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Synthesis and Coordination Behavior of a Bipyridine Platinum(II) Complex with Thioglucose

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ABSTRACT

A mononuclear platinum(II) complex with two monodentate-\(S\ H_4tg^-\) ligands, [Pt(\(H_4tg-\kappa S\))\(_2\)(bpy)] (1), was newly synthesized by the reaction of [PtCl\(_2\)(bpy)] (bpy = 2,2’-bipyridyl) with NaH\(_4tg\) (NaH\(_4tg\) = 1-thio-\(\beta\)-D-glucose sodium salt) in water. Complex 1 reacted with additional [PtCl\(_2\)(bpy)] in water to give an S-bridged dinuclear complex, [Pt\(_2(\mu_2-\mu_2\ H_4tg-\kappa 1\ S;\kappa 1\ S)_2(bpy)_2\)]\(^{2+}\) ([2]\(^{2+}\)), in which a square-planar [Pt(\(H_4tg\)\(_2\)(bpy))] unit binds to a [Pt(bpy)]\(^{2+}\) moiety through two thiolato groups. Treatments of 1 with Cu\(^{2+}\) and Ni\(^{2+}\) in water in the presence of bpy produced S-bridged dinuclear complexes, [PtCu(\(\mu_2-\mu_2\ H_4tg-\kappa 1\ S;\kappa 2\ O,S)_2(bpy)_2\)]\(^{2+}\) ([3]\(^{2+}\)) and [PtNi(\(\mu_2-\mu_2\ H_4tg-\kappa 1\ S;\kappa 2\ O,S)_2(bpy)_2\)]\(^{2+}\) ([4]\(^{2+}\)), respectively, in which a square-planar [Pt(\(H_4tg\)\(_2\)(bpy))] unit binds to a [M(bpy)]\(^{2+}\) (M = Cu\(^{II}\), Ni\(^{II}\)) moiety through two thiolato and two hydroxyl groups to form a chiral [M(\(N\)\(_2\)(O)\(_2\)(S)\(_2\))] octahedron with the \(\Delta\) configuration. On the other hand, similar treatment with Cd\(^{2+}\) in the presence of bpy resulted in the formation of an S-bridged trinuclear complex, [Cd{Pt(\(\mu_2-\mu_2\ H_4tg-\kappa 1\ S;\kappa 2\ O,S)(\mu_2-\mu_2\ H_4tg-\kappa 1\ S;\kappa 1\ S)_2}\(bpy\))\_2\}]\(^{2+}\) ([5]\(^{2+}\)), in which each of two square-planar [Pt(\(H_4tg\)\(_2\)(bpy))] units binds to a Cd\(^{II}\) ion through two thiolato and one hydroxyl groups to form a chiral [Cd(O)\(_2\)(S)\(_4\)] octahedron with the \(\Lambda\) configuration. Of two geometrical configurations, syn and anti, which arise from the relative arrangement of two \(\beta\)-D-pyranose moieties, [2]\(^{2+}\) adopts the syn configuration with symmetric bridging sulfur atoms, while each of [3]\(^{2+}\), [4]\(^{2+}\), and [5]\(^{2+}\) has the anti configuration with \(R\) configurational bridging sulfur atoms. All the complexes were fully characterized by electronic absorption, CD, and NMR spectroscopies, along with single-crystal X-ray crystallography.
INTRODUCTION

