CORE-CM develops teams across CNS, Engineering and Nano-Medicine

Center mission

The CORE-CM mission is to promote excellence in materials research on the MSU campus through high quality multidisciplinary group research, infrastructure development, training programs and technological innovation.

Developing groups competitive for federal group and center calls, along with shared infra-structure projects are primary goals. Both academic and industry based groups are promoted.

Some Research Groups (lead faculty):
Thermoelectric materials (Morelli)
Organic solar (Lunt, McCusker)
Nanotoxicology (Worden)
Ultrafast diffraction (Ruan)
Battery materials (Drzal)
Fuel cells (Barton, Reguera, Promislow)
Materials in extreme environments (Bollen)
Ruan - Designing and building the worlds fastest e-microscope

Other members of microscope MRI development team: (Berz, Crimp, Duxbury)
Also affiliated with NSCL/FRIB materials groups
New Frontiers in Sciences Enabled by fs-EM

**Imaging Material Transformation**
- Electron holography of superconductor (Lichte, Tonomura)
- Graphite to diamond transition through photoexcitation (Ruan, Tomanek, Drzal, MSU)

**Imaging Molecular Function**
- Free energy landscape of molecules
- Photo-interconversion of two isomers of azobenzene (Barret, McGill)
- Au Nanocatalyst Supported on TiO2 (Zilin Wu, ORNL, Landman, Georgia Tech)

**Imaging Electron Dynamics**
- Photovoltaic processes in nanoparticle (Kilmov, LANL)
- Carrier dynamics in quantum nanostructures (Kim, Culobia; McEuen, Cornell)

**Imaging Biological Function**
- Solvated protein in nanocavity (water) (H. Grubmüller, Max Planck Inst.)
- Protein folding dynamics (Dobson, Cambridge)

**Key areas**
- High-Tc superconductor
- Laser machining
- Water splitting
- Photo-catalysis
- Nano-electronics
- Photo-voltaics
- Protein-folding
- Drug discovery
FRIB/NSCL – CORE-CM materials group (SPG)
Leverage unique MSU expertise and capabilities to address extreme materials problems of critical local and national importance

- **Develop and study key materials** under extreme radiation, stress, temperature conditions (Graphite, Diamond, Ti alloys)
- **Develop a broad MSU team** to leverage NSCL (and future FRIB) beam conditions and MSU’s expertise in material science and engineering

- **Physics and Astronomy** (Duxbury, Ruan)
- **Electrical and Computer Engineering** (Grotjohn)
- **Chemical Engineering and Material Science** (Boehlert)
- **National Superconducting Cyclotron Laboratory** (Mittig, Stolz)
- **Facility for Rare Isotope Beams** (Bollen, Pellemoine, Ronningen)

Use swift heavy ions to for nanolithography and to create nanostructured materials
Segregation of nanoparticles to polymer thin film surfaces and interfaces

Neutron reflectivity of PS NP in PS – Segregation to substrate due to entropy

TEM functionalized CdSe quantum dots in PS – NPs segregate to free surfaces due to dispersion forces
Physical interactions which control nanoparticle segregation

- Entropy is gained when nano-particles segregate to surfaces – e.g. relaxation of polymer confinement at hard substrate.
- Long range dispersion forces play a key role. Systems with similar refractive index have an effective attraction. (Monotonic dielectric ordering is favored).
- CdSe nanoparticles have a low n alkane layer at the surface. This leads to an effective dielectric constant which is less than PS (1.59), favoring segregation to the air/PS surface.
- CdSe with Pyridine coating anneal to a SiO2 (4nm) on Si substrate.
Characterization of NP-Polymer morphology

- NP concentration profile: Reflectometry
- Characteristic domain size: SANS

Kiel et al., Nanoparticle agglomeration..., PRL 2011

3-D real space models: Constrained optimization

\[ H = - \sum_{ij} J_{ij} S_i S_j + a \int dz |\rho^{ex}(z) - \rho^{m}(z)| + b|p_m - p_{ex}| \]

