Inorganic-organic hybrids constructed of bis(undecatungstogerma-nate)lanthanates polyoxoanions and oxalate-bridged dinuclear copper complexes and their magnetic properties

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A series of inorganic-organic hybrids $K_2Na_{m-n}[\{Ln(GeW_{11}O_{39})_2\}\{Cu_2(bpy)_2(\mu-ox)\}]\cdot nH_2O$ (bpy = 2,2-bipyridine and ox = oxalate; Ln = La, Nd, Sm, Eu, Gd; $n = 19, 17, 22, 20, 19$; $m = 4, 4, 4, 9, 2$) were isolated after reacting in a potassium acetate buffer. X-ray structural analyses show that compounds 1–5 are isomorphic and consist of $[Ln(GeW_{11}O_{39})_2]^{13-}$ polyoxoanion building blocks and oxalate-bridged dinuclear copper metalorganic complex with a 1D chain structure. The 1D chain further connects into the 3D framework by $\pi-\pi$ interactions with neighboring bpy groups. The magnetic susceptibility data indicate that antiferromagnetic coupling between the neighboring Cu$^{2+}$ ions in the structure and the rare earth ions affects magnetic property of the structure.

polyoxometalates, inorganic-organic hybrid, dinuclear copper, rare earth, magnetic property

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Recently, the successful design and synthesis of polyoxometalates(POMs)-based hybrids has attracted considerable interest because of their diverse structures and topological properties, as well as their potential applications in catalysis [1–4]. Polyoxometalates with a strong ability to accept electrons, have been extensively used as inorganic building blocks for the construction of these materials containing organic complexes. An effective approach is to covalently link transition-metal-organic ligands with POMs [5–7]. The transition-metal (TM) copper centers can have various coordination numbers and elements as well as coordination modes. They can form expansive structures by a combination of donors containing N or carboxyl groups (such as acetates and oxalates) [8,9] or by bridging Cl atoms to obtain highly nuclear magnetic clusters [10,11]. More attention has recently been given to the POM-based hybrids that have been modified by dinuclear copper complexes containing carboxyl ligands. The carboxylate bridging ligand can mediate magnetic exchange interactions between metal ions [12–16]. Many oxalate-bridged dinuclear copper hybrids have been constructed using classical polyoxoanions such as the monosubstituted Keggin type $[SiW_{12}O_{40}Cu]^+$ and the Anderson type $[M(OH)_{7}Mo_{6}O_{17}]^2-$ (M=Al, Cr) by Reinoso et al. [16] and our group [17]. However, the use of bis(undecatungstogermanate)lanthanates as inorganic building blocks has not been widely explored, possibly because they are difficult to assemble because of their large size. Only our group has reported a similar building block based on bis(undecatungstophosphate) lanthanates polyoxoanion inorganic-organic hybrids. Herein, we successfully isolated 5 inorganic-organic hybrids $K_2Na_{m-n}[\{Ln(GeW_{11}O_{39})_2\}\{Cu_2(bpy)_2(\mu-ox)\}]\cdot nH_2O$ (bpy = 2,2-bipyridine and ox=oxalate; Ln=La, Nd, Sm, Eu, Gd; $n = 19, 17, 22, 20, 19$; $m = 4, 4, 4, 9, 2$) by introducing dinuclear copper (II)-oxalate into the bis(undecatungstogermanate)lanthanates cluster. X-ray structural analyses show

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that compounds 1–5 are isomorphous and consist of [Ln(\(\text{GeW}_{11}\text{O}_{39}\))\(_2\)]\(^{13-}\) polyoxoanion building blocks and oxalate-bridged dinuclear copper metalorganic complex with a 1D chain structure. The 1D chain further connects to the 3D framework by π–π interactions of the bpy groups with neighboring chains. The magnetic susceptibility data indicate that antiferromagnetic coupling between the neighboring Cu\(^{2+}\) ions in the structure and the rare earth ions influences the magnetic property of the structure.

1 Experimental

1.1 Materials and instrumentation

The compound K\(_x\)Na\(_x\)GeW\(_{11}\text{O}_{39}\)·9H\(_2\)O was prepared as previously described [19]. All the chemicals were used as purchased without further purification. Elemental analyses (C, H, and N) were performed using a Perkin-Elmer 2400 CHN elemental analyzer; Cu, W, K, Na and Ln were determined using PLASMA-SPEC (I) ICP atomic emission spectrometer. Elemental analyses were obtained from 400–4000 cm\(^{-1}\) on an Alpha Centauri FT/IR spectrophotometer using KBr pellets. Thermal stability analyses were performed on a Perkin-Elmer TGA-7 instrument in flowing N\(_2\) at a heating rate of 10°C/min. Magnetic susceptibility data were collected from 2–300 K under a magnetic field of 1000 Oe on a Quantum Design MPMS-5 SQUID magnetometer. Diamagnetic corrections were estimated using Pascal’s constants.

