Accepted Manuscript

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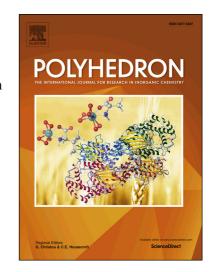
PII: S0277-5387(18)30249-3

DOI: https://doi.org/10.1016/j.poly.2018.05.011

Reference: POLY 13158

To appear in: Polyhedron

Received Date: 22 March 2018 Accepted Date: 6 May 2018



Please cite this article as: Y-Y. Nie, W-M. Wang, X-Y. Chu, H-M. Chen, S-X. Huang, Z. Liu, J-Z. Cui, H-L. Gao, Near-infrared luminescence and solvent modulation of the magnetic relaxation behavior of dinuclear lanthanide complexes, *Polyhedron* (2018), doi: https://doi.org/10.1016/j.poly.2018.05.011

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ACCEPTED MANUSCRIPT

Near-infrared luminescence and solvent modulation of the magnetic relaxation behavior of dinuclear lanthanide complexes

Yao-Yao Nie,^a Wen-Min Wang,^b Xiao-Ya Chu,^a Hong-Man Chen,^a Shao-Xia Huang,^a Zhen Liu,^a Jian-Zhong Cui*^{a,b} and Hong-Ling Gao*^{a,b}

Abstract

Seven new dinuclear centrosymmetric complexes, [Ln₂(dbm)₂L₂(DMF)₂]·2DMF (**5**), Yb (Ln = Nd (1), Eu (2),Tb Dy (4), Er (3), $[Dy_2(dbm)_2L_2(CH_3OH)_2] \cdot 2CH_3OH$ (7) (Hdbm = 1,3-diphenyl-1,3-propanedione, H₂L = 2-[(2-(amino)benzhydrazide)-methyl]-8-hydroxylquinoline, DMF = dimethylformamide) have been synthesized and structurally characterized. Dynamic magnetic studies reveal the slow magnetic relaxation of complexes 4 and 7 are influenced by a fast quantum tunneling relaxation of the magnetization (QTM). However, the different magnetic relaxation behaviors were observed apparently in the two Dy₂ complexes under a 3000 Oe dc field and modulated by the subtle change of solvents from DMF to CH₃OH. For complex 7, the temperature dependence curves of in-phase (γ') and out-of-phase (\(\chi''\)) show better frequency-dependent signals, suggesting a slow relaxation of magnetization, typical of SMM behavior. Fitting the dynamic magnetic data of complex 7 to the Arrhenius law gives the energy barrier $U_{\text{eff}}/k_{\text{B}} = 44.06 \text{ K}$ with the pre-exponential factor $\tau_0 = 2.17 \times 10^{-7}$ s under a 3000 Oe dc field.

Keywords: 8-hydroxyquinoline Schiff base derivative; dinuclear lanthanide complexes; single-molecule magnets; magnetic relaxation behavior.

1. Introduction

A great deal of single-molecule magnets (SMMs) exhibiting slow magnetic relaxation behaviors have been reported because of their huge potential applications

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