



Nanostructured Organic and Hybrid Solar Cells

Jonas Weickert, Ricky B. Dunbar, Holger C. Hesse, Wolfgang Wiedemann, and Lukas Schmidt-Mende

Department of Physics & Center for NanoScience (CeNS) Ludwig-Maximilians University (LMU) Munich

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Heidi Potts



Outline

- Introduction
- Operating principle
- OPV and HSC
- Device Morphologies
- Outlook



Solar Cell Efficiencies



Operating principle



- Cell absorbs light
- Electron-hole pair (exciton)
- Binding energy of exciton:
 - ~ 0.5 eV
- Heterojunction with offset between HOMO und LUMO
- Charge separation
- Charge extraction by electrodes



OPVC versus HSC

- Organic PhotoVoltaic Cell
- Organic polymers:
 - light absorption
 - charge transport
- Change absorption spectra:
 - polymer length
 - functional groups
- Flexible and low cost
- Low efficiency and stability

- Hybrid Solar Cell
- Organic polymers:
 - light absorption (donor)
 - hole transport
- Semiconductors:
 - acceptor
 - electron transport
- Example: DSSC



Optimal device morphology

- Domain size of donor and acceptor material should be small compared to the exciton diffusion length (~10 nm for polymers)
- Electrochemical potential drop at donor-acceptor interface must be large enough to overcome exciton binding energy (~0.5 eV)
- Exciton must split into free charges before recombination
- Electron and hole mobility must be high enough to extract charges (and they should be equal)
 - => Minimize charge carrier recombination
 - => Optimal morphology still unknown



Approaches for OPVC



- Small molecules
- Di-Block Co-Polymers
- Nanoimprint Lithography



[4] J. Björk et. al. *Physical Chemistry Chemical Physics.* **2010**, 12, 8815–8821

Approaches for HSC



P3HT within TiO_2 nanowires on FTO glass

Poly(3-hexylthiophen-2,5-diyl) (P3HT)

- Absorbing material: dye
- N type nanostructures:
 - metal oxides: ZnO or TiO₂
- P type semiconductors:
 - e.g. P3HT
- Nanowires
 - regularity and orientation
- Nanutubes
 - higher surface area



Core-shell structures



Idealized core-shell metal oxide-P3HT HSC

- High potential for HSC
 - charge transport
 - avoid recombination
- Properties
 - rutile TiO₂ nanowire
 - Transparent concuctive Oxide
 - Anastase TiO₂ for hole blocking
 - TiCl₄ treatment
 - Dye for absorption
 - Nanostructures ~15 nm
 - PEDOT for hole-collection
 - active layer thickness ~1 μm



Open questions

- Exciton separation only in the direction of the external field (given by the work functions of the metal electrodes)?
- Interface design to efficiently enable charge injection but hinder charge recombination?
- Maximum size of the nanostructure to allow efficient charge transport?
- Additional effects based on the nanodimensions when getting smaller than 10 nm?
- Can efficiency be increased by light trapping approaches?

=> Defined nanostructures might give the answers!



Thank you for your attention.

