Magnetic blocking and large coercivity in lanthanide-based single-molecule magnets

Selvan Demir

Department of Chemistry, Michigan State University, East Lansing, MI 48824, USA

12 de diciembre de 2018

Salón de Actos Edificio B (Matemáticas) Facultad de Ciencias Campus San Francisco 12:00 p.m.

Molecules that possess an energy barrier to spin inversion have intriquing potential applications in areas such as magnetic refrigeration, molecular spintronics and high-density information storage. For these applications, however, key performance characteristics such as large spin-relaxation barriers and high magnetic blocking temperatures are required. Lanthanides have been proven to be particularly well-suited for the design of singlemolecule magnets owing to their large magnetic moments and magnetic anisotropy that stem from strong spinorbit coupling of the 4f orbitals. By using lanthanide ions such as Tb³+, Dy³+, and Er³+ which possess intrinsically large orbital angular momentum, significantly higher barriers and blocking temperatures can be achieved. A general methodology to enhance single-molecule magnet properties in mononuclear lanthanide complexes comprises matching the ligand field symmetry with the anisotropic electron density distribution of the maximal MJ state. Employing this methodology, we will present the synthesis of mononuclear rare-earth metallocene complexes that function as new f-metal-based single-molecule magnets. Another particularly successful approach to improve blocking temperatures is to generate strong magnetic exchange between lanthanide centers through the employment of radical bridging ligands. We will further present the synthesis of multiple bimetallic radical-bridged lanthanide-based single-molecule magnets and describe effective suppression of quantum tunneling pathways using various organic bridging radical ligands. In addition, we combine both methodologies and demonstrate with the first series of N23- radical-bridged metallocene complexes that the combination of axial magnetic anisotropy provided by the cyclopentadienyl ligands with the strong magnetic exchange coupling enabled by the inorganic N23- radical results in exceptionally large magnetic hysteresis loops which remain open up to high temperatures.