1-Thio-β-D-glucose (H₅tg = C₆H₁₂O₅S) is one of the simplest thiol-containing hexoses.¹ It has been used as starting material for a variety of synthetic S-glycosides in organic chemistry and as a coating reagent for gold nanoparticles and electrodes in material science.¹⁻³ In coordination chemistry, the development of transition-metal complexes with H₅tg was initially motivated by the finding that some Au¹-thiolate compounds show an anti-rheumatoid activity.⁴ During the late 1920’s, Landé and Pick independently reported that Aurothioglucose ([Au(H₄tg-S)]ₙ) has an effect on rheumatic disease.⁵a,b The subsequent studies established an excellent anti-rheumatoid activity for this complex, which has been attributed to its hydrophilicity and biocompatibility of the sugar backbone.⁴,⁵ The investigation of this class of complexes was extended to include a phosphine compound as a co-ligand, and a number of related Au¹ complexes such as [Au(H₄tg-S)(PET₃)], [Au₂(H₄tg-S)₂(dppe)] (dppe = 1,2-bis(diphenylphosphino)ethane), and [Au(Ac₄tg-S)(PET₃)] (auranofin) were prepared and their anti-rheumatoid activities were studied.⁴,⁶ On the other hand, reports on the synthesis of coordination compounds from H₅tg in combination with a transition metal ion other than Au¹ are relatively limited,⁷⁻¹⁰ despite the presence of both soft thiol and hard hydroxyl groups in H₅tg available for coordination to a variety of metal centers. In particular, only a few transition metal complexes that are fully characterized have been presented to date,⁸ and to the best of our knowledge, [Co(H₃tg-S,O)(en)₂]⁺ (en = ethylenediamine) and [ReO(SSS)(H₄tg-S)] (SSS = 2,2'-thiodiethanethiolate) are the only transition metal complexes that have been structurally characterized by single-crystal X-ray crystallography.⁹,¹⁰ This is most likely because of the significant hydrophilicity of glycosyl moiety in H₅tg, which makes it difficult to isolate target species in a pure form from a reaction mixture in solution.¹¹
As part of our continuing interest in chiral coordination compounds having thiolato donor groups, which potentially function as an S-donating chiral metalloligand,\textsuperscript{12-16} we started to investigate the rational preparation of an inert thiolato platinum(II) system by using H$_5$tg as a chiral ligand. Our strategy is the introduction of a hydrophobic diimine-type co-ligand in a platinum(II) coordination system, which should compensate the hydrophilicity of glycosyl moieties to facilitate the isolation and crystallization of a platinum(II) complex from a polar reaction medium that is required to dissolve H$_5$tg. In this study, we employed 2,2’-bipyridyl (bpy) as a diimine-type co-ligand. This ligand is expected to show intermolecular π-π interactions that also contribute to decrease the solubility of a platinum(II) complex in a polar solvent.\textsuperscript{17} Indeed, we were able to isolate a new transition metal complex with two H$_4$tg$^-$ ligands, [Pt(H$_4$tg-S)$_2$(bpy)] ([1]), by a simple 1:2 reaction of [PtCl$_2$(bpy)] and H$_4$tg$^-$ in water (Scheme 1). Furthermore, [1] was found to react with a second transition metal ion (M = Pt$^{II}$, Cu$^{II}$, Ni$^{II}$, Cd$^{II}$), giving S-bridged dinuclear or trinuclear complexes with retention of the mononuclear structure in [1] (Scheme 2). In this paper, we report on the synthesis and characterization of these new coordination compounds containing H$_4$tg$^-$ ligand. Notably, all the four dinuclear or trinuclear complexes obtained were structurally characterized by single-crystal X-ray diffraction. This study demonstrates the functionality of [1] as a versatile chiral metalloligand available for the chiralselective construction of S-bridged multinuclear structures, as well as the effectiveness of the introduction of a diimine-type co-ligand to isolate coordination compounds with hydrophilic glycosyl moieties.


EXPERIMENTAL SECTION

Preparation of [Pt(H$_4$tg-κ$S$)$_2$(bpy)] (1). To a yellow suspension of [PtCl$_2$(bpy)]$^{18}$ (0.50 g, 1.18 mmol) in water (300 mL) was added a solution of NaH$_4$tg (0.59 g, 2.70 mmol) in water (20 mL). The mixture was stirred at 60°C for 1 h to give an orange solution. After a small amount of unreacted [PtCl$_2$(bpy)] was filtered off, the filtrate was evaporated to dryness using a rotary evaporator. The orange residue was dissolved in 20 mL of water, followed by allowing it to stand at room temperature overnight. The resulting orange fiber product was collected by filtration.

