

New structural phase transition may broaden the applicability of photo-responsive solids

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A team of scientists from Waseda University in Tokyo and Rigaku Corporation has discovered a new type of organic crystal structural phase transition called the photo-triggered phase transition. Hideko Koshima, a visiting professor at Waseda's Research Organization for Nano & Life Innovation and leading author of the study, says, "Phase transition mechanisms are widely used in memory, switch, and actuation materials, and we believe that this discovery of a new phase transition has potential for both basic science and application fields."

Their study was published in *Communications Chemistry* on February 20, 2019.

Induced by <u>external stimuli</u> such as temperature, pressure, electromagnetic fields and <u>light</u>, a structural <u>phase</u> transition is a phenomenon that changes the physical properties and functions of solidstate materials. For instance, shape memory alloys, which have applications in robotics and in automotive, aerospace and biomedical industries, recover their shape upon heating due to martensitic transitions. In recent years, organic crystals have been considered candidates as materials for next-generation actuators because of their softness and light weight.

Prior to this study, the team reported a mechanical crystal that bends with exposure to light, as well as a robotic crystal which 'walks and rolls' when heated and cooled. The actuation of these crystals can be respectively explained by a photochromic reaction, known as



photoisomerization, and structural phase transition. To diversify the movements of such crystals, scientists have been looking for organic crystals that exhibit both phenomena.

Finding such crystals is no easy task, requiring trial and error. However, when the team was studying the photochromic chiral salicylideneamine crystal, not only did they find that it exhibits both phenomena, but also discovered the new structural phase transition. "We accidentally stumbled upon the photo-triggered phase transition of the photochromic chiral salicylideneamine crystal, which exhibits a thermal phase transition that is reversible upon heating and cooling," explains Professor Koshima. "When irradiating this crystal with ultraviolet light at -50 degrees C, a temperature lower than its thermal transition temperature (40 degrees C), we found from an X-ray crystallographic analysis that the crystal undergoes transformation identical to that of a thermal phase transition."

The team also learned that the photo-triggered phase transition occurs because of the strain of molecules produced by photoisomerization, and Koshima adds that the photo-triggered phase transition differs from a photo-induced phase transition, which has appeared in other publications. "The crystal phase due to the photo-induced phase transition appears only by light irradiation, which changes the electric and/or magnetic properties of the crystals within femto- or picoseconds. In the photo-triggered phase transition, the crystal phase triggered by light is identical to that via thermal phase transition, induced by heating, but unique with respect to its molecular conformation," she says.

Because the photo-triggered phase transition is induced by light irradiation and does not require heating and cooling for the structural phase transition to occur, the team's findings may 'lead to a new strategy to broaden applicability of photo-responsive solids' and contribute to research and development of next-generation sensing, switching,



memory, and actuators that enable remote control and/or local operation by light.

The team is now planning to measure and quantitatively evaluate the magnitude of the crystal's strain caused by photoisomerization, systematically investigate whether the photo-triggered phase transition occurs in other <u>crystals</u> using materials informatics, and clarify its conditions.

More information: Takuya Taniguchi et al, Photo-triggered phase transition of a crystal, *Communications Chemistry* (2019). DOI: 10.1038/s42004-019-0121-8

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