Controlling thermal conductivity of polymers with light
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"Polymers are used extensively in engineered systems, but these materials have almost always been considered thermally static. Discovery of polymers that can be optically triggered to quickly switch between thermally conducting and insulating states will open up entirely new opportunities in thermal engineering," explained Paul Braun, a materials science and engineering (MatSE) professor and director of the Illinois Materials Research Laboratory.

"To the best of our knowledge, this is the first observation of a light-triggered reversible crystal-liquid transition in any polymeric material. The particularly notable finding in this study is the fast, reversible 3-fold change in thermal conductivity associated with the phase transition," explained Jungwoo Shin, a MatSE Ph.D. student at Illinois.

The thermal switching polymer developed by the Illinois research team demonstrates a powerful control of the thermophysical properties of a polymer in response to light. This ability originates from a photo-responsive molecule, azobenzene, which can be optically excited by ultraviolet (UV) and visible light.

"We synthesized a complex polymer functionalized with light-responsive azobenzene groups. By illuminating with UV and visible light, we could change the shape of the azobenzene group, modulate interchain bonding strength and drive a reversible transition between crystal and liquid," said Jaeuk Sung, a MatSE Ph.D. student at Illinois.

To capture the thermal conductivity transitions of azobenzene polymers under light illumination, the Illinois research team used a technique called time-domain thermoreflectance (TDTR) developed by David Cahill, a MatSE professor at Illinois.

"The way heat is carried in polymer is related to the diffusion of vibrational modes. In ordered crystals, these vibrational modes travel much further than..."
what is observed in disordered liquids. As a result, an extreme change in molecular ordering of the polymer can significantly alter the thermal conductivity,” David Cahill said.

This extreme change in macromolecular ordering, e.g., crystal-to-liquid, is rare in nature, and has not been reported previously for any polymer system in response to light. Thus, unravelling the mechanism of the light-triggered phase transition was critical to understand the polymer’s unique behavior.

"We could observe that, upon light exposure, this material quickly switches from one state to another with dramatically distinct heat transport properties. We used synchrotron-based X-ray scattering to elucidate the structure associated with each state during the transformation, closing the synthesis-characterization-function loop for this sophisticated polymer,” added Cecilia Leal, a MatSE professor at Illinois.

"Keeping an electrical device warm is as equally important as keeping it cold. Materials with such switchable thermal conductivity would enable ways to keep electrified systems safe, reliable and efficient even under extreme condition." explained Andrew Alleyne, the director of the National Science Foundation (NSF) sponsored Engineering Research Center for Power Optimization of Electro-Thermal Systems (POETS) which supported this work, and a professor in Mechanical Science and Engineering at Illinois.

"The ability to rapidly switch the thermal properties of a polymer by exposure to light opens up exciting new routes for control of thermal transport and energy conversion at the molecular level," added Nancy Sottos, a MatSE professor at Illinois.

This finding provides a striking example of how light can be used to control the thermal conductivity of polymers. A better understanding of the physical relationship between thermal conductivity and macromolecular ordering would also help push the limits of traditional polymers.


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