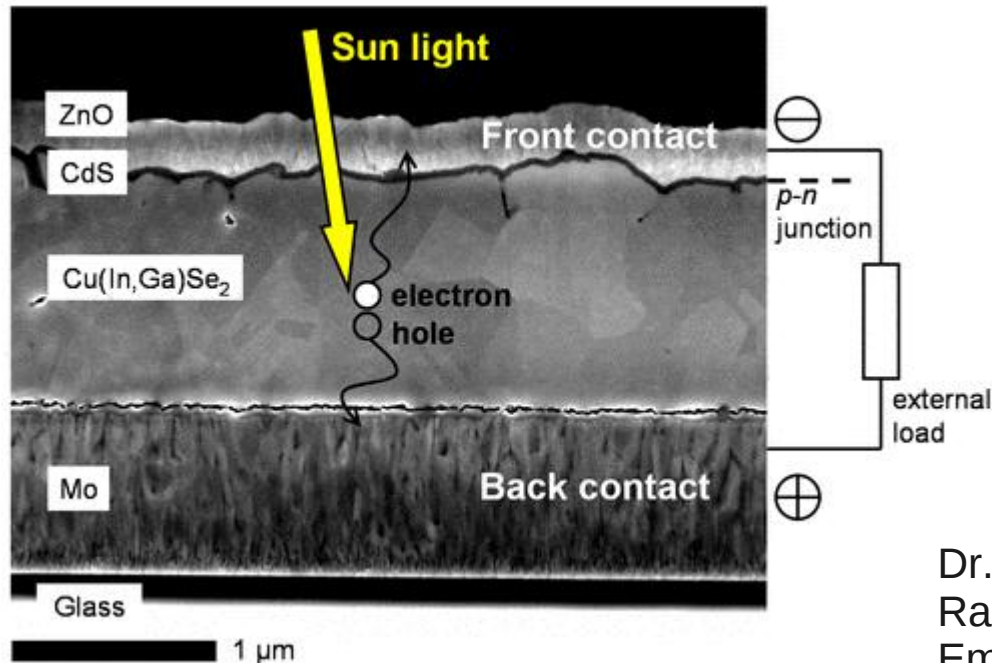
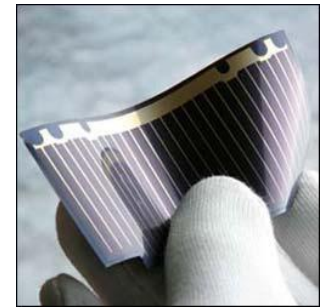
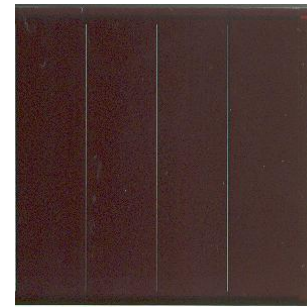
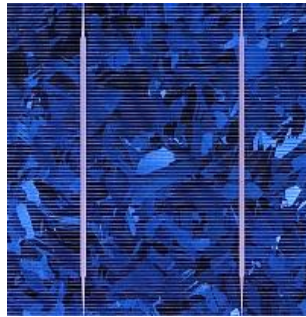


## Dye Sensitized Solar Cells (27027-01)

(Dienstag, 8:00-10:00 Departement Physik, Seminarzimmer 3.12)



Dr. Thilo Glatzel  
 Raum 3.04  
 Email: [thilo.glatzel@unibas.ch](mailto:thilo.glatzel@unibas.ch)



## Übersicht der Vorlesung

<b>22.02.2011</b>	allg. Einführung in die Solarenergie
<b>01.03.2011</b>	Physikalische Grundlagen der Photovoltaik I
<b>08.03.2011</b>	Physikalische Grundlagen der Photovoltaik II
<b>15.03.2011</b>	(Fastnachtsferien)
<b>22.03.2011</b>	Photochemische und photoelektrische Methoden der Energiewandlung
<b>29.03.2011</b>	Aufbau der Farbstoffsolarzelle, vgl. org. Solarzelle
<b>05.04.2011</b>	TiO <sub>2</sub> Nanopartikel als Substrat der Farbstoffsolarzelle
<b>12.04.2011</b>	Geeignete molekulare Farbstoffe zur Sensibilisierung
<b>19.04.2011</b>	Funktionsweise und Alternativen für den Elektrolyten
<b>26.04.2011</b>	(Osterferien)
<b>03.05.2011</b>	(FANAS meeting)
<b>10.05.2011</b>	Experimentelle Methoden zur Solarzellen-Charakterisierung
<b>17.05.2011</b>	Experimentelle Methoden zur Solarzellen-Charakterisierung
<b>24.05.2011</b>	Bau und Charakterisierung eigener Solarzellen
<b>31.05.2011</b>	



# Aufbau der Farbstoffsolarzelle und Vergleich mit organischen Solarzellen

- Nanoporous structure of the DSSC
- Sensitization by dye molecules
- Absorption of dye molecules on rutile  $\text{TiO}_2$
- Principle of operation
  
- Organic solar cells
- Setup of a device
- Working principle
- Structural properties / KPFM measurements

## Seminars

Adrian Hodel

*Chem. Rev.* **1995**, *95*, 49–68

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### Light-Induced Redox Reactions in Nanocrystalline Systems

Anders Hagfeldt<sup>†</sup> and Michael Grätzel<sup>\*</sup>

*Institut of Physical Chemistry, Swiss Federal Institute of Technology, CH-1015 Lausanne, Switzerland*

*Received June 9, 1994 (Revised Manuscript Received August 8, 1994)*

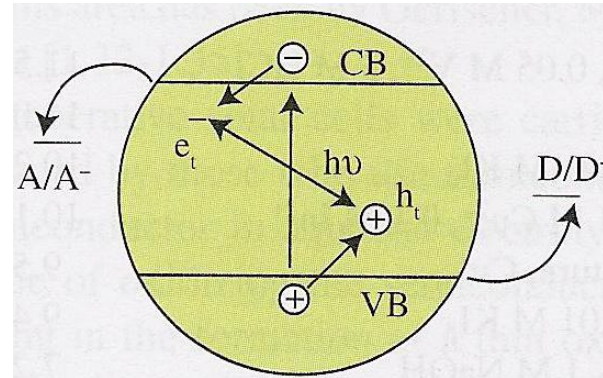
Clevin Handschin



### Photoelectrochemical complexes for solar energy conversion that chemically and autonomously regenerate

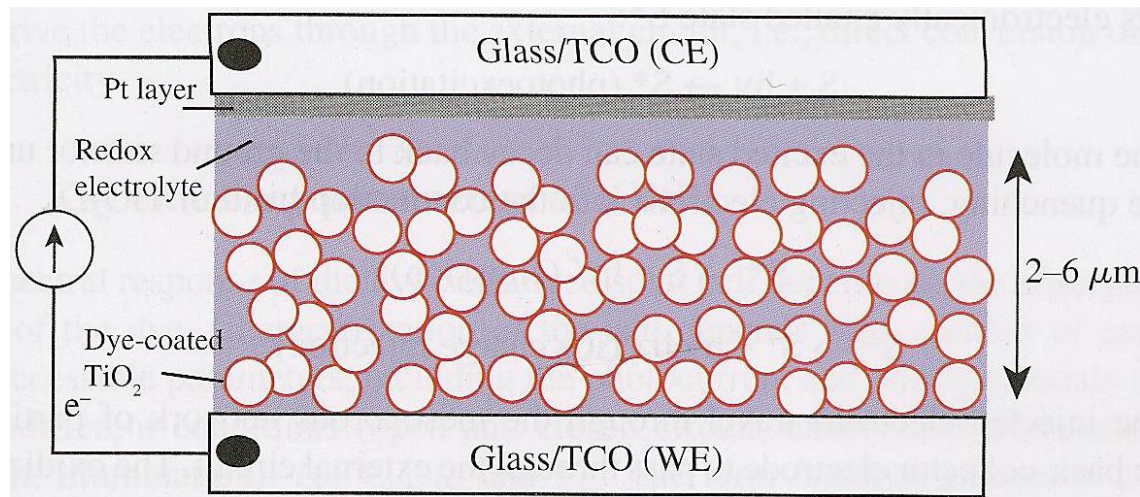
Moon-Ho Ham<sup>1†</sup>, Jong Hyun Choi<sup>2†</sup>, Ardemis A. Boghossian<sup>1†</sup>, Esther S. Jeng<sup>1</sup>, Rachel A. Graff<sup>1</sup>, Daniel A. Heller<sup>1</sup>, Alice C. Chang<sup>1</sup>, Aidas Mattis<sup>3</sup>, Timothy H. Bayburt<sup>3</sup>, Yelena V. Grinkova<sup>3</sup>, Adam S. Zeiger<sup>4</sup>, Krystyn J. Van Vliet<sup>4</sup>, Erik K. Hobbie<sup>5</sup>, Stephen G. Sligar<sup>3</sup>, Colin A. Wraight<sup>3</sup> and Michael S. Strano<sup>1\*</sup>