Yield: 0.75 g (81%). Anal. Calcd for [Pt(H$_4$tg-κ$S$)$_2$(bpy)]$\cdot$2H$_2$O ([1]$\cdot$2H$_2$O = C$_{22}$H$_{34}$N$_2$O$_{12}$PtS$_2$): C, 33.98; H, 4.41; N, 3.60%. Found: C, 33.82; H, 4.26; N, 3.59%. $^1$H NMR (D$_2$O): $\delta$ 9.54 (2H, d, $J = 4.9$ Hz, bpy), 8.29 (2H, d, $J = 7.6$ Hz, bpy), 8.22 (2H, t, $J = 7.8$ Hz, bpy), 7.66 (2H, t, 6.7 Hz, bpy),...
Preparation of [Pt₂(μ₂-H₄tg-κ¹S:κ¹S)$_2$(bpy)$_2$](ClO₄)$_2$ ([2](ClO₄)$_2$).

Method A. To an orange suspension of [1]·2H₂O (0.05 g, 0.06 mmol) in water (50 mL) was added a solid sample of [PtCl₂(bpy)] (0.03 g, 0.07 mmol). The mixture was stirred at 50°C for 2.5 h to give a yellow-brown solution. After a small amount of unreacted [PtCl₂(bpy)] was filtered off, the filtrate was concentrated to dryness using a rotary evaporator. The yellow green residue was dissolved in 10 mL of water, and an aqueous NaClO₄ solution (0.5 mL, 5 M) was added to it. The solution was allowed to stand at room temperature for 10 days. The resulting yellow block crystals were collected by filtration. Yield: 0.06 g (64%). One of the crystals thus obtained was used for X-ray analysis. Anal. Caled for [Pt₂(μ₂-H₄tg-κ¹S:κ¹S)$_2$(bpy)$_2$](ClO₄)$_2$·9H₂O ([2](ClO₄)$_2$·9H₂O = C₃₂H₅₆N₄Cl₂O₂₇Pt₂S₂): C, 26.43; H, 3.88; N, 3.85%. Found: C, 26.37; H, 3.69; N, 3.89%. $^1$H NMR (D₂O): δ 8.92 (d, 2H, J = 5.65 Hz, bpy), 8.77 (d, 2H, J = 5.34 Hz, bpy), 8.23 (t, 4H, J = 7.55 Hz, bpy), 8.10 (d, 4H, J = 8.09 Hz, bpy), 7.80 (t, 2H, J = 6.48 Hz, bpy). $^{13}$C{¹H} NMR (D₂O): δ 157.5 (bpy), 157.4 (bpy), 151.0 (bpy), 150.2 (bpy), 144.8 (bpy), 144.7 (bpy), 130.6 (bpy), 130.5 (bpy), 126.7 (bpy), 126.5 (bpy), 86.56 (H₄tg⁻), 83.07 (H₄tg⁻), 79.61 (H₄tg⁻), 76.81 (H₄tg⁻), 71.30 (H₄tg⁻), 62.76 (H₄tg⁻). ESI-MS (in MeOH) m/z: 546.1 ([M]$^{2+}$).

Method B. To a yellow suspension of [PtCl₂(bpy)] (0.10 g, 0.24 mmol) in water (60 mL) was added a solution of NaH₄tg (0.06 g, 0.27 mmol) in water (5 mL). The mixture was stirred at 50°C for 3 h to give a yellow solution. After a small amount of unreacted [PtCl₂(bpy)] was filtered off, the filtrate was concentrated to ca. 20 mL using a rotary evaporator. An aqueous NaClO₄ solution (0.5 mL, 5 M) was added to the yellow concentrated solution, followed by allowing it to
stand at room temperature for 5 days. The resulting yellow block crystals of [2](ClO₄)₂·9H₂O were collected by filtration. Yield: 0.12 g (70%).

