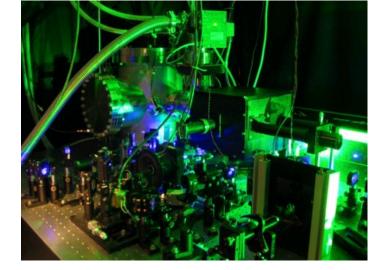
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Dipl. 0D. PhiPostdoInstitu	ation and Appoir Chem. ETH 1997, ETH I. Oxon 2001, Oxford octoral Fellow 2001-2004, ute of Standards and Tech rsity of Colorado, Boulder)	JILA (National nology and	 Physic Chemi Chemi Dynan Energy Instru Spectri 	arch Int al cal Physics cal Reaction nics/Kinetics, y Science ment Develo coscopy/mole ces and Solid	'Interactions pment ecular Struct			
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organic electronic devices such as organic photovoltaic cells, organic thermoelectrics or organic spin valves. Their function is largely controlled by interfacial processes such as exciton dissociation, polaron formation, geminate recombination and heat and carrier transport. Interfaces are intrinsically complex environments, with typically high defect densities and complex structure. Heterogeneity makes it difficult to extract the underlying physics of charge generation and transport in real operating devices. As a consequence, the development of solar cells with significantly higher power conversion efficiencies or more efficient spin valves and thermoelectrics presents a major challenge.



Research in LabMonti seeks to elucidate the chemistry and physics of carriers in organic semiconductors at interfaces *on the short length- and time-scales* present in organic photovoltaic, spin and thermoelectric devices. We develop and use novel forms of optical microscopy in combination with ultrafast photoelectron spectroscopy, scanned probe microscopy and state-of-the-art experiments at synchrotrons to study the electronic structure and dynamics of organic semiconductors under highly controlled conditions.

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Using this experimental approach, we are pursuing research in several complementary directions:

1. Interface formation and charge transport at organic/metal boundaries for organic solar cells, organic light emitting diodes and organic thin film transistors. We use two-photon photoemission spectroscopy and a gamut of photoelectron spectroscopies in conjunction with low-temperature scanning tunneling microscopy to study charge generation and transport at highly controlled interfaces. This approach lets us investigate both occupied (HOMO) and unoccupied (LUMO, higher lying excited states and sometimes transport levels) states at organic semiconductor interfaces, giving insight e.g. to the role of the interface on molecular energy levels and intermolecular interactions. Recent work has focused on the role of short- and long-range electrostatic fields at the interface, showing dramatic effects on interfacial electronic structure.

2. We study the complex interactions at controlled organic/inorganic semiconductor interfaces in order to understand the role of bandgap, defects and dopants on charge injection events in organic solar cells and models of dye-sensitzied solar cells. We use a variety of time-resolved and steady-state photoemission spectroscopies as well as synchrotron-based excited-state and time-resolved spectroscopies in order to unravel charge-transfer and chemical bond formation at these interfaces and its effect on charge-transfer events. Recent work has focused on the role of the band structure of layered semiconductors and transparent oxides on the interfacial electronic structure.

3. When organic semiconductors are in contact with a ferromagnetic electrode such as Co or certain oxides, charge-transfer can be spin-selective. This effect can be used for organic spin-valves, the basis of magnetic memory devices in computers. The process of spin-selection is dominated by the *spinterface*, the interface between electrode and organic semiconductor. We use time-, spin- and momentum-resolved photoelectron spectroscopy to investigate this process and develop an understanding of the fundamental processes at work. Recent work has looked at dynamical reasons for spin-selective charge injection into the organic semiconductor.

4. In order to obtain a detailed picture of the exciton dissociation process, we reduce the complex bulk heterojunction typically used in organic solar cells to a model system, where charge transfer occurs from a *single molecule* to a single crystalline wide bandgap semiconductor under highly defined conditions. Ensemble measurements often obscure some of the most intriguing aspects of such small-scale systems. This requires us to look at individual molecules in order to capture the complete picture and allows us to investigate effects of surface state densities, defect densities, distance etc. on exciton dissociation efficiencies. These experimental results are also accessible to high level ab inito calculations.

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