# Photoredox reactions of colloidal semiconductors and particulates



Both forms of photo-generated charge carriers reach the surface  
Low cost efficient system for degrading toxic waste

## Schematic Representation of a DSSC



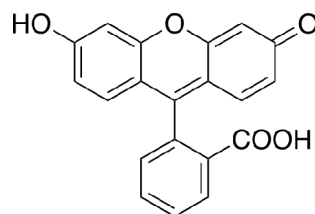
Metal oxide semiconductor nanoparticles:  
ZnO, TiO<sub>2</sub>, SnO<sub>2</sub>, In<sub>2</sub>O<sub>3</sub>, SrTiO<sub>3</sub>

Sensitized with: ruthenium polypyridyl complexes, or organic dyes  
Rhodamine B, rose bengal (xanthenes), fluorescein, and  
alkylthiacarbocyanines

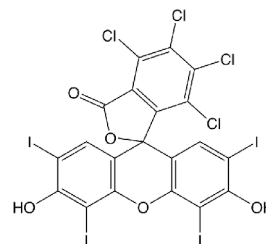


## Common dye molecules

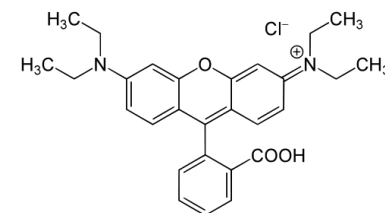
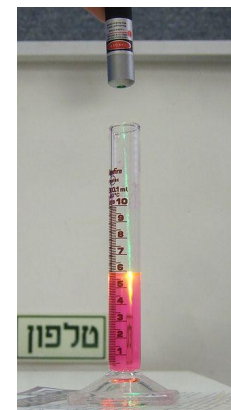
- The Ru-complexes are very stable to light, electricity, and heat because the bonds between the central metal ion and polypyridine ligands are usually very strong



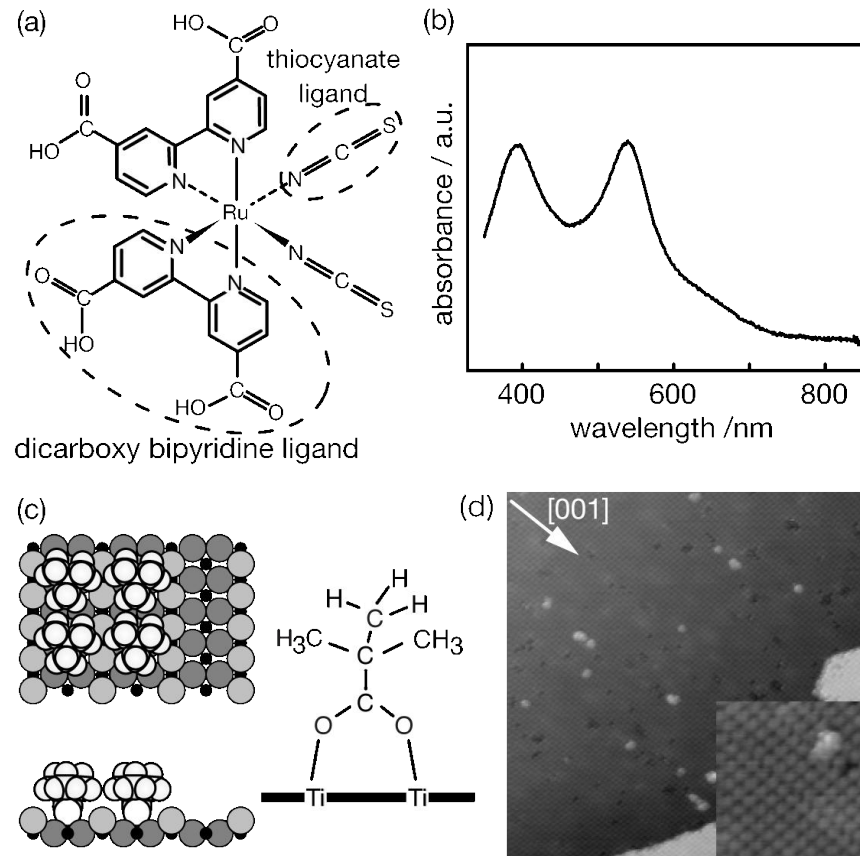
Fluorescein



Rose Bengal



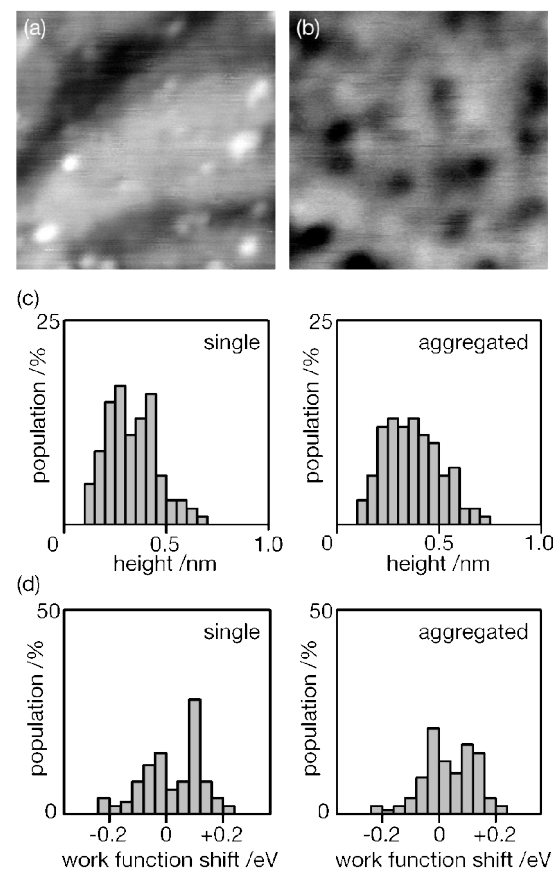
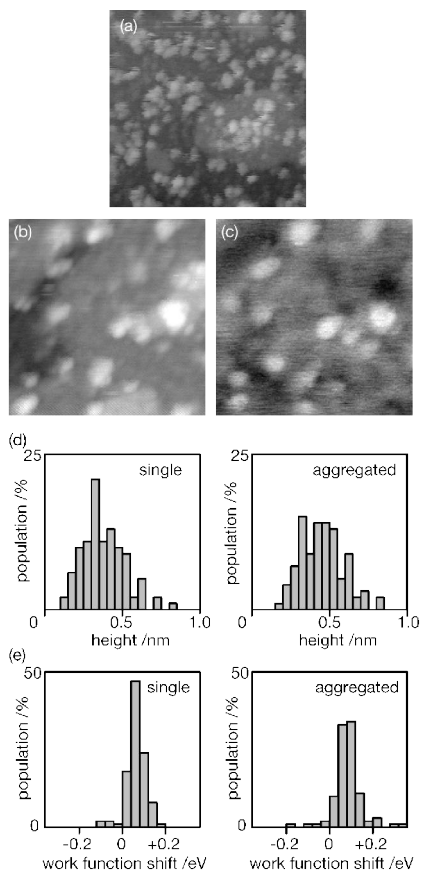
Rhodamin B

KPFM analysis of dye molecules on  $\text{TiO}_2$ 

N3, was adsorbed on an atomically flat  $\text{TiO}_2$  (110)



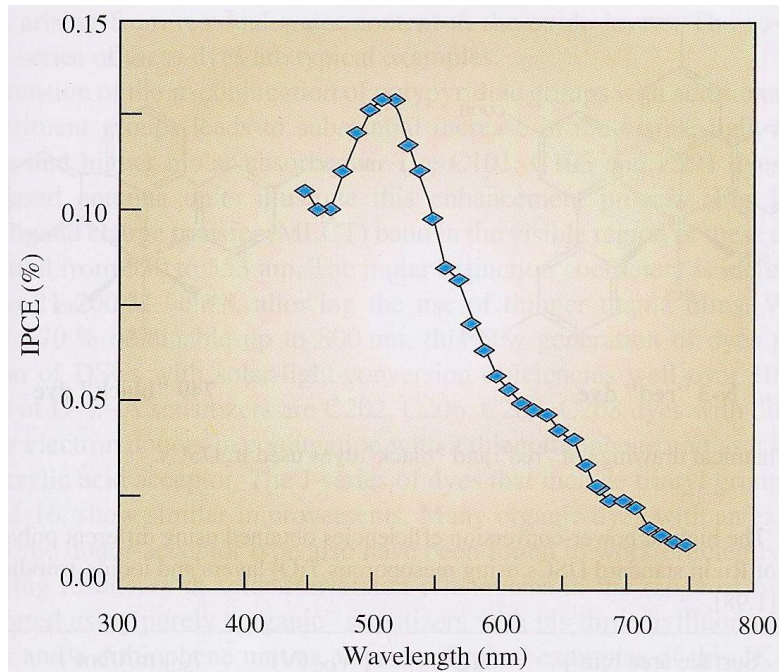
## KPFM analysis of dye molecules on $\text{TiO}_2$



