



**Presentation of Light-Matter Interactions for Middle School Students**  
**West Las Vegas Middle School, May 2011**  
**Daniel Casillo, Alexandr Fonari, Gary Angles, and Tatiana Timofeeva,**  
**NMHU, DMR 0934212**



**A series of presentations on light-matter interactions have been given at the West Las Vegas Middle School. These have served to promote connections between local schools and PREM program.**

**Three NMHU PREM students organized four different science presentations, each approximately one hour in length for 80 students total for the science class of Mr. Michael Tenorio (7th grade).**



**Topics covered:**

- Economic problems of electric power generation
- Light waves-particles
- Light sources
- Solar cells
- Flexible screens

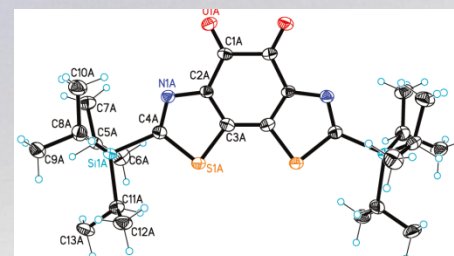
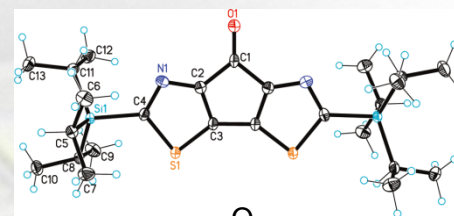
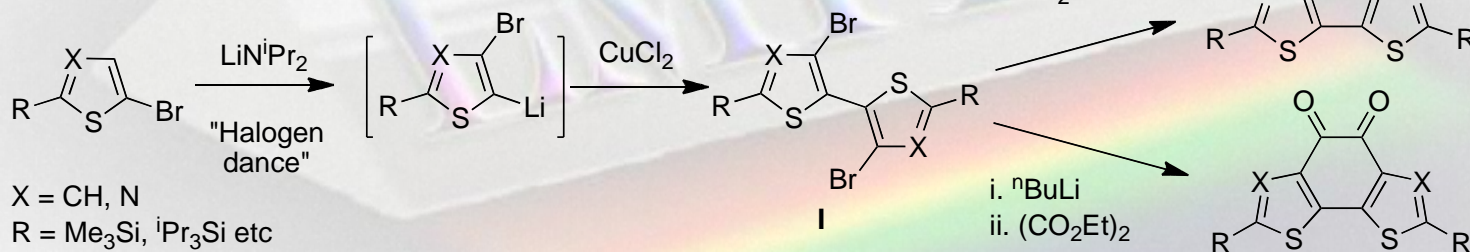


## Base-Catalyzed Halogen-Dance Route to (Di)Keto-Bridged Bithiophene & Bithiazole Building Blocks for Optoelectronics

Yulia Getmanenko,<sup>1</sup> Seth Marder,<sup>1</sup> Bhupinder Sandhu,<sup>2</sup> Paul Tongwa,<sup>2</sup>  
and Tatiana Timofeeva,<sup>2</sup> <sup>1</sup>GIT and <sup>2</sup>NMHU, DMR 0934212



Planar electron-accepting  $\pi$ -systems are useful building blocks for both nonlinear optical and organic electronic applications. We have recently developed a convenient synthesis of dibromo bithiophenes and bithiazoles of type I (see below) that relies upon the base-catalyzed "Halogen dance", in which the lithiation initially takes place adjacent to the bromine atom, before changing places with it. Compounds I can be converted to a variety of tricyclic species including the keto and diketone species shown on the right. The structures of representative intermediates and products have been confirmed by X-ray crystallography.



These species are strong electron acceptors, with dithiazole diketone (lower right) being particularly easy to reduce (−0.9 V vs. ferrocene) suggesting possible applications in  $n$ -channel organic field-effect transistors. We are currently working on incorporating these cores into more extending  $\pi$  acceptor structures.



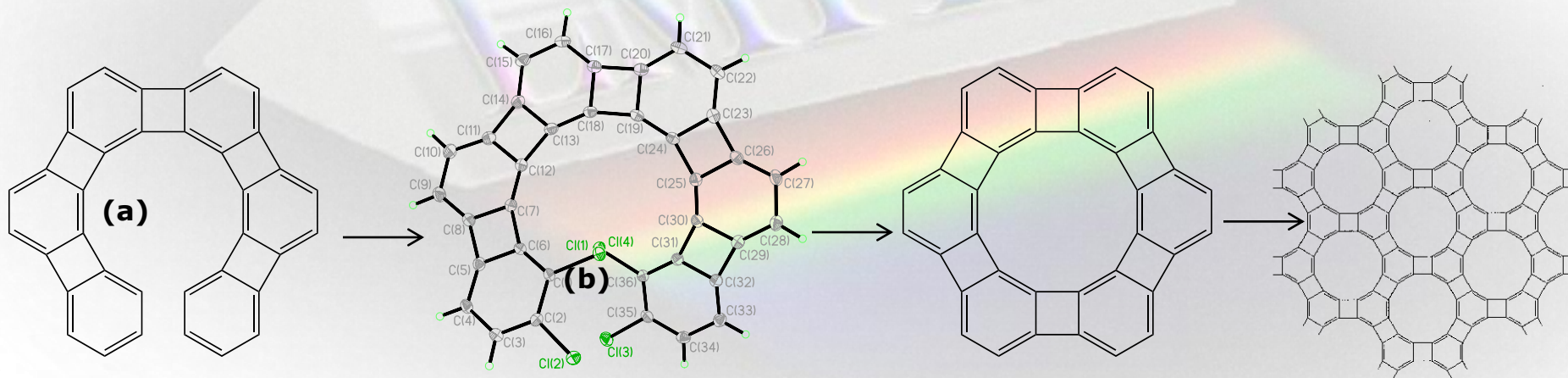


**Angular tetrachloro[6]phenylene:  
a step toward circular phenylenes**

Alexandr Fonari,<sup>1</sup> Tatiana Timofeeva,<sup>1</sup> Hao Shen<sup>2</sup> and K. Peter C. Vollhardt<sup>2</sup>

<sup>1</sup>NMHU, <sup>2</sup>Department of Chemistry, University of California at Berkeley, Berkeley.

Carbon-rich compounds and materials have garnered much attention recently due to their small band gaps and thus applications as organic conductors and ferromagnets that form monolayers in solid-phase state. One such type of materials that can form carbon monolayers are [N]phenylenes, hydrocarbons built of fused alternating aromatic benzene and anti-aromatic cyclobutadiene rings. Vollhardt's group has developed efficient synthetic routes to this family of compounds, including helical angular phenylenes.



As a precursor for circular phenylenes, PhenyleneCl<sub>4</sub> crystals were studied. Scheme above shows transformation of molecular material into monolayers with potential conducting properties.