1.2 Synthesis of K\(_2\)Na\(_4\)H\(_3\)[(La(\(\text{GeW}_{11}\text{O}_{39}\))\(_2\)][Cu\(_2\)(bpy)\(_2\)-(\(\mu\)-ox)]·19H\(_2\)O (1)

A cationic complex [Cu\(_2\)(2,2-bpy)\(_2\)-(\(\mu\)-ox)]\(^{2+}\) (A) and the bis(undecatungstogermanate) anion [La(\(\text{GeW}_{11}\text{O}_{39}\))\(_2\)]\(^{13-}\) (B) were prepared. For A: to an aqueous solution (20 mL) of CuCl\(_2\)·2H\(_2\)O (68 mg, 0.4 mmol) was successively added a solution of 2,2-bipyridine (62 mg, 0.4 mmol) in ethanol (10 mL) and oxalic acid (25 mg, 0.2 mmol) in water (10 mL). For B: to an aqueous solution (20 mL) of LaCl\(_3\)·7H\(_2\)O (37 mg, 0.1 mmol) was added a solution of K\(_x\)Na\(_x\)GeW\(_{11}\text{O}_{39}\)·9H\(_2\)O (610 mg, 0.2 mmol) in 1 mol/L (20 mL) potassium acetate buffer at pH 4.8. Then, solution A was added dropwise to solution B and a blue precipitate appeared. The resulting mixture was heated to 80°C for 30 min, after cooling to room temperature, and the precipitate was removed by filtration. The filtrate was allowed to stand at room temperature for crystallization. Blue block-shaped crystals of compound I were obtained after about a week. Yield: 60% (based on W). Elemental Anal. Calcd. for C\(_{22}\)N\(_4\)H\(_{65}\)Ge\(_2\)W\(_{22}\)SmCu\(_2\)Na\(_4\)K\(_2\)O\(_{104}\) (%): C, 3.95; N, 0.84; H, 0.99; W, 59.81; Eu, 2.25; Sm, 2.03; Cu, 2.12; K 1.29; Na 2.85. IR (KBr, cm\(^{-1}\)): 3423 (br), 1640 (m), 1600 (m), 1497 (m), 1472 (m), 1445 (m), 1314 (m), 1169 (m), 1100 (m), 946 (m), 872 (s), 816 (s), 755(s), 719 (m), 526 (m), 467 (m).

1.3 Synthesis of K\(_2\)Na\(_4\)H\(_3\)[(Nd(\(\text{GeW}_{11}\text{O}_{39}\))\(_2\)][Cu\(_2\)(bpy)\(_2\)-(\(\mu\)-ox)]·17H\(_2\)O (2)

The synthesis of compound 2 followed the procedure described above except that LaCl\(_3\)·7H\(_2\)O was replaced by NdCl\(_3\) (25 mg, 0.1 mmol). Yield: 61% (based on W). Elemental Anal. Calcd. for C\(_{22}\)N\(_4\)H\(_{65}\)Ge\(_2\)W\(_{22}\)NdCu\(_2\)Na\(_4\)K\(_2\)O\(_{104}\) (%): C, 4.01; N, 0.85; H, 0.84; W, 61.36; Nd, 2.19; Cu, 1.93; K 1.19; Na 1.40. Found (%): C, 4.23; N, 0.79; H, 0.95; W, 59.51; Nd, 2.35; Cu, 1.76; K 1.22; Na 1.15. IR (KBr, cm\(^{-1}\)): 3423 (br), 1642 (m), 1496 (m), 1472 (m), 1445 (m), 1314 (m), 1169 (m), 1100 (m), 946 (m), 872 (s), 816 (s), 755(s), 719 (m), 526 (m), 466 (m).