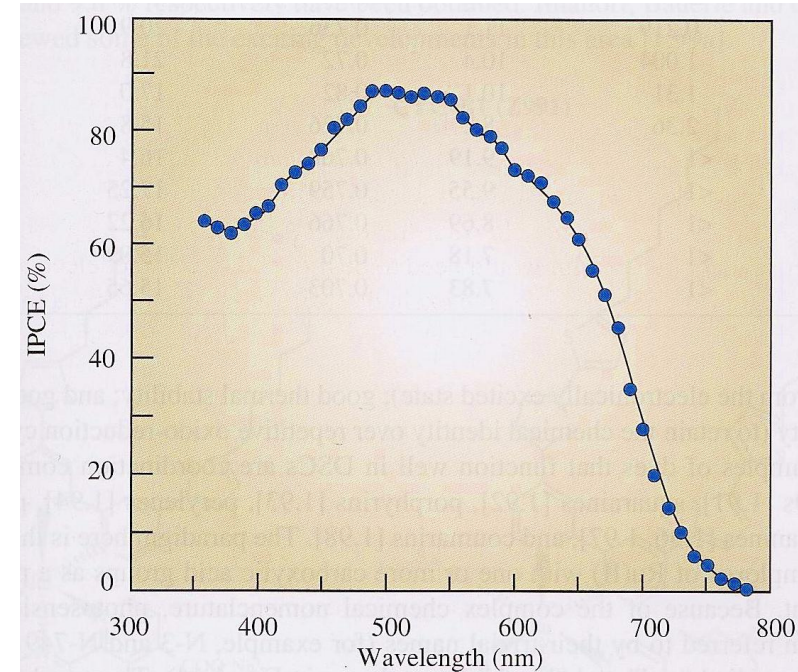
## Comparison of IPCE single crystal/mesoporous

Ruthenium N719 dye, deposited on:

Single crystal  $\text{TiO}_2$  (anatase)



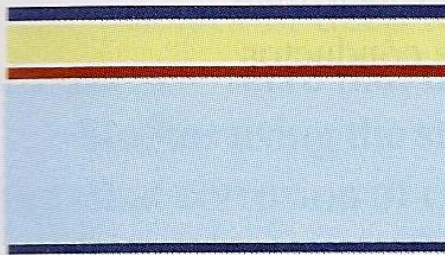
Mesoporous  $\text{TiO}_2$



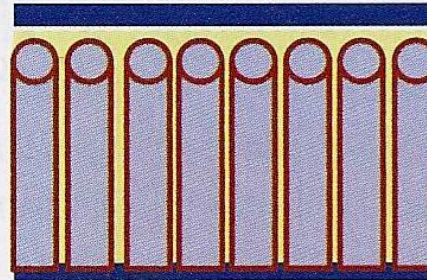


## Nanostructuring of the $\text{TiO}_2$

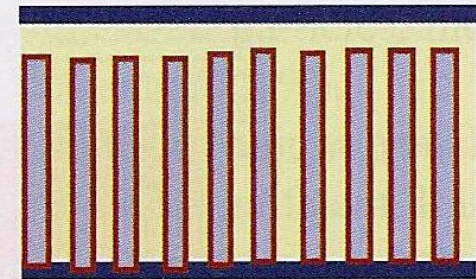
1. Bulk crystalline



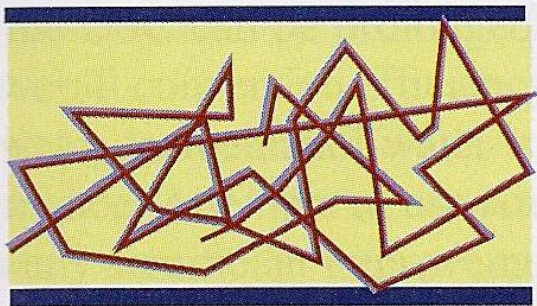
2. Nanotubes



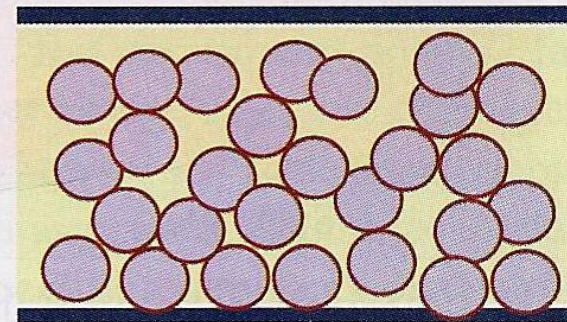
3. Nanowires



4. Interpenetrating polymer  
(bulk heterojunction)

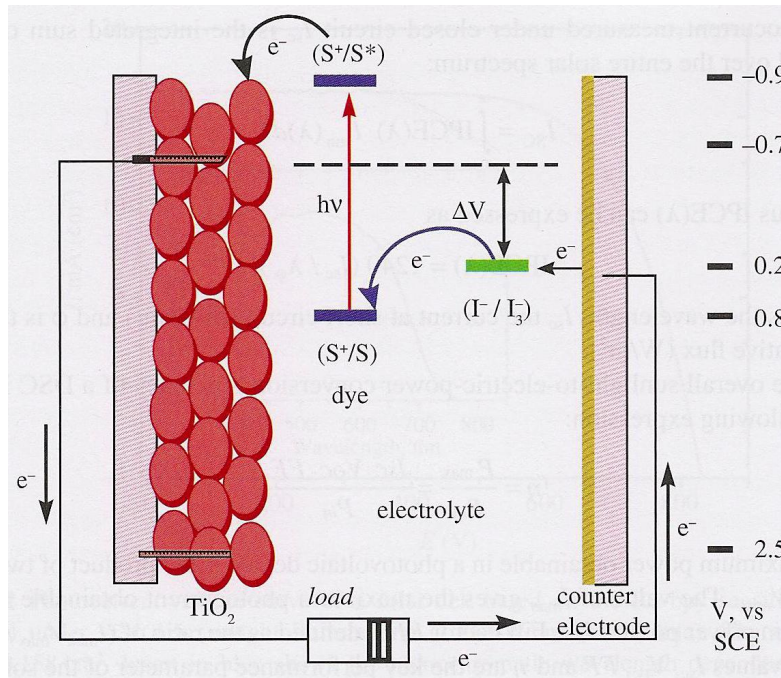


5. Mesoporous assembly  
based on nanosized colloids



Morphologies available for increasing the surface area and degree of interaction

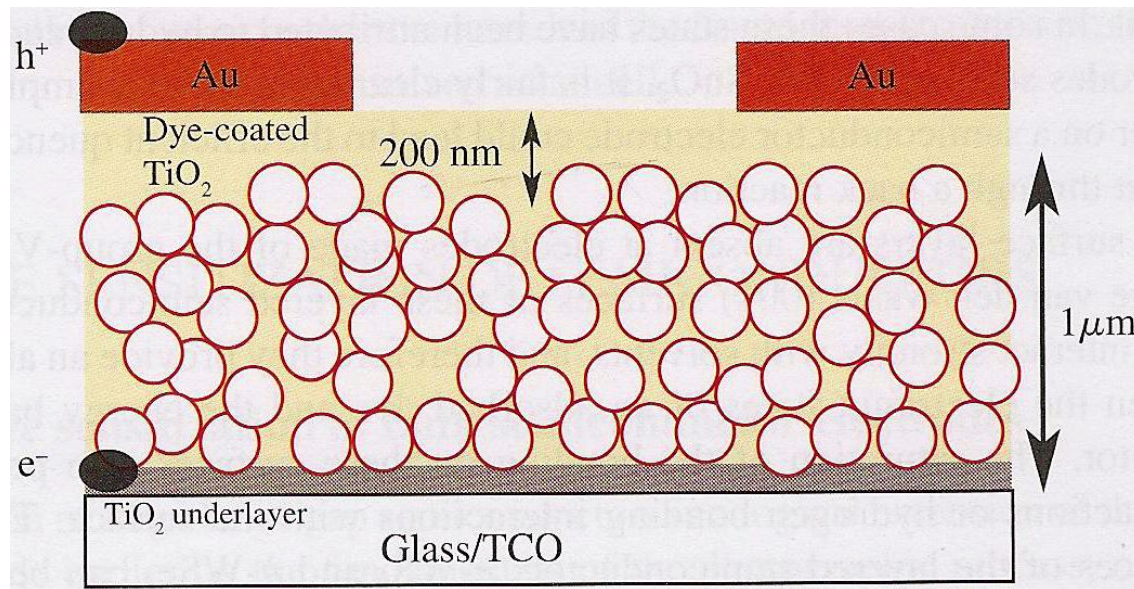
## Principle of Operation



- $S + h\nu \rightarrow S^*$   
(photoexcitation)
- $S^* \rightarrow S + h\nu$   
(emission)
- $S^* \rightarrow S^+ + e^- - cb$   
(TiO<sub>2</sub> charge injection)
- $2S^+ + 3I^- \rightarrow 2S + I_3^-$   
(regeneration of S)
- $S^+ + e^-(TiO_2) \rightarrow S$   
(recombination)
- $I_3^- + 2e^- \rightarrow 3I^-$   
(regeneration of I<sup>-</sup>)



