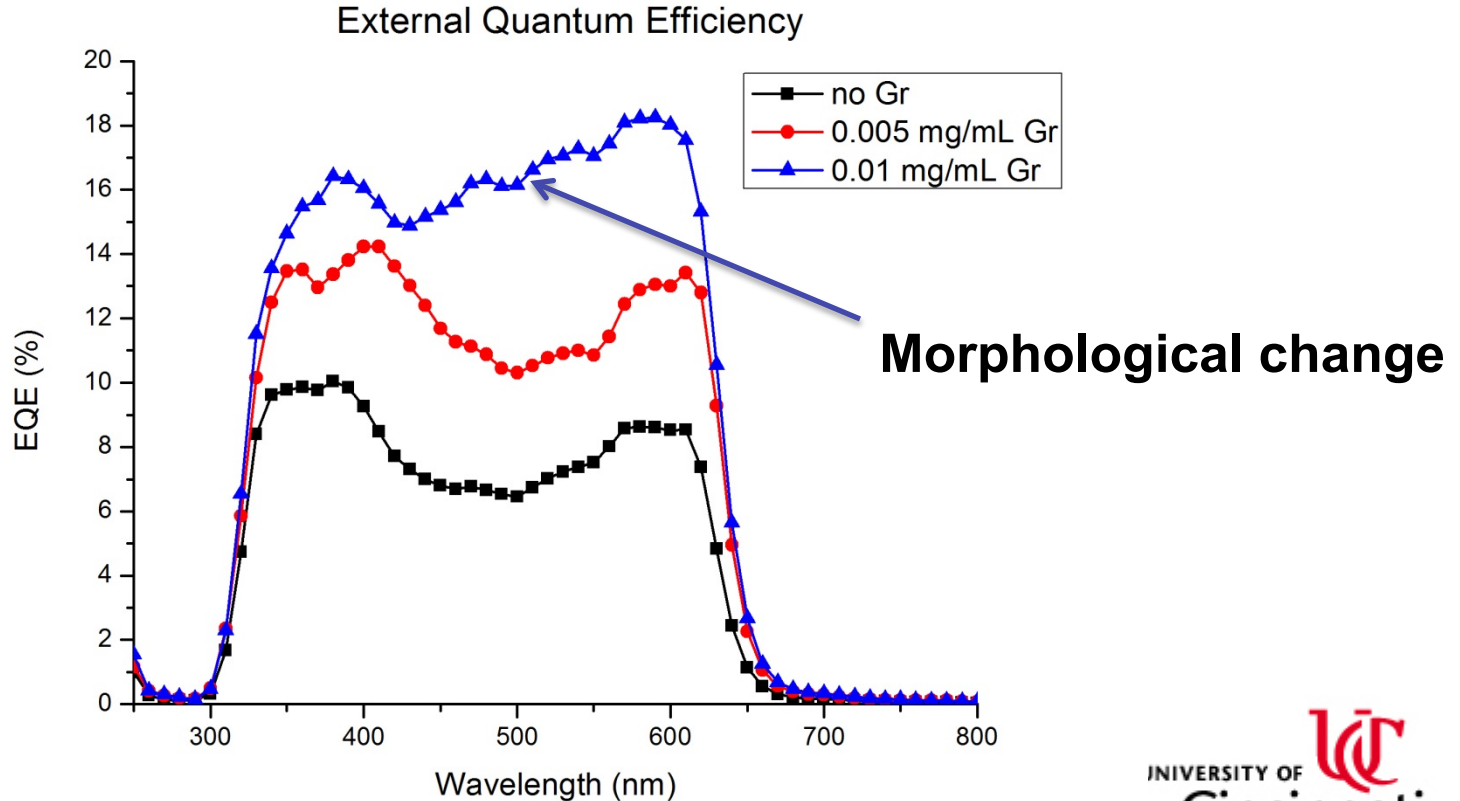
## ➤ Cell performance summary(cont.)





# Device Characterization(cont.)

## ➤ External Quantum Efficiency(EQE)\*



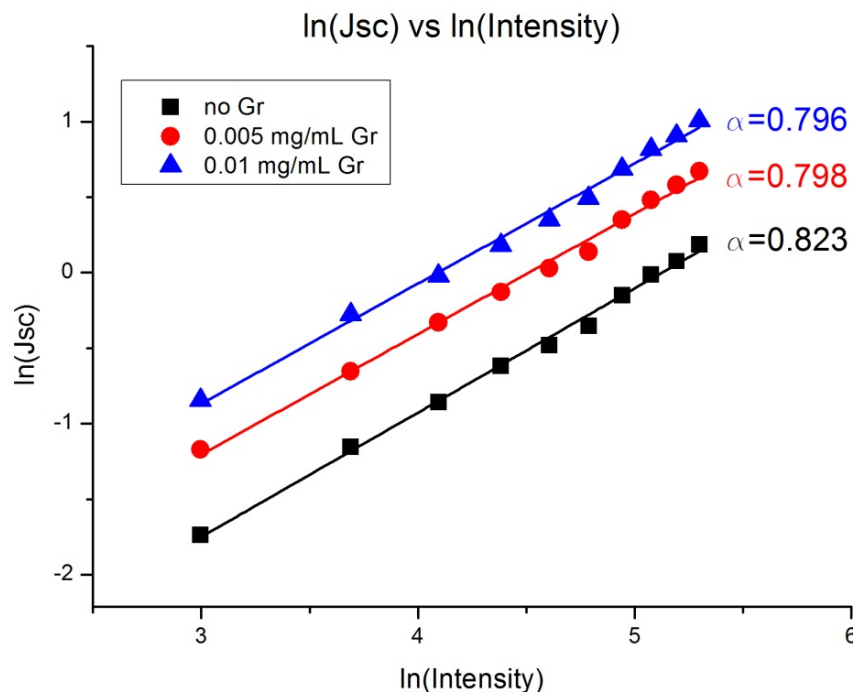
# Device Characterization(cont.)

## ➤ Recombination mechanism

$$J_{sc} \sim P_{In}^{\alpha}$$


$\alpha=1$ : monomolecular(geminate) recombination

$\alpha=0.5$ : bimolecular(non-geminate) recombination



greater bimolecular recombination

# Conclusions

- Adding small fraction of graphene greatly enhances charge transport and leads to much better  $J_{sc}$  and 
- Cells with more than 90% P3HT are viable
- Introduction of graphene in active layer leads to change of morphology
- Device physics change with increasing graphene fraction

# Future Work

- Better dispersed and oriented graphene via morphological control
- Increase FF by reducing interfacial roughness
- Stability and device encapsulation

**FY and VKK thank UC and the URC for funding and support**

# Thank you!