1.4 Synthesis of K\(_2\)Na\(_4\)H\(_3\)[(Sm(\(\text{GeW}_{11}\text{O}_{39}\))\(_2\)][Cu\(_2\)(bpy)\(_2\)-(\(\mu\)-ox)]·22H\(_2\)O (3)

The synthesis of 3 followed the procedure described above except that LaCl\(_3\)·7H\(_2\)O was replaced by SmCl\(_3\) (26 mg, 0.1 mmol). Yield: 55% (based on W). Elemental Anal. Calcd. for C\(_{22}\)N\(_4\)H\(_{65}\)Ge\(_2\)W\(_{22}\)SmCu\(_2\)Na\(_4\)K\(_2\)O\(_{104}\) (%): C, 3.95; N, 0.84; H, 0.98; W, 50.8 (m), 25.2; Sm, 2.03; Cu, 2.12; K 1.29; Na 1.22. IR (KBr, cm\(^{-1}\)): 3423 (br), 1640 (m), 1496 (m), 1471 (m), 1445 (m), 1312 (m), 1170 (m), 1100 (m), 944 (m), 872 (s), 818 (s), 756 (s), 728 (m), 523 (m), 467 (m).

1.5 Synthesis of K\(_2\)Na\(_4\)H\(_3\)[(Eu(\(\text{GeW}_{11}\text{O}_{39}\))\(_2\)][Cu\(_2\)(bpy)\(_2\)-(\(\mu\)-ox)]·20H\(_2\)O (4)

The synthesis of 4 followed the procedure described above except that LaCl\(_3\)·7H\(_2\)O was replaced by EuCl\(_3\) (26 mg, 0.1 mmol). Yield: 57% (based on W). Elemental Anal. Calcd. for C\(_{22}\)N\(_4\)H\(_{65}\)Ge\(_2\)W\(_{22}\)EuCu\(_2\)Na\(_4\)K\(_2\)O\(_{102}\) (%): C, 3.91; N, 0.83; H, 0.83; W, 59.81; Eu, 2.25; Cu, 1.88; K 1.16; Na 3.06. Found(%): C, 3.86; N, 0.96; H, 0.95; W, 62.23; Eu, 2.01; Cu, 1.79; K 1.03; Na 2.85. IR (KBr, cm\(^{-1}\)): 3422 (br), 1638 (m), 1600 (m), 1496 (m), 1472 (m), 1446 (m), 1313 (m), 1170 (m), 1100 (m), 944 (m), 871 (s), 818 (s), 755 (s), 728 (m), 520 (m), 467 (m).

1.6 Synthesis of K\(_2\)Na\(_2\)H\(_2\)[(Gd(\(\text{GeW}_{11}\text{O}_{39}\))\(_2\)][Cu\(_2\)(bpy)\(_2\)-(\(\mu\)-ox)]·19H\(_2\)O (5)

The synthesis of 5 followed the procedure described above except that LaCl\(_3\)·7H\(_2\)O was replaced by GdCl\(_3\) (26 mg, 0.1 mmol). Yield: 52% (based on W). Elemental Anal. Calcd. for C\(_{22}\)N\(_4\)H\(_{65}\)Ge\(_2\)W\(_{22}\)GdCu\(_2\)Na\(_2\)K\(_2\)O\(_{101}\) (%): C, 4.01; N, 0.85; H, 0.93; W, 61.32; Gd, 2.38; Cu, 1.93; K 1.19; Na 0.70.
Found (%): C, 3.94; N, 0.81; H, 0.97; W, 63.82; Gd, 2.40; Cu, 1.90; K 1.29; Na 0.53. IR (KBr, cm⁻¹): 3431 (br), 1637 (m), 1600 (m), 1496 (m), 1473 (m), 1439 (m), 1314 (m), 1113 (m), 1100 (m), 943 (m), 869 (s), 819 (s), 755 (s), 711 (m), 528 (m), 465 (m).

1.7 X-ray single crystal diffraction analysis

The diffraction intensities for compounds 2, 3 and 5 were collected on a SMART CCD diffractometer (Rigaku R-AXIS RAPID IP diffractometer for 1 and 4) equipped with a graphite monochromatic Mo-Kα radiation source (λ = 0.71073 Å) at 293 K. The structures were solved by the direct method and refined by the full-matrix least-squares method on F² using the SHELXTL crystallographic software package [20]. The hydrogen atoms were placed geometrically on the 2,2’-bipyridine ligands.