## Quasi Solid State DSSC



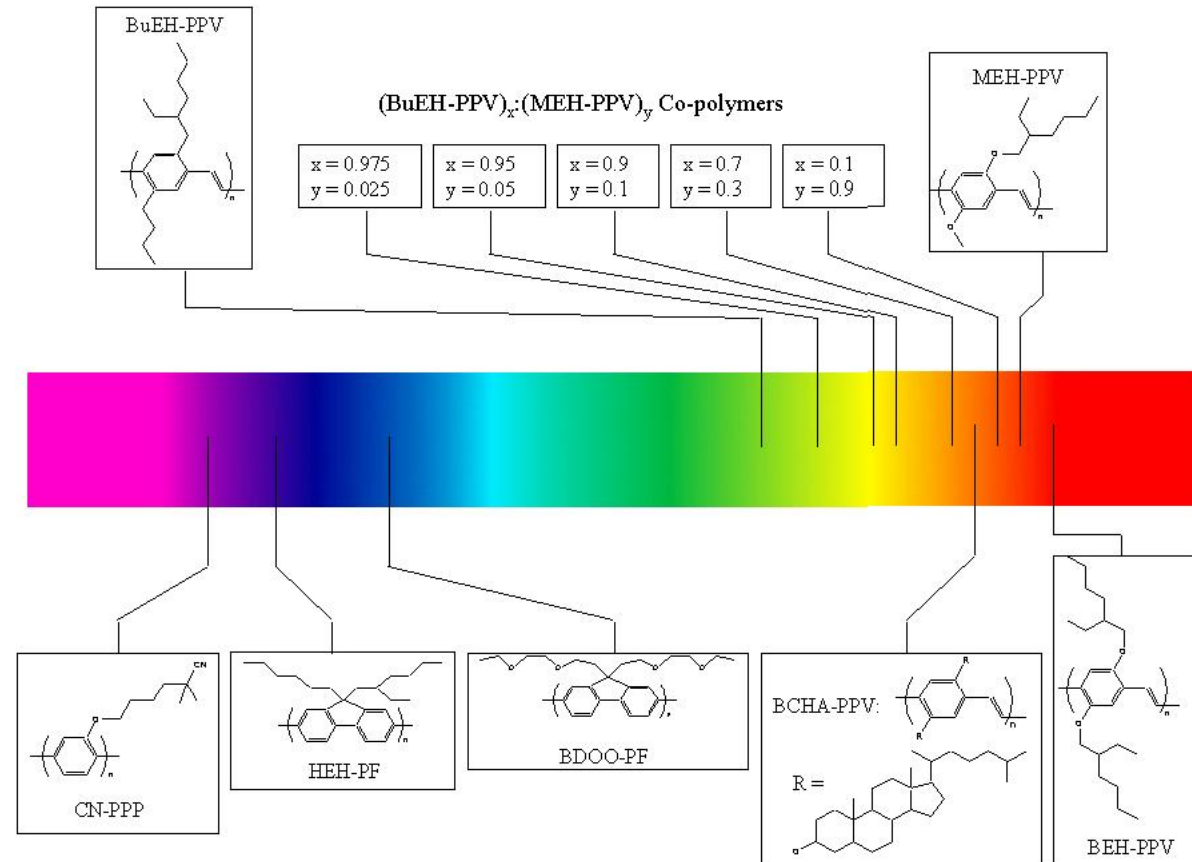
Organic hole transporter: triarylamine

## Nobelprize for Chemistry 2000



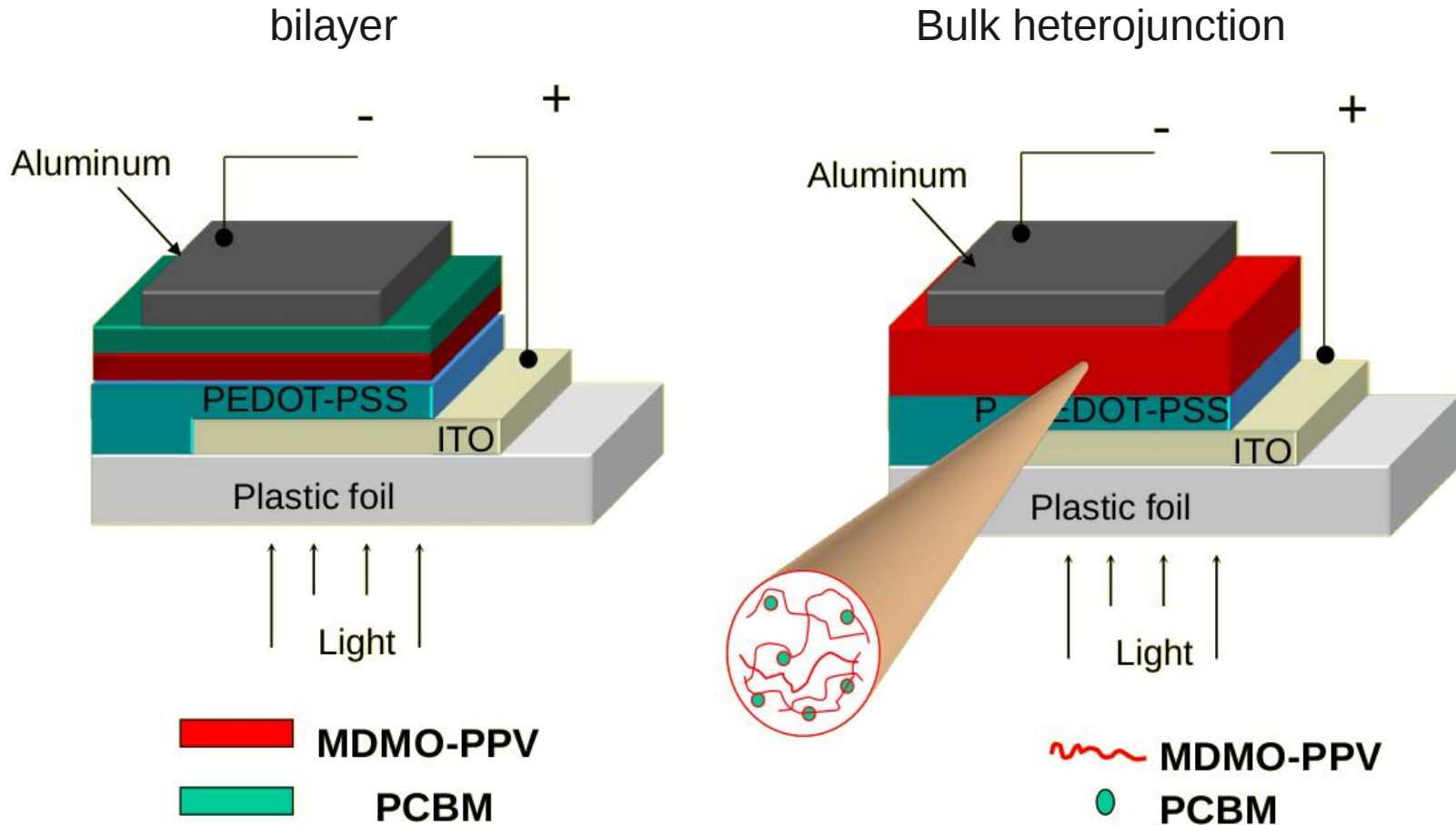
- Alan Heeger, Alan MacDiarmid (†) and Hideki Shirakawa
- 1974: Discovery of metallic conductivity in iodine doped trans-polyacetylene  $(CH)_x$

## Color Variations: Band Gap Engineering



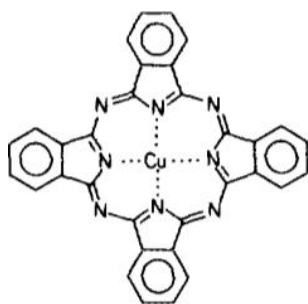
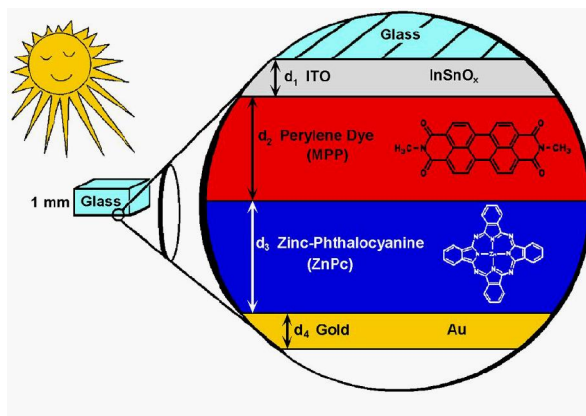


## Organic Solar Cell Device Geometries

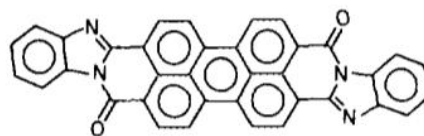


PEDOT-PSS: is used as a transparent, conductive polymer  
 MDMO-PPV: Donor  
 PCBM: Acceptor