**Preparation of** [PtCu(μ₂-H₄tg-κ¹S₂κ²O₅S₂)(bpy)₂](NO₃)₂ ([(3)(NO₃)₂]. To a suspension of [1]·2H₂O (0.05 g, 0.06 mmol) in water (10 mL) was added a solution of [CuCl₂(bpy)]¹⁹ (0.02 g, 0.07 mmol) in water (5 mL). The mixture was stirred at room temperature for 1 h, and an aqueous NaNO₃ solution (1 mL, 5 M) was added to it. The solution was allowed to stand at room temperature for 2 days to give green block crystals, which were collected by filtration and washed with a small amount of water. Yield: 0.04 g (55%, based on Pt). Anal. Calcd for [PtCu(μ₂-H₄tg-κ¹S₂κ²O₅S₂)(bpy)₂](NO₃)₂: C, 34.27; H, 3.77; N, 7.49%. Found: C, 34.29; H, 3.69; N, 7.50%.

Single crystals of [(3)(ClO₄)₂·2H₂O used for X-ray analysis were obtained by the following method: To a suspension of [PtCl₂(bpy)] (0.08 g, 0.19 mmol) in water (10 mL) was added a solution of NaH₄tg (0.09 g, 0.41 mmol) in water (2 mL). The mixture was stirred at room temperature for 4 h to give an orange suspension, to which a solution of [CuCl₂(bpy)] (0.06 g, 0.21 mmol) in 5 mL of water was added. The mixture was stirred at room temperature for 2 h to give a green suspension. After insoluble materials were filtered off, to the green filtrate was added a small amount of saturated NaClO₄ aqueous solution. The solution was allowed to stand at room temperature for 3 days to give green block crystals of [(3)(ClO₄)₂·2H₂O.

**Preparation of** [PtNi(μ₂-H₄tg-κ¹S₂κ²O₅S₂)(bpy)₂](NO₃)₂ ([(4)(NO₃)₂]. To a suspension of [1]·2H₂O (0.05 g, 0.06 mmol) in water (10 mL) was added a solution of Ni(NO₃)₂·6H₂O (0.02 g, 0.07 mmol) in water (2 mL). The mixture was stirred at 40ºC for 1 h to give a yellowish brown solution. A solid sample of bpy (0.01 g, 0.06 mmol) was added to it, and the mixture was stirred at 40ºC for 2 h to give a yellow-brown solution. To this reaction solution was added an aqueous
NaNO₃ solution (1 mL, 5 M), followed by allowing it to stand at room temperature for 12 days. The resulting pale green crystals were collected by filtration and washed with a small amount of water. One of the crystals thus obtained was used for X-ray analysis. Yield: 0.04 g (57%, based on Pt). Anal. Calcd for [PtNi(μ₂-H₄tg-κ¹S:\kappa²O,S)₂(bpy)₂](NO₃)₂·H₂O ([4](NO₃)₂·H₂O = C₃₂H₄₀N₆NiO₁₇PtS₂): C, 34.99; H, 3.67; N, 7.65%. Found: C, 34.74; H, 3.68; N, 7.69%.

Preparation of [Cd{Pt(μ₂-H₄tg-κ¹S:\kappa²O,S)(μ₂-H₄tg-κ¹S:\kappa¹S)(bpy)}₂](ClO₄)₂ ([5](ClO₄)₂).

