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## Dinuclear cage-core [Co<sub>2</sub>]/[Ni<sub>2</sub>] oxo-clusters supported by Sb(III) tartrate scaffolds: Synthesis, structure and magnetic properties

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### Abstract

Two dimeric metal-oxo clusters, {Na<sub>4</sub>[Co<sub>2</sub>Sb<sub>12</sub>(μ<sub>3</sub>-O)<sub>6</sub>(μ<sub>3</sub>-OH)<sub>2</sub>(μ<sub>4</sub>-O)<sub>3</sub>(tartrate)<sub>6</sub>·19.5H<sub>2</sub>O]<sub>2</sub> (**1**) and {KNa<sub>3</sub>[Ni<sub>2</sub>Sb<sub>12</sub>(μ<sub>3</sub>-O)<sub>5</sub>(μ<sub>3</sub>-OH)<sub>3</sub>(μ<sub>4</sub>-O)<sub>3</sub>(tartrate)<sub>6</sub>·20H<sub>2</sub>O]<sub>2</sub> (**2**) were obtained from a simple reaction in aqueous solution without heating. Structural analyses revealed that complexes **1** and **2** feature a cage compound {Sb<sub>12</sub>} with an encapsulated {Co<sub>2</sub>}/{Ni<sub>2</sub>} core, implying the structural diversity in the {Sb<sub>12</sub>} cage assemblies constructed from potassium antimonyl tartrate as the ligand in aqueous solution. The temperature-dependent magnetic susceptibilities indicated that there are weak ferromagnetic intermolecular or/and intramolecular interactions in **1** and strong ferromagnetic behavior in **2**, owing to the strong spin-orbital coupling (**1**) and Jahn-Teller effect (**2**).

**Keywords:** Dipotassium bis(μ-tartrato)-diantimony(III) ligand, Dinuclear cobalt(II)/nickel(II) cage-core structure, Metal-oxo clusters, Magnetic study

### 1. Introduction

Metal-oxo clusters have attracted considerable attention in recent years as a result of their intriguing architectures [1-5] as well as their fascinating properties resulting in potential applications in many fields, such as magnetism, gas adsorption and separation, catalysis and electrochemistry [6-12]. Although a diversity of self-assembly procedures have been synthesized for the construction of metal clusters by homo- and heterometallic complexes, the design and synthesis metal-oxo clusters continue to be interesting and challenging [13-16]. One feasible

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