## Small Molecular Organic Solar Cells "Tang-Cell"



CuPc



PV

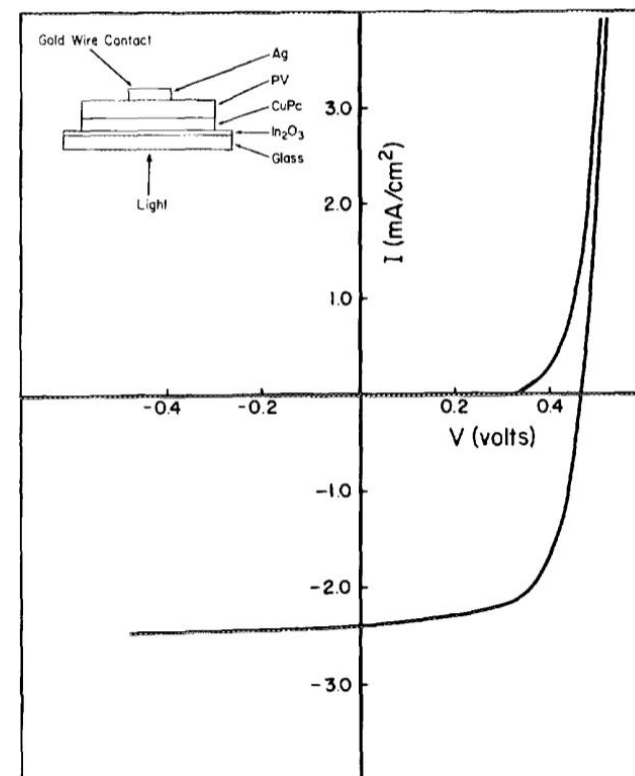
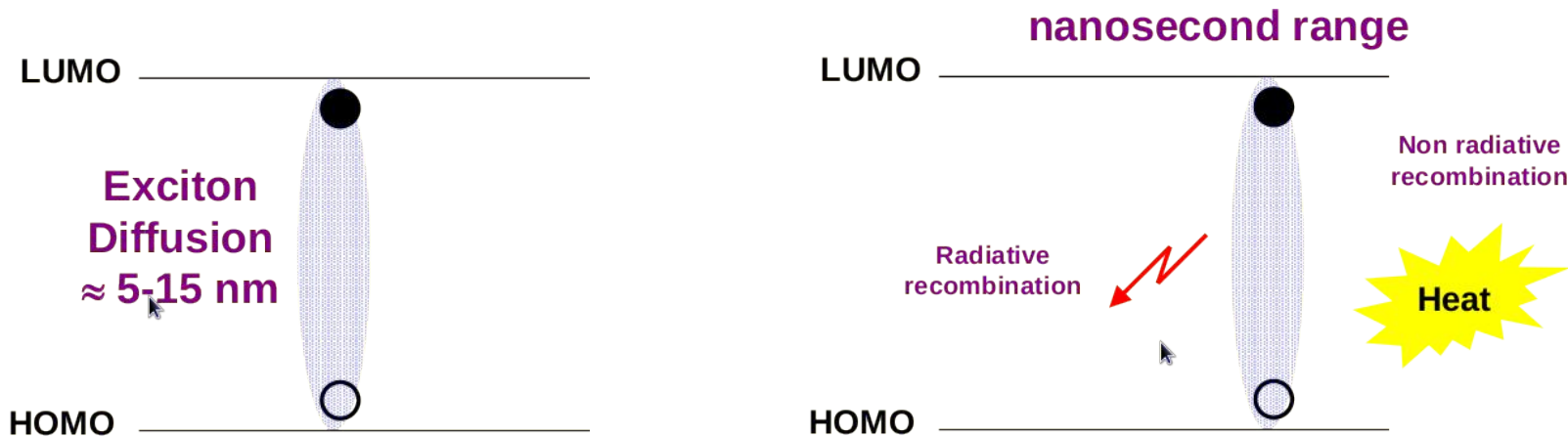


FIG. 1. Configuration and current-voltage characteristics of an ITO/CuPc (250 Å)/PV (450 Å)/Ag cell.

## Working Principle Photoexcitation in conjugated polymers

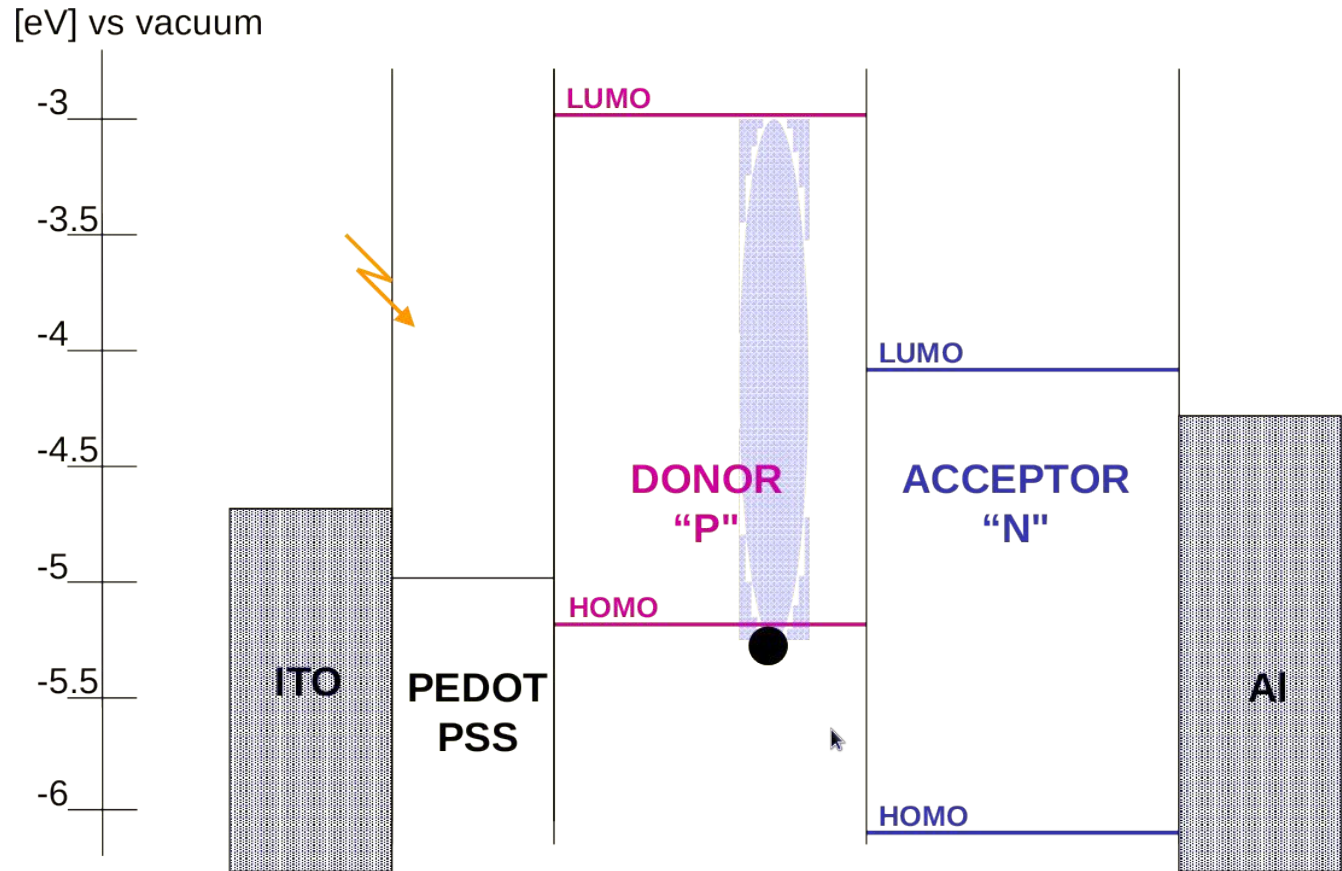
### Conjugated Polymer



Need to dissociate the exciton...