Method A. To an orange suspension of [1]·2H₂O (0.025 g, 0.03 mmol) in water (5 mL) was added a solution of CdCl₂·2.5H₂O (0.008 g, 0.04 mmol) in water (1 mL). The mixture was stirred at 40°C for 2 h to give a yellow solution. A solid sample of bpy (0.005 g, 0.03 mmol) was added to it, and the mixture was stirred at 40°C for 1 h to give a yellow suspension. After a small amount of colorless solid was filtered off, to the yellow filtrate was added an aqueous NaClO₄ solution (10 drops, 1 M). The solution was allowed to stand at room temperature for 5 days. The resulting yellow block crystals were collected by filtration and washed with a small amount of water. Yield: 0.005 g (16%, based on Pt). Anal. Calcd for [Cd{Pt(μ₂-H₄tg-κ¹S:\kappa²O,S)(μ₂-H₄tg-κ¹S:\kappa¹S)(bpy)}₂](ClO₄)₂·6H₂O ([5](ClO₄)₂·6H₂O = C₄₄H₇₂N₄CdCl₂O₃₄Pt₂S₄): C, 27.77; H, 3.81; N, 2.94%. Found: C, 27.88; H, 3.70; N, 2.95%.

Method B. To a suspension of [1]·2H₂O (0.05 g, 0.06 mmol) in water (10 mL) was added a solution of CdBr₂ (0.009 g, 0.03 mmol) in water (2 mL). The mixture was stirred at room temperature for 15 min to give a yellow solution. To this solution was added an aqueous NaClO₄ solution (8 drops, 5 M), followed by allowing it to stand at room temperature for 6 days. The resulting yellow block crystals of [5](ClO₄)₂·6H₂O were collected by filtration. One of the crystals thus obtained was used for X-ray analysis. Yield: 0.044 g (72%, based on Pt).
Caution! Perchlorate salts of metal complexes are potentially explosive and should be handled with great care.

**Measurements.** The electronic absorption spectra were recorded with JASCO Ubest-55, V-570, or V-660 spectrophotometer, and the CD spectra were recorded with JASCO J-820, J-700, or J-600 spectrophotometer at room temperature. The diffuse reflection spectra in the solid state were recorded on a JASCO V-570 spectrophotometer using MgSO₄. The IR spectra were measured with a JASCO FT/IR-550 infrared spectrophotometer using KBr disks. The $^1$H and $^{13}$C{1H} NMR spectra were recorded with a JEOL EX-270 or a GSX-500 spectrometer at the probe temperature in D₂O. Sodium 4,4-dimethyl-4-silapentane-1-sulfonate (DSS) was used as the internal reference. Elemental analyses (C, H, N) were performed with Yanaco CHN Corder MT-5. The X-ray fluorescence analyses were made on a HORIBA MESA-500 spectrometer. The molar conductivities were measured with a HORIBA DS12 conductivity meter. Electrospray-ionization (ESI) mass spectra were recorded in a positive-ion mode on ABsciex QSTAR Elite controlled by Analyst QS 2.0 software package. The data of electronic absorption, CD, IR, and $^1$H NMR spectra together with molar conductivity are summarized in Tables S6 and S7.