2 Results and discussion

2.1 Synthesis and structure

Compounds 1–5 were successfully isolated after the solution reaction of [Cu₂(bpy)₂(μ-ox)]²⁺ and bis(undecatungstogermanate)[Ln(GeW₁₁O₃₉)₂]¹³⁻. It is worth noting that the role of the potassium acetate buffer as an agent can be confirmed by the formation of bis(undecatungstogermanate) [Ln(Ge-W₁₁O₃₉)₂]¹³⁻ polyoxoanions. When the reaction was carried out without a potassium acetate buffer, the M/[α-GeW₁₁O₃₉] (M =Nd, Sm, Y, Yb) series of compounds formed [21]. It is obvious that the steady pH provided by the buffer solution is significant.

Single-crystal X-ray diffraction analysis revealed that compounds 1–5 are isomorphic only with slight differences in bond length, bond angle and the amount of lattice water (Table 1). Compound 1 is described as an example below.

Table 1

| Compound | M (g/mol) | Crystal system | Space group | a (Å) | b (Å) | c (Å) | α (°) | β (°) | γ (°) | V (Å³) |
|----------|---------|----------------|-------------|-------|-------|-------|-------|-------|-------|--------|
| 1        | 6585.55 | Monoclinic     | C2/c       | 15.737(3) | 19.923(4) | 37.805(8) | 90.000(0) | 101.69(3) | 90.000(0) | 11607(4) |
| 2        | 6626.92 | Monoclinic     | C2/c       | 15.920(2) | 20.038(2) | 38.106(5) | 90.000(0) | 109.00(0) | 90.000(0) | 11891(2) |
| 3        | 6687.08 | Monoclinic     | C2/c       | 15.8644(13) | 20.0581(13) | 38.060(3) | 90.000(0) | 101.981(2) | 90.000(0) | 11440.0  |
| 4        | 6762.57 | Monoclinic     | C2/c       | 15.964(3) | 19.965(4) | 37.694(8) | 90.000(0) | 101.8660(10) | 90.000(0) | 11424.0  |
| 5        | 6595.96 | Monoclinic     | C2/c       | 15.6336(13) | 20.0523(13) | 38.027(3) | 90.000(0) | 101.74(3) | 90.000(0) | 11864.0  |

Table 1 Crystal data and structure refinement parameters for compounds 1–5

| Empirical formula | C₂₂N₄H₄Ge₂W₂₂LaCu₂Na₂K₂O₉₉ | C₂₂N₄H₄Ge₂W₂₂NdCu₂Na₂K₂O₉₉ | C₂₂N₄H₄Ge₂W₂₂SmCu₂Na₂K₂O₉₉ | C₂₂N₄H₄Ge₂W₂₂EuCu₂Na₂K₂O₉₉ | C₂₂N₄H₄Ge₂W₂₂GdCu₂Na₂K₂O₉₉ |
|-------------------|------------------------------|-----------------------------|-----------------------------|-----------------------------|-----------------------------|
| M                 | 6585.55                      | 6626.92                     | 6687.08                     | 6762.57                     | 6595.96                     |
| Crystal system    | Monoclinic                   | Monoclinic                  | Monoclinic                  | Monoclinic                  | Monoclinic                  |
| Space group       | C2/c                         | C2/c                        | C2/c                        | C2/c                        | C2/c                        |
| θ range(°)        | 3.00–25.00                   | 1.09–25.00                  | 1.09–25.00                  | 1.09–25.00                  | 1.09–25.00                  |
| a (Å)             | 15.737(3)                    | 15.920(2)                   | 15.8644(13)                 | 15.964(3)                   | 15.6336(13)                 |
| b (Å)             | 19.923(4)                    | 20.038(2)                   | 20.0581(13)                 | 19.965(4)                   | 20.0523(13)                 |
| c (Å)             | 37.805(8)                    | 38.106(5)                   | 38.060(3)                   | 37.694(8)                   | 38.027(3)                   |
| α (°)             | 90.000(0)                    | 90.000(0)                   | 90.000(0)                   | 90.000(0)                   | 90.000(0)                   |
| β (°)             | 101.69(3)                    | 101.981(2)                  | 101.8660(10)                | 101.74(3)                   | 101.8660(10)                |
| γ (°)             | 90.000(0)                    | 90.000(0)                   | 90.000(0)                   | 90.000(0)                   | 90.000(0)                   |
| V (Å³)            | 11607(4)                     | 11891(2)                    | 11852.3(16)                 | 11762(4)                    | 11678.2(15)                 |
| Z                 | 4                             | 4                           | 4                           | 4                           | 4                           |
| F(000)            | 11748.0                      | 11440.0                     | 11424.0                     | 11864.0                     | 11336.0                     |
| D(ρ g cm⁻³)       | 3.838                         | 3.660                       | 3.667                       | 3.823                       | 3.698                       |
| Abs coeff (mm⁻¹)  | 23.130                       | 22.644                      | 22.769                      | 23.010                      | 23.171                      |
| Total data collected | 44497                     | 29976                       | 29939                       | 44165                       | 29419                       |
| Unique data       | 10227                        | 10463                       | 10435                       | 10364                       | 10283                       |
| Rm                | 0.1914                       | 0.076                       | 0.0682                      | 0.1226                      | 0.0703                      |
| GOF               | 1.027                        | 1.043                       | 1.052                       | 1.063                       | 1.057                       |
| R(I > 2σ(I))      | 0.0574                       | 0.0537                      | 0.0518                      | 0.0562                      | 0.0543                      |
| wR(Cl all data)   | 0.1253                       | 0.1434                      | 0.1384                      | 0.0922                      | 0.1444                      |