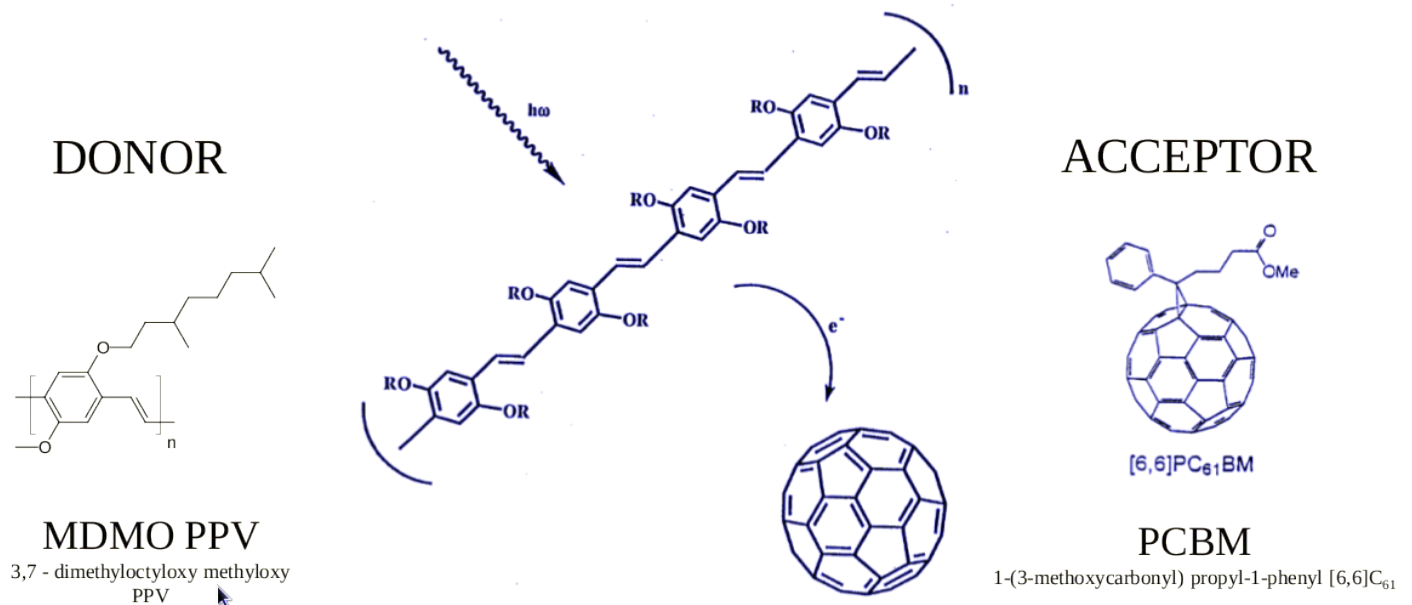


## Bi-layer polymer solar cells



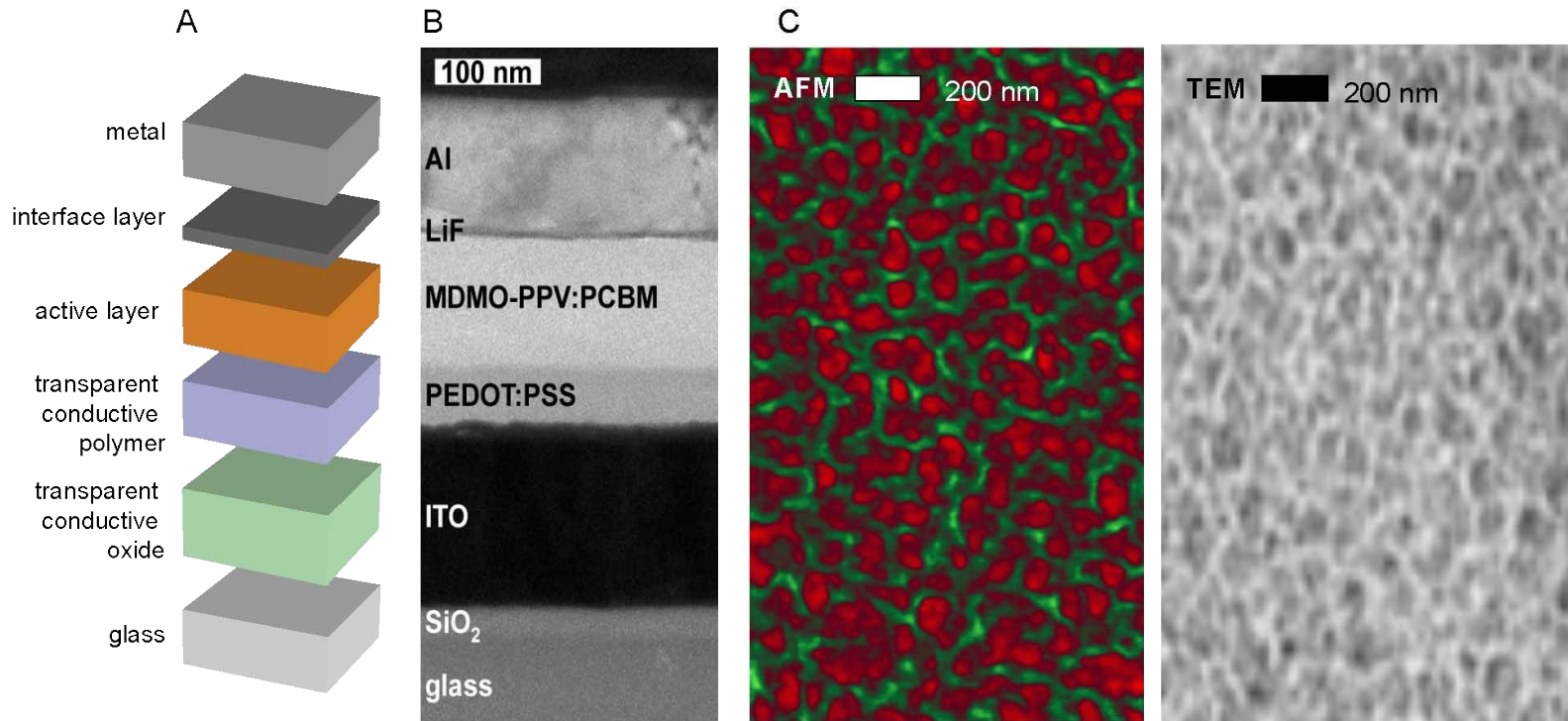
## Photoinduced Charge Generation

An ultrafast  $e^-$  **transfer** occurs between Conjugated Polymer / Fullerene composites upon illumination. The transition time is less than 40 fs. The Internal Quantum efficiency of charge generation is therefore  $\sim 100\%$ .



N. S. Sariciftci, L. Smilowitz, A. J. Heeger and F. Wudl., *Science* **258**, 1474 (1992)

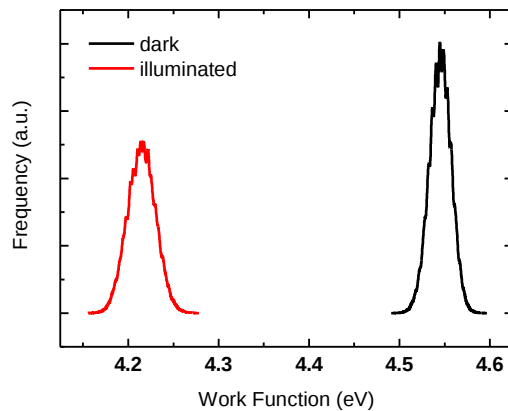
## Bulk Heterojunction Device Structure



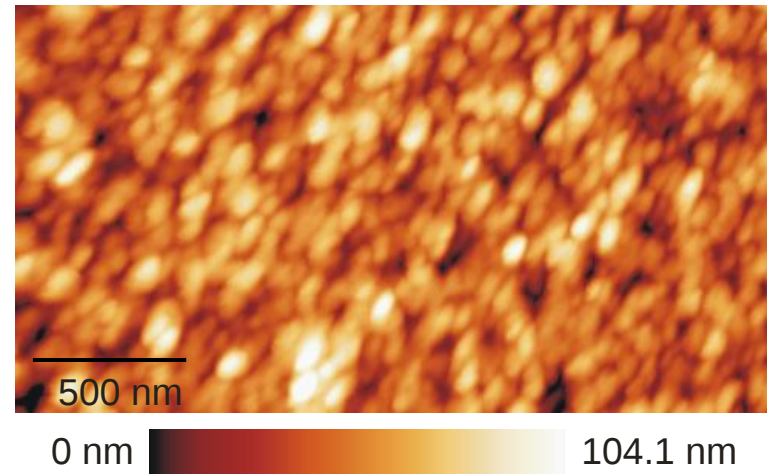


## KPFM: Chlorobenzene Blend – MDMO-PPV/PCBM

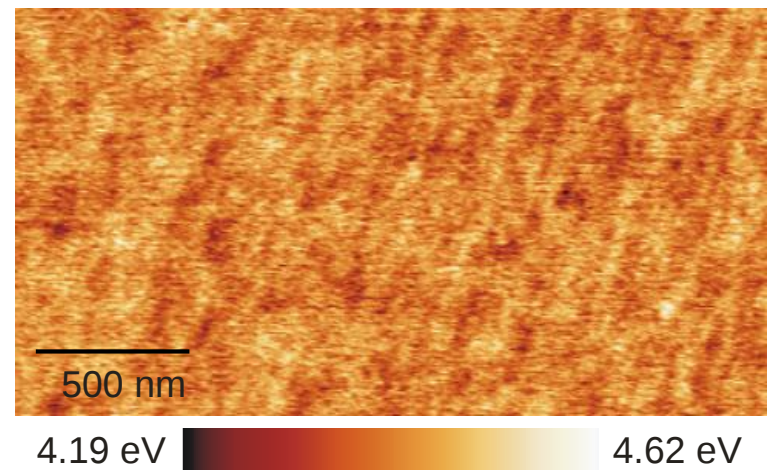
- Deposition and transport within Ar<sup>+</sup>-atmosphere
- Tip calibration on HOPG
- Laser diode illumination (~70mW/ 675nm)



