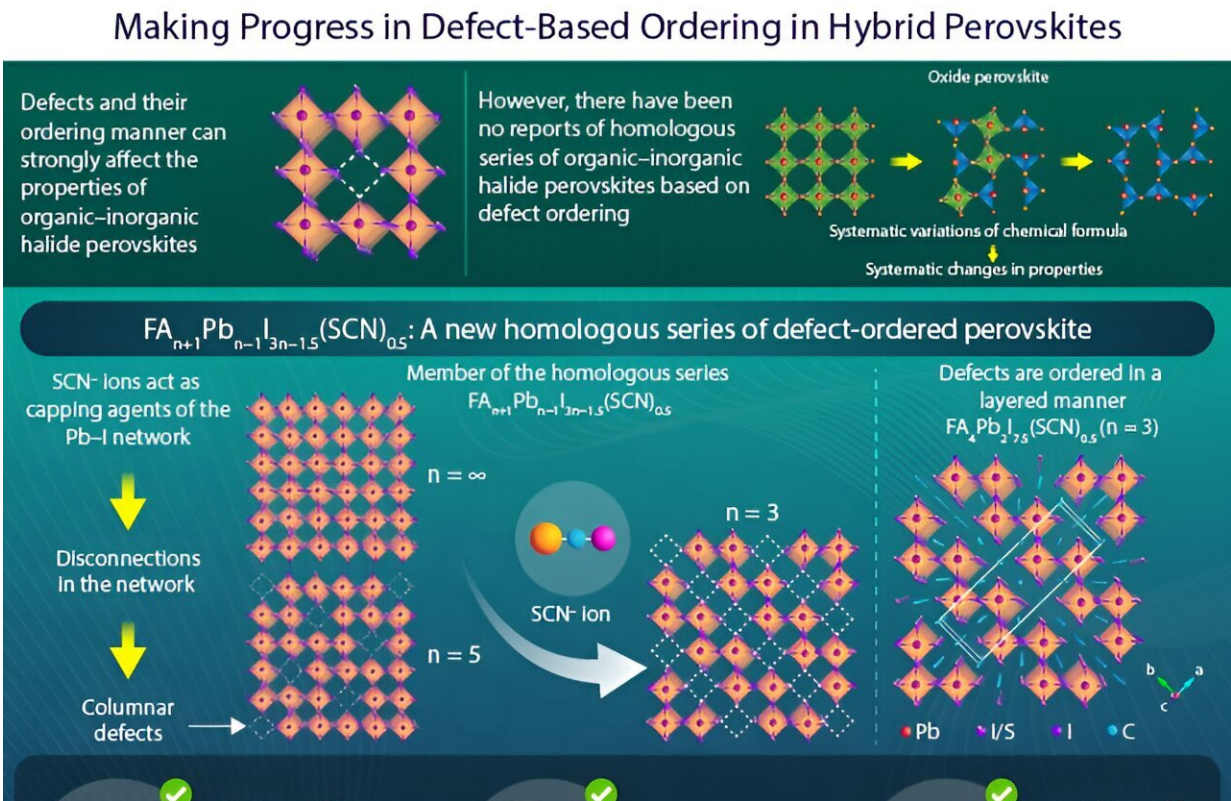


From defects to order: Spontaneously emerging crystal arrangements in perovskite halides

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Defect order engineering represents a promising strategy to control the optical properties of perovskites. Credit: Tokyo Tech

Perovskites are among the most extensively studied materials in modern

materials science. Their often unique and exotic properties, which stem from perovskite's peculiar crystal structure, could find revolutionary applications in various cutting-edge fields. One intriguing way of realizing such properties is through the precise ordering of a perovskite's defects, such as vacancies or substitutions.

In oxide chemistry, scientists have known for a long time that oxide [defects](#) can spontaneously and consistently arrange themselves throughout the crystal lattice, once they reach certain concentrations (e.g. integer ratio). This emerging order can give rise to attractive properties. While defect ordering has been observed numerous times in [perovskite oxides](#), the same cannot be said about hybrid halide perovskites, composed of an organic cation, a metal cation, and a halide anion.

In a study published in [ACS Materials Letters](#), a research team including Associate Professor Takafumi Yamamoto from Tokyo Institute of Technology discovered a new defect-ordered layered halide perovskite, shedding light on how order can emerge through defects in these compounds.

This work was inspired by a previous finding reported by the researchers, namely the formation of 'defect columns' obtained by introducing thiocyanate ion (SCN^-) into the [crystal lattice](#) of FAPbI_3 to obtain $\text{FA}_6\text{Pb}_4\text{I}_{13.5}(\text{SCN})_{0.5}$.

"We hypothesized that, if the concentration of SCN in the lattice increased, the amount of the PbI columnar defects would also increase, leading to different types of defect ordering, as seen in vacancy-ordered perovskite oxides," explains Dr. Yamamoto.

The team synthesized FAPbI_3 perovskite powders and single crystals via solid-state reactions using precisely defined concentrations of starting

materials, including specific ratios of SCN^- . They found that when an appropriately high ratio of SCN^- was used, the obtained perovskite was represented by the formula $\text{FA}_4\text{Pb}_2\text{I}_{7.5}(\text{SCN})_{0.5}$.

This layered compound, like the previously reported one, also exhibited columnar defects spanning all stacked layers. However, unlike $\text{FA}_6\text{Pb}_4\text{I}_{13.5}(\text{SCN})_{0.5}$, in which one-fifth of the PbI columns were orderly defected, one-third of all columns in the new $\text{FA}_4\text{Pb}_2\text{I}_{7.5}(\text{SCN})_{0.5}$ were defects.

The main novelty of this discovery is that the new compound, alongside the previous one, forms what's known as a "homologous series." This means that systematic variations of the compound's chemical formula, which can be represented using integer variables, result in systematic changes in its properties. In this case, the researchers found that the optical bandgap of the material increased with the concentration of ordered defects in the lattice.

Worth noting, this work presents the first homologous series based on defect ordering found for hybrid organic–inorganic perovskites. "This study provides a new playground for defect engineering in organic–inorganic hybrid perovskite compounds. We believe that this new field has the potential to develop by an analogy to the defect-ordering already seen in perovskite oxides," remarks Dr. Yamamoto.

"We have also provided a new strategy to control the defect orderings for tuning a perovskite's optical properties by incorporating SCN^- ."

The researchers hope these findings will translate to progress in an exciting area of materials science, ultimately leading to new perovskites with useful qualities for next-generation technologies.

More information: FA4Pb2I7.5(SCN)0.5: n = 3 Member of Perovskite Homologous Series FAn+1Pbn−1I3n−1.5(SCN)0.5 with Columnar Defects, *ACS Materials Letters* (2024). [DOI: 10.1021/acsmaterialslett.3c01514](https://doi.org/10.1021/acsmaterialslett.3c01514)

Provided by Tokyo Institute of Technology

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