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Novel heterometallic polymeric lanthanide acetylacetonates with bridging cymantrenecarboxylate groups - synthesis, magnetism and thermolysis

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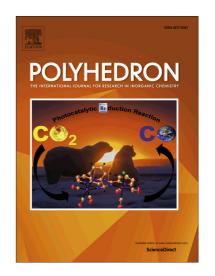
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## **ACCEPTED MANUSCRIPT**

# Novel heterometallic polymeric lanthanide acetylacetonates with bridging cymantrenecarboxylate groups – synthesis, magnetism and thermolysis

Andrey V. Gavrikov <sup>a</sup>,\*, Pavel S. Koroteev <sup>a</sup>, Zhanna V. Dobrokhotova <sup>a</sup>, Andrey B. Ilyukhin <sup>a</sup>, Nikolay N. Efimov <sup>a</sup>, Denis I. Kirdyankin <sup>a</sup>, Mikhail A. Bykov <sup>b</sup>, Mikhail A. Ryumin <sup>a</sup>, Vladimir M. Novotortsev <sup>a</sup>

New isostructural complexes  $[Ln(CymCOO)(acac)_2(H_2O)]_n$  (Ln = Eu (1), Gd (2), Tb (3), Dy (4), Ho (5), Er(6), acac – acetylacetonate (pentane-2,4dionate) anion) having a polymeric structure based on CymCOO-bridged {Ln(acac)<sub>2</sub>(H<sub>2</sub>O)} fragments were prepared by exchange reactions of hydrated Ln acetylacetonates with CymCOOH (Cym =  $(\eta^5-C_5H_4)Mn(CO)_3$ )) in a CHCl<sub>3</sub>-EtOH-H<sub>2</sub>O mixture. For the synthesized complexes comprising diamagnetic cymantrenyl moieties the temperature dependences of the dc magnetic susceptibility were studied in the temperature range of 300-2 K under an applied magnetic field of 5 kOe. Slow magnetic relaxation was found in complexes 4 and 6. The thermal decomposition of the complexes was studied by TGA and DSC in the 30 – 600°C temperature range in air and under Ar atmosphere. In the former case, the reaction affords LnMnO<sub>3</sub> phases. The first step of the thermal decomposition of 6 was studied by the kinetic analysis. For the Dy and Ho complexes, standard thermodynamic functions were calculated from adiabatic calorimetry data. Manganites LnMnO<sub>3</sub> (Ln = Eu, Gd, Tb, Dy) were synthesized from the corresponding complexes, and their magnetic properties were studied.

**Keywords**: cymantrenecarboxylate,  $\beta$ -diketonate, magnetic properties, thermal decomposition, lanthanide manganites

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