Topography

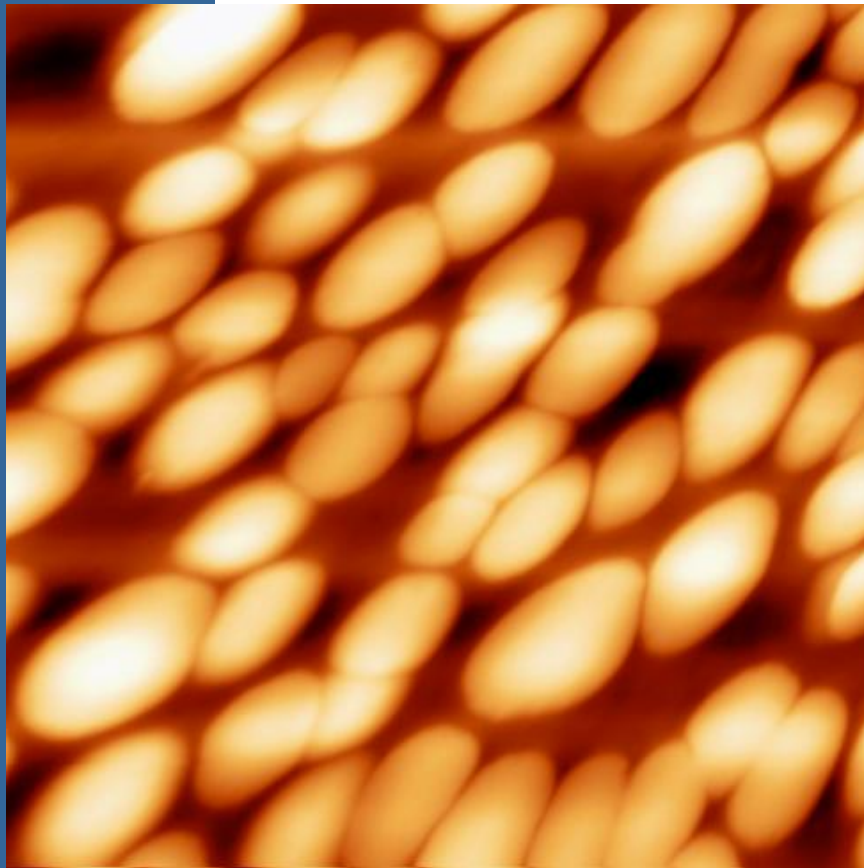


Work function, dark

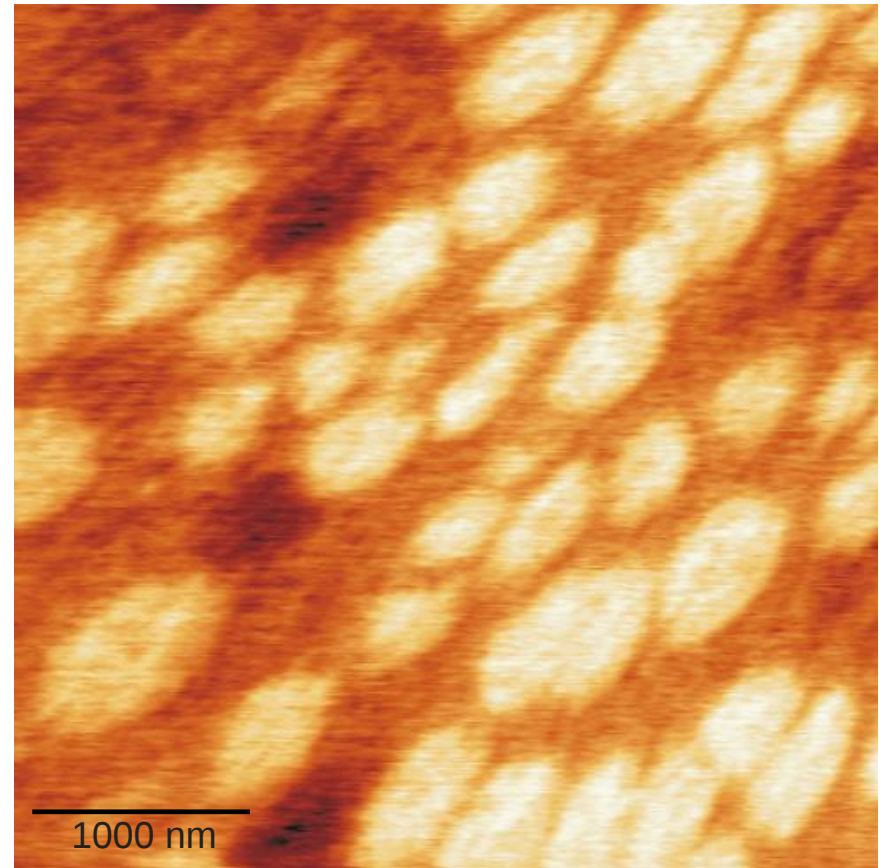




## KPFM: Toluene Blend – dark



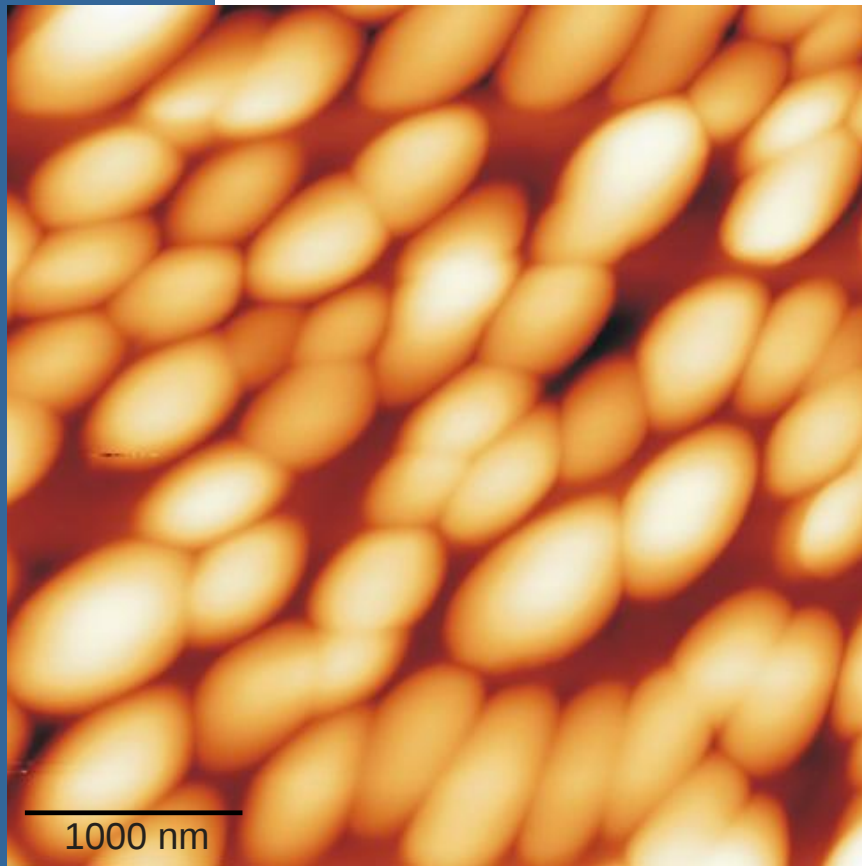
0 nm  104.1 nm



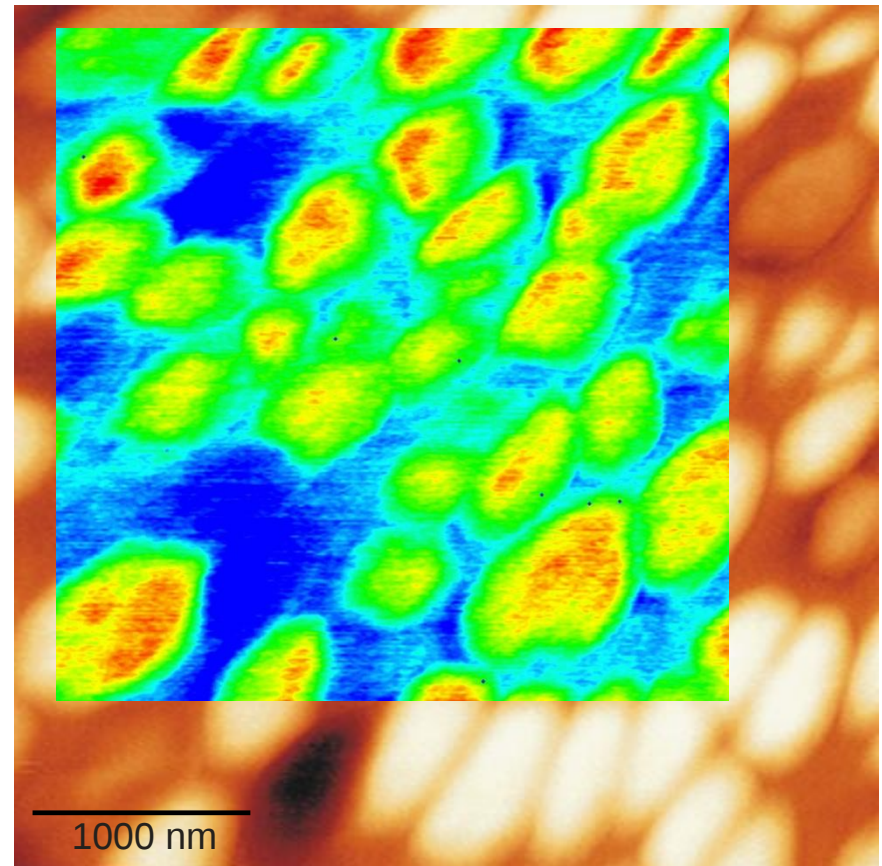
1000 nm



4.36 eV  4.57 eV

## KPFM: Toluene Blend – 675nm



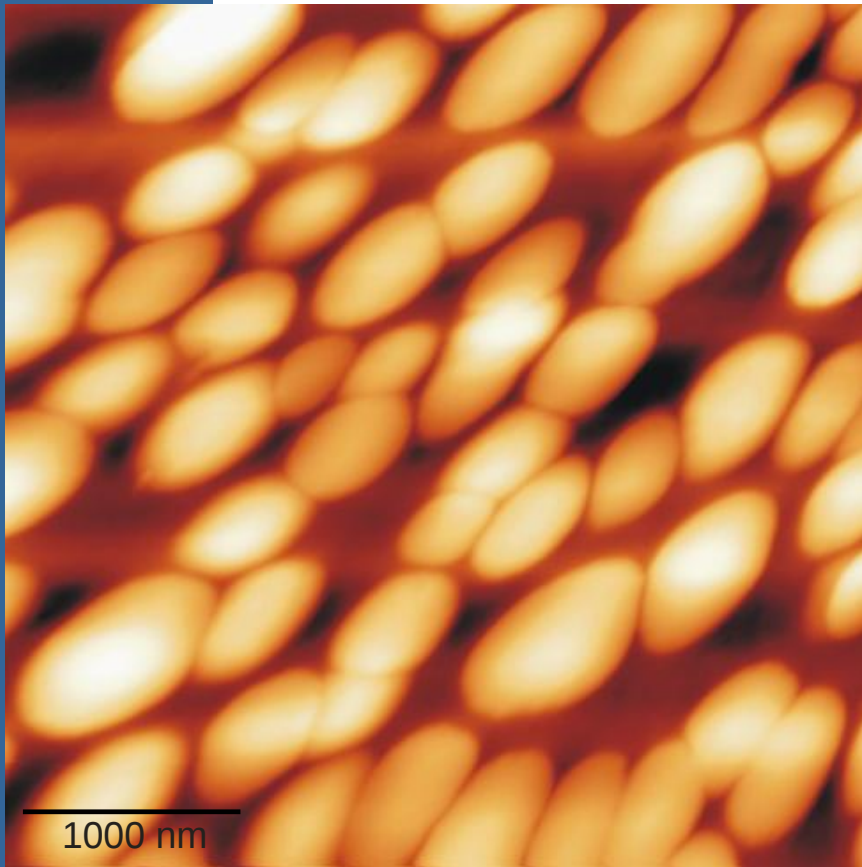