a) R₁ = Σ||Fo|−|Fc||/Σ|Fo|; b) wR₂ = [Σw(Fo²−Fc²)²/Σw(Fo²)]¹².
Table 2  Ln–O bond lengths in compounds 1–5

| Compound | Ln–O bond length |
|----------|-----------------|
| 1 (La<sup>3+</sup>) | 2.526–2.552 |
| 2 (Nd<sup>3+</sup>) | 2.440–2.507 |
| 3 (Sm<sup>3+</sup>) | 2.403–2.479 |
| 4 (Eu<sup>3+</sup>) | 2.398–2.467 |
| 5 (Gd<sup>3+</sup>) | 2.394–2.464 |

connected by two oxalate oxygen atoms and two bipyridine nitrogen atoms at the equatorial positions and two terminal oxygen atoms from the [Ln(GeW<sub>11</sub>O<sub>39</sub>)<sub>2</sub>]<sup>13−</sup> polyoxoanions at the axial positions with a Cu–Cu distance close to the average of 5.2 Å. The connection between [Cu<sub>2</sub>(bpy)<sub>2</sub>(μ-ox)]<sup>2+</sup> and La(GeW<sub>11</sub>O<sub>39</sub>)<sub>2</sub> gives a 1D chain (Figure 1). Adjacent bipyridine layers further result in a 3D structure by π–π stacking interactions (distance between the bpy layers are approximately 3.7 Å) (Figure 2).

### 2.2 IR spectroscopy

The IR spectra of compounds 1–5 are similar, as shown in Figure 3, there are 4 characteristic asymmetric vibrations that result from the [Ln(GeW<sub>11</sub>O<sub>39</sub>)<sub>2</sub>]<sup>13−</sup> polyoxoanions, namely, mas(W–O<sub>a</sub>), mas(W–O<sub>b</sub>), mas(W–O<sub>c</sub>) and mas(Ge–O<sub>a</sub>) that appear at 943, 872, 755 and 816 cm<sup>−1</sup>, respectively. By comparing the IR spectra of the polyoxoanions with those of α-H<sub>2</sub>GeW<sub>12</sub>O<sub>40</sub> [23], shifts and splits are present because of the lower symmetry. The characteristic peaks that are present between 1100 and 1600 cm<sup>−1</sup> are attributed to the 2,2′-bipyridine ligand, and these are of low intensity with respect to those of the polyoxoanions. The features at about 755 cm<sup>−1</sup> are assigned to ν<sub>as</sub>(CO) and those at 1315 and 1360 cm<sup>−1</sup> are assigned to ν<sub>s</sub>(CO) while the broad absorption at 1642 cm<sup>−1</sup> corresponds to ν(OCO) of the oxalate ligand in bisbidentate bridging mode [24].

### 2.3 Thermogravimetric analyses

For compounds 1–5, the amount of lattice water was determined by thermal gravimetric analysis. The curve of compound 1 gives a total weight loss of 11.69% between 25–750°C, which agrees with the calculated value of 11.85%. The weight loss of 5.17% at 25–260°C corresponds to the loss of all non-coordinated and coordinated water molecules (calcd. 5.19%). The weight loss of 6.52% at 260–750°C arises from the decomposition of organic molecules (calcd. 6.66%). Compounds 2–5 also undergo a two-step weight loss.

