

Changing the handedness of molecules

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The transient chirality inversion in this study. A right-handed helical cobalt(III) complex underwent the chirality inversion to give the left-handed form before complete racemization upon removal of the chiral source. Credit: Kanazawa University

Researchers at Kanazawa University report in *Proceedings of the National Academy of Sciences* a responsive molecular system that, through chemical reactions, inverses its chirality before becoming



racemic.

Molecules that have the ability to change their structure in response to a chemical or physical stimulus are called responsive <u>molecules</u>. This type of molecule plays an important role in signal transduction at the nanoscale. The typical time profile of a structural change of a responsive molecule follows an exponential relaxation. However, <u>molecular systems</u> with non-typical time responses, such as e.g. chemical oscillators (whose structure switches periodically between two states), offer advanced functionalities and are also intensively investigated. Shigehisa Akine and colleagues from Kanazawa University have now designed a particular responsive molecule in which the chirality ('handedness') changes in a non-exponential fashion. The achievement is a breakthrough in the field of responsive systems as the chirality change happens in a unimolecular system—not, as has often been the case before, in supramolecular assemblies.

The researchers' responsive molecule has six exchangeable sites; it can be written as $[LCo_3X_6]_3^+$, where X stands for a ligand at each of the six sites. The molecule has two forms, a 'left-handed' (abbreviated 'M') and a 'right-handed' (abbreviated 'P') version. In solution, the two forms will occur, in a given ratio. Akine and colleagues started from the molecule where X is a particular chiral amine labeled A (chiral means that the ligand and its mirror image cannot be superimposed, and amine refers to a type of molecular group containing nitrogen). In a methanol solution, the P/M ratio of $[LCo_3X_6]_3^+$ was 88:12, meaning that the right-handed version was dominant. The scientists then looked at what happened when exchanging the chiral A groups with piperidine (another, but achiral, amine).

Because of the achirality of the piperidine groups, the resulting $[LCo_3(piperidine)_6]_3^+$ solution should become 'racemic', which means that any effects of chirality are compensated. This is indeed what



happened, but the researchers discovered that before reaching the racemic state after two days, the solution first switched from originally P-dominant to M-dominant after seven minutes, with maximum M-dominance after 60—120 minutes. Remarkably, a similar transient chirality inversion was not observed for the reverse reaction, $[LCo_3(piperidine)_6]_3^+$ to $[LCo_3X_6]_3^+$, for which the solution changed monotonically from racemic to P-dominant.

Akine and colleagues note that their $[LCo_3X_6]_3^+$ responsive molecule is the first unimolecular platform displaying a transient chirality inversion, and that the unique <u>chirality</u> change happens on the timescale of minutes to hours, which could be potentially useful for time-dependent functional materials related to human activity. Quoting the scientists: "this result will provide an important insight into the science of autonomously driven materials."

More information: Yoko Sakata et al, Transient chirality inversion during racemization of a helical cobalt(III) complex, *Proceedings of the National Academy of Sciences* (2022). DOI: 10.1073/pnas.2113237119

Provided by Kanazawa University

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