Olds and Duxbury, Percolating bulk heterostructures..., PRE 2012
Dynamic Monte Carlo with First Reaction Method

In our simulations:

• Excitons are generated uniformly in P3HT at a rate extracted from experiment and diffuse until they dissociate or decay (exciton decay length).
• When excitons find a donor/acceptor interface before decaying, they form a geminate pair.
• Geminate pairs diffuse on the interface until they recombine or dissociate into free carriers hole (pair recombination length).
• Free carriers diffuse until they recombine or are collected at an electrode (free carrier recombination length).
Submicrosecond Time Resolution Atomic Force Microscopy for Probing Nanoscale Dynamics

C-AFM images of annealed (a-b) and non-annealed (c-d) P3HT films. (a,c) show the topology, and (b,d) show the corresponding current maps.

The c-AFM image showing topography of annealed sample and IV-characteristic at fixed spots A and B.

C-AFM current maps and IV relations:

Nanofibers are clearly observed in annealed samples.

Different currents are measured on and off fiber.

Experimental Result for P3HT (J. Sun, at. al, 2012, preprint)
3D Continuum model of Hole Injection System

Poisson’s Equation

\[ \nabla \cdot (\varepsilon \nabla \psi) = -qn_p \]

Continuity Equations at steady state

\[ \nabla \cdot J_p = 0 \]

Drift-Diffusion Equations

\[ J_{p,\zeta} = -qn_p \mu_{p,\zeta} \frac{\partial \psi}{\partial \zeta} - qD_{p,\zeta} \frac{\partial n_p}{\partial \zeta} \]

\[ \vec{\mu}_p(\vec{r}) = \sum_{\zeta=1}^{3} \mu_{p,\zeta}(\vec{r}) \vec{e}_{\zeta} \]

where \( \zeta = x, y, z \)

Einstein relation of charged particles,

\[ D_{p,\zeta} = \mu_{p,\zeta} \frac{k_b T}{q} \]

Implementing fiber into the morphology, the experimental result can fit to the same value of background hole mobility around

\[ \mu_o \approx 4.15 \times 10^{-4} \text{cm}^2\text{V}^{-1}\text{s}^{-1} \]

MSU group, preprint 2012
Mesoscale Priority Research Direction
Taming complexity from atoms to the mesoscale: NP/organic hybrids

Opportunity
Nanoparticle (NP)/organic hybrids are the basis of many energy harvesting, storage and utilization systems. The functionality of these systems is controlled by NP dispersion and segregation, interface structure, organic crystallinity etc. Taming this complexity through control of synthesis from the atomic to micrometer (A->M) scales, guided by high precision static and dynamic characterization and modeling would be transformative across a broad range of materials systems and devices.

Meso Challenge
- Synthesis and processing to control A->M structure
- Characterization methods to generate 3-D interconnected A->M models consistent with experimental data.
- Characterization of the dynamics of charge transport across A->M length scales and fs to seconds time scales.
- Modeling of charge generation and reactive transport in experimentally consistent A->M architectures.
- Optimize A->M architectures for applications

Approach
Direct self-assembly using dispersion forces, entropy and electrostatics toward either equilibrium or strongly metastable structures. Polymer architecture, solvent and/or thermal annealing etc to further control A->M structure. Combine scattering, reflectometry, AFM, SEM, TEM, nano-CT etc with modeling to find percolating 3-D real space A->M models. Use integrated Ab-initio, Kinetic MC, and continuum approaches with A->M model architectures to correlate performance with morphology and hence with synthesis and processing.

Impact
Success in multiscale synthesis, integrated A->M characterization and multiscale modeling would open many new opportunities for materials and device optimization. NP/organic hybrids are a good place to start as they have a broad knowledge base, they exhibit a wide range of novel physical and chemical phenomena, and they are of interest for a broad range of energy and other applications.

Contact: Phil Duxbury, MSU

M.C. Dennler, G. Scharber, C.J. Brabec, Polymer- Fullerene Bulk heterostructure solar cells, Advanced Materials, 21, 1232 (2009)