### 2.4 Magnetic properties

Oxalate dinuclear copper compounds are known to exhibit strong antiferromagnetic interactions [25]. When rare earth ions are added to the dinuclear compounds, the crystal...
structure becomes complex as the two copper (II) ions and the free Ln(III) make the analysis of the observed magnetic behavior difficult. For example, despite the presence of strong antiferromagnetic interactions, no relative maximum is observed on the susceptibility versus temperature curve. It is possible that the expected maximum is outside the measured temperature range but it is also possible that the maximum is enveloped by the contributions of the Ln(III) ions. According to the change in the rare earth ion f-orbital electronic number from zero to partially full to half full, the thermal evolution of the magnetic molar susceptibility and the $\chi_m T$ product of 1, 2 and 5 in the temperature range 2–300 K are shown selectively in Figure 4. For compound 1, from 300 to 2 K, the $\chi_m$ value increases gradually with a decrease in temperature and no maximum is observed. The $\chi_m T$ value is equal to $0.609 \text{ cm}^3 \text{ K mol}^{-1}$ at 300 K, which is lower than the expected value of $0.750 \text{ cm}^3 \text{ K mol}^{-1}$ for the two $S = 1/2$ Cu$^{2+}$ centers with $\chi_m T = \mu_{\text{eff}}^2/8$ (considering $g = 2$). Upon cooling the samples, the product $\chi_m T$ decreases from 300 to 50 K, and $\chi_m T$ remains constant around $0.0869 \text{ cm}^3 \text{ K mol}^{-1}$ when the temperature is lower than 50 K and this may be attributed to a paramagnetic impurity. The decrease in $\chi_m T$ indicates the antiferromagnetic coupling of the bridged-oxalate dinuclear copper; and the rare earth ion (La$^{3+}$, 4f$^0$) does not contribute to the magnetic property of the whole structure. Thus the oxalate-bridged dinuclear copper complex has intrinsic magnetism in the inorganic-organic hybrids. For compound 2, the $\chi_m T$ curve is different to that of the former because of the existence of paramagnetic Nd$^{3+}$ ions. This fact indicates the predominance of antiferromagnetic interactions in compound 2, as confirmed by the rapid decrease in the effective magnetic moment with a decrease in temperature from 300 to 2 K. Moreover, the exchange coupling must certainly be strong considering that Curie-Weiss behavior is not observed even at a high temperature on the reciprocal susceptibility curve. The $\chi_m T$ value at room temperature is equal to $1.789 \text{ cm}^3 \text{ K mol}^{-1}$, which is lower than the expected value of $2.386 \text{ cm}^3 \text{ K mol}^{-1}$ for two $S = 1/2$ Cu$^{2+}$ centers with $\chi_m T = \mu_{\text{eff}}^2/8$ (considering $g = 2$) and one free Nd$^{3+}$ ion in the $^4I_{9/2}$ ground state ($g' = 8/11$). Upon cooling the samples, the product $\chi_m T$ decreases very sharply and reaches a minimum ($0.475 \text{ cm}^3 \text{ K mol}^{-1}$) at 2 K, and this may be attributed to crystal field effects because of the intrinsic nature of the lanthanide series [26]. For compound 5, the magnetic coupling is dependent on the exchange transfer integral between the 3d orbital of Cu (II) and the 4f orbital of Gd (III); however, the long Cu–Gd distance (10.527 Å) may affect the magnetic interactions in the whole structure. According to the Curie-Weiss law, there is a dominant antiferromagnetic coupling in oxalate-bridging Cu(II) dimers. Similar magnetic curves were found for compounds 2 and 5. For compound 5, the $\chi_m T$ curve is different to that of the former because of the existence of half-full Gd$^{3+}$ (4f$^7$) ions. The relationship between $1/\chi_m$ and T from 2–300 K fits the Curie-Weiss law.
The dominant antiferromagnetic coupling exists in the structure. Earth ions play a role in the magnetic character of this from 300 K to 2 K in a near straight line because the rare earth ions affects strong antiferromagnetic coupling in the dinuclear copper oxalate-bridged dinuclear copper complexes by conventional methods. Magnetic susceptibility studies reveal a strong antiferromagnetic coupling in the dinuclear copper complexes and the choice of different rare earth ions affects the magnetic property of the structure.

3 Conclusions

In this paper, we synthesized five novel inorganic-organic hybrid compounds using bis(undecatungstogermanate) and oxalate-bridged dinuclear copper complexes by conventional methods. Magnetic susceptibility studies reveal a strong antiferromagnetic coupling in the dinuclear copper complexes and the choice of different rare earth ions affects the magnetic property of the structure.

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