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A-site Magnetism in Perovskites $\text{CaCu}_3\text{B}_4\text{O}_{12}$ ($B = \text{Ge}, \text{Ti}, \text{Sn}$).
TAKASHI SAITO, HIROSHI SHIRAKI, YUICHI SHIMAKAWA, Kyoto University, MASAICHIRO MIZUMAKI, Japan Synchrotron Radiation Research Institute — A-site-ordered perovskites $\text{CaCu}_3\text{Ge}_4\text{O}_{12}$ and $\text{CaCu}_3\text{Sn}_4\text{O}_{12}$, both isostructural to antiferromagnetic $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$, were found to be ferromagnets, which are very rare in cuprates. All of these materials may be called “A-site magnets”, since they contain magnetic species only at the A-site of the perovskite ABO_3 structure. The ferromagnetism of $\text{CaCu}_3\text{B}_4\text{O}_{12}$ ($B = \text{Ge}, \text{Sn}$) is attributed to the ferromagnetic direct exchange interaction, whereas antiferromagnetic superexchange interaction, due to the $\text{Cu}(3d)\text{-O}(2p)\text{-Ti}(3d)$ orbital hybridization, is dominant in antiferromagnetic $\text{CaCu}_3\text{Ti}_4\text{O}_{12}$. The A-site magnetism is controlled by the electronic structure of the non-magnetic B site. Solid solutions $\text{CaCu}_3(\text{Ge},\text{Ti})_4\text{O}_{12}$ and $\text{CaCu}_3(\text{Ti},\text{Sn})_4\text{O}_{12}$ display phase boundary between ferromagnetic and antiferromagnetic phases. [1] H. Shiraki, T. Saito, Y. Shimakawa et al., *Phys. Rev. B*, **76**, (2007) 140403. [2] Y. Shimakawa, H. Shiraki and T. Saito, *J. Phys. Soc. Jpn.*, **77**, (2008) 113702.

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