0 nm  105.6 nm



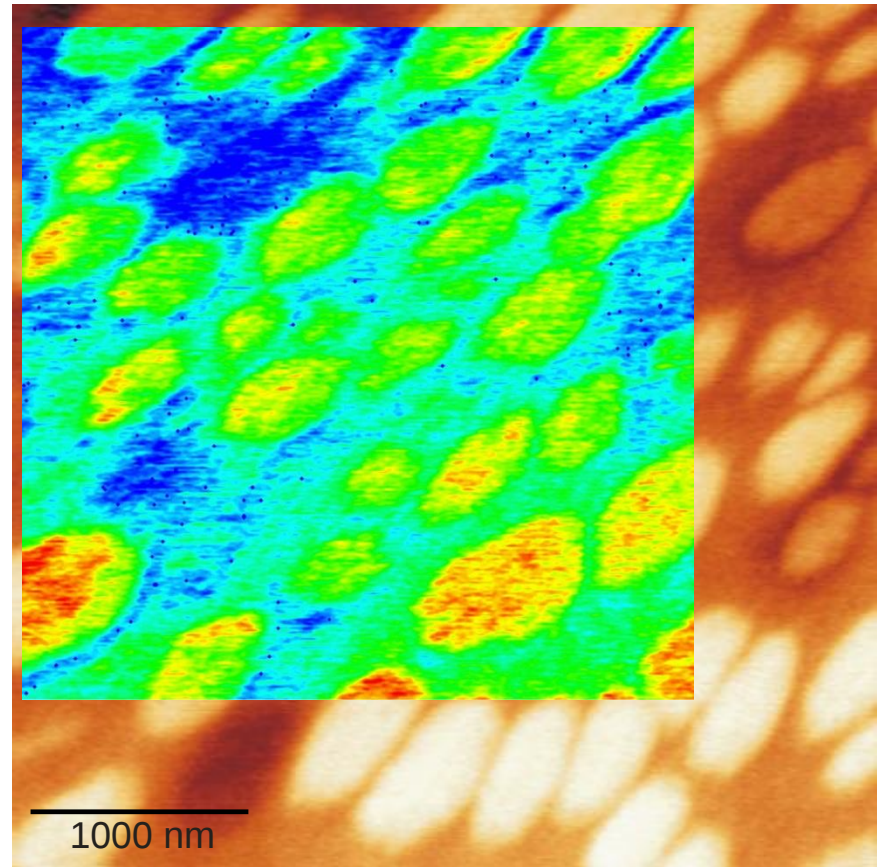
4.19 eV  4.62 eV  
-50 mV  220mV



## KPFM: Toluene Blend – 442nm



0 nm  106.8 nm

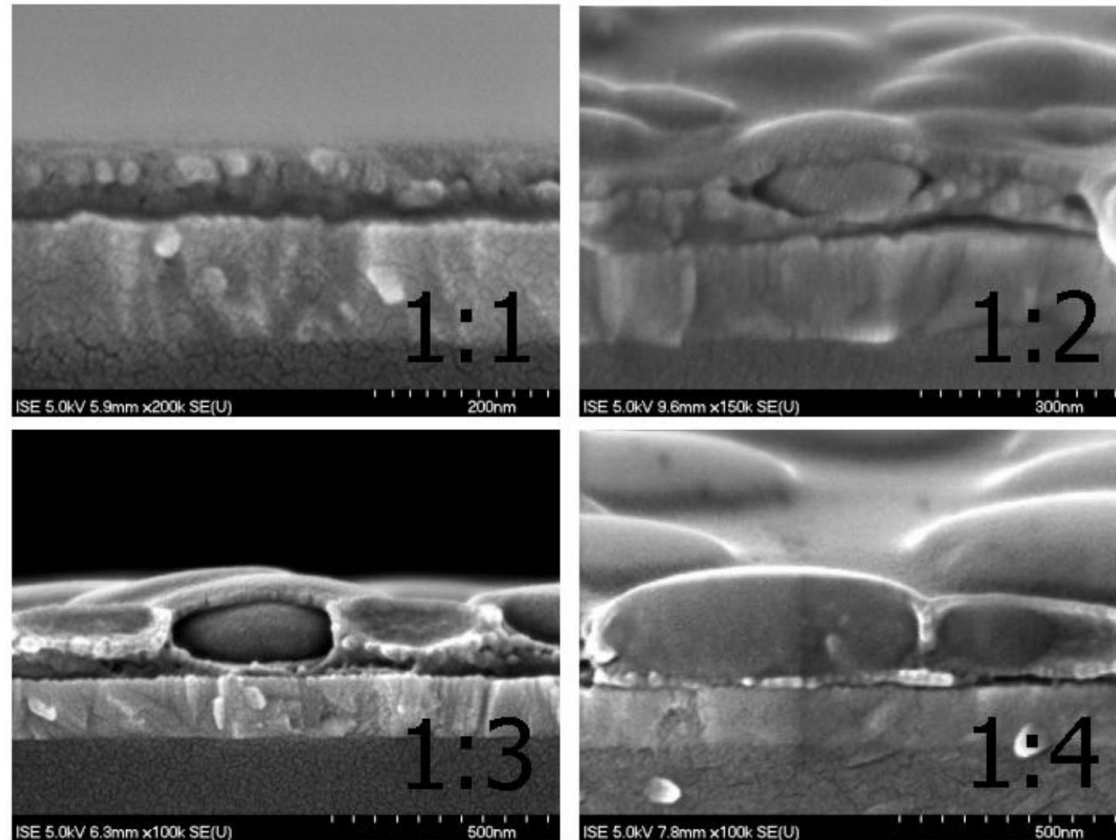


4.34 eV  4.70 eV

-30 mV  150mV

## Nanomorphology Effects-SEM Studies

### Chlorobenzene blend



### Toluene blend

