## Accepted Manuscript

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PII:	\$0277-5387(17)30605-8
DOI:	https://doi.org/10.1016/j.poly.2017.09.028
Reference:	POLY 12838
To appear in:	Polyhedron
Received Date:	22 June 2017
Revised Date:	14 September 2017
Accepted Date:	17 September 2017



Please cite this article as: Y-S. Xue, J-C. Bian, M-M. Wu, P-Y. Cheng, W-M. Wang, Z-L. Wu, M. Fang, Ligand field tuning single-molecule magnet behaviors of two dysprosium dinuclear compounds, *Polyhedron* (2017), doi: https://doi.org/10.1016/j.poly.2017.09.028

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## ACCEPTED MANUSCRIPT

### Ligand field tuning single-molecule magnet behaviors of two

#### dysprosium dinuclear compounds

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#### Abstract

In this paper, in search of an approach for tuning the single-molecule magnet behaviors in lanthanide polynuclear compounds, two new dinuclear dysprosium compounds based on an 8-hydroxyquinoline Schiff base and two different  $\beta$ -diketonate ligands,  $[Dy_2(hfac)_4L_2]$  (1) and  $[Dy_2(tfac)_4L_2]$ (2)(HL 2-[[(4-iodophenyl)imino]methyl]-8-hydroxyquinoline, hfac hexafluoroacetylacetonate and tfac = trifluoroacetylacetonate), have been synthesized, structurally and magnetically characterized. The two Dy<sub>2</sub> compounds have very similar structures, however, magnetic studies reveal that 1 and 2 showed different SMM behaviors with energy barriers of 15.70 K for 1 and 6.57 K for 2. The different SMM behaviors mainly result from the different coordination environments of the central  $Dy^{III}$  ions in 1 and 2.

*Keywords*: ligand field; dysprosium dinuclear compounds;  $\beta$ -diketonate; structures; single-molecule magnet behaviors

#### 1. Introduction

In recent years, the design and synthesis of single-molecule magnets (SMMs) have attracted significant interest in the field of molecular nanomagnets [1-3]. For such molecules magnetic materials, most of them display in-phase ( $\chi'$ ) and out-of-phase ( $\chi''$ ) ac signals and/or magnetic hysteresis-loops below the blocking temperature ( $T_B$ ) [4].

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