

Relaxor ferroelectric-like relaxation based on supramolecular rotor in a ferromagnetic $[\text{MnCr}(\text{oxalate})_3]^-$ salt

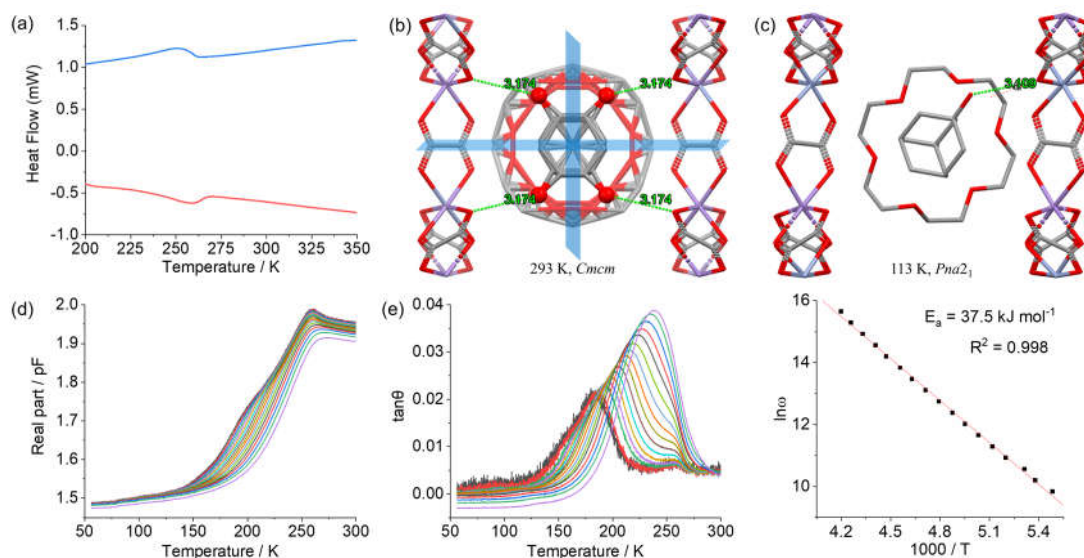
(¹Research Institute for Electronic Science, Hokkaido University, ²Graduate School of Environmental Science, Hokkaido University) ○Rui-Kang Huang,¹ Jia-Bing Wu,² Kiyonori Takahashi,¹ Kenta Kokado,¹ Takayoshi Nakamura.¹

Keywords: dielectric relaxation; supramolecular rotor; ferromagnetic; Relaxor ferroelectric; oxalate coordinates

Metal oxalates have attracted extensive attention for discovering novel multi-functional materials due to its diverse intrinsic properties.¹ Multiferroics is one of the most focusing kinds in multi-functional materials due to its wide applications in sensor, actuators, memories, *etc.*² In this study, the dielectric relaxation resulted from the frozen-rotation of the supramolecular rotor cation, $[(\text{HADA})(18\text{C6})]^+$ (HADA = 3-hydroxy-1-adamantylammonium, 18C6 = 18-crown-6), in the ferromagnetic $[\text{MnCr}(\text{oxalate})_3]^-$ anion is studied.

As shown in **Figure a**, $[(\text{HADA})(18\text{C6})][\text{MnCr}(\text{oxalate})_3]$ undergoes a phase transition at around 260 K (**LTP**&**HTP**). Single crystal X-ray diffraction show that **LTP** crystalized in polar $Pna2_1$ with ordered cation and **HTP** crystalized in nonpolar $Cmcm$ with 4-fold disordered cation. During phase transition, this material displays a dielectric relaxation in a board range with real part similar to relaxor ferroelectrics. The E_a of this relaxation calculated by imagery part is 37.5 kJ mol⁻¹, which is comparable to the E_a of hydrogen bond.

These results indicate the potential multiferroicity of this materials. Further studies is ongoing.



1. Clemente-Leon, M.; Coronado, E.; Marti-Gastaldo, C.; Romero, F. M., *Chem. Soc. Rev.* 2011, 40.
2. Fiebig, M.; Lottermoser, T.; Meier, D.; Trassin, M., *Nat. Rev. Mater.* 2016, 1.