**X-ray Structure Determinations.** Single-crystal X-ray diffraction measurements for [2](ClO₄)₂·9H₂O, [4](NO₃)₂·2H₂O and [5](ClO₄)₂·6H₂O were made on a Rigaku RAXIS RAPID imaging plate diffractometer with a graphite monochromated Mo-Kα radiation ($\lambda = 0.7107$ Å) at 293 K, 200 K and 200 K, respectively. Unit cell parameters were determined by a least-squares refinement. The intensity data were collected by the $\omega$ scan mode up to $2\theta_{\text{max}} = 54.94^\circ$ for [2](ClO₄)₂·9H₂O, to $2\theta_{\text{max}} = 54.98^\circ$ for [4](NO₃)₂·2H₂O and to $2\theta_{\text{max}} = 54.96^\circ$ for [5](ClO₄)₂·6H₂O. Empirical absorption corrections were also applied. The 10740, 8915 and 7149
independent reflections with \( I > 2\sigma(I) \) of the measured 23133, 29367 and 40029 reflections were considered as “observed” and used for the structure determinations of (2)(ClO\(_4\))\(_2\)\( \cdot \)9H\(_2\)O, (4)(NO\(_3\))\(_2\)\( \cdot \)2H\(_2\)O and (5)(ClO\(_4\))\(_2\)\( \cdot \)6H\(_2\)O, respectively. The structures were solved by the direct method with SHELXS-97 and expanded using Fourier techniques.\(^{20}\) The non-hydrogen atoms except oxygen atoms of water and ClO\(_4\)\(^-\) molecules were refined anisotropically by full-matrix least-squares techniques on \( F^2 \) using SHELXL-97.\(^{20}\) The asymmetric unit contained a half of complex-cation for (5)(ClO\(_4\))\(_2\)\( \cdot \)6H\(_2\)O. In (2)(ClO\(_4\))\(_2\)\( \cdot \)9H\(_2\)O and (5)(ClO\(_4\))\(_2\)\( \cdot \)6H\(_2\)O, one of two 1-thio-\( \beta \)-D-glucose moieties was disordered over two positions ((2)(ClO\(_4\))\(_2\)\( \cdot \)9H\(_2\)O: C32A-O10A and C32B-O10B, (5)(ClO\(_4\))\(_2\)\( \cdot \)6H\(_2\)O: C12A-O10A and C12B-O10B). Several solvated water molecules in (5)(ClO\(_4\))\(_2\)\( \cdot \)6H\(_2\)O were disordered.

A single-crystal X-ray diffraction measurement for (3)(ClO\(_4\))\(_2\)\( \cdot \)2H\(_2\)O was made on a Rigaku AFC-7R four-cycle diffractometer with a graphite monochromated Mo-K\( \alpha \) radiation at room temperature. The intensity data were collected by the \( \omega \)-2\( \theta \) scan mode up to 2\( \theta \)\( _{\text{max}} \) = 55.08\(^\circ\). The intensities were corrected for Lorentz and polarization effects. Empirical absorption correction based on \( \Psi \) scans was also applied. The 5593 independent reflections with \( I > 2\sigma(I) \) of the measured 5593 reflections were considered as “observed” and used for the structure determination. The structure was solved by the direct method with SHELXS-97 and expanded using Fourier techniques.\(^{20}\) The non-hydrogen atoms except oxygen and chlorine atoms of water and ClO\(_4\)\(^-\) molecules were refined anisotropically by full-matrix least-squares techniques on \( F^2 \) using SHELXL-97.\(^{20}\) Several solvated water molecules in (3)(ClO\(_4\))\(_2\)\( \cdot \)2H\(_2\)O were disordered.

All calculations were performed using the Crystal Structure crystallographic software package and SHELXL-97 program. Hydrogen atoms except those of water molecules were located at calculated positions and refined as riding models.
The selected bond distances and angles of [2](ClO₄)₂·9H₂O, [3](ClO₄)₂·2H₂O, [4](NO₃)₂·2H₂O, and [5](ClO₄)₂·6H₂O, as well as their crystallographic data, are summarized in Supporting Information (Tables S1-S5).

RESULTS AND DISCUSSION

Synthesis and Characterization of [Pt(H₄tg-κS)₂(bpy)] ([1]). The reaction of [PtCl₂(bpy)] with NaH₄tg in a 1:2 ratio in water gave an orange solution, from which an orange fiber product ([1]) was isolated in a good yield. The elemental analytical data of [1] were consistent with the formula for [Pt(H₄tg)(bpy)], and its neutral charge was implied by the molar conductivity in water (3.7 S cm² mol⁻¹). The presence of H₄tg⁻ and bpy ligands in [1] in a 2:1 ratio was indicated by its ¹H NMR spectrum in D₂O (Figure S1), which shows aliphatic proton signals (δ 3.3-3.8) and aromatic proton signals (δ 7.6-9.6) in an integration ratio of 7:4. The electronic absorption spectrum of [1] in water shows an intense visible band at 392 nm (ε = 2.44 x 10³ M⁻¹ cm⁻¹), besides more intense near-UV band at 298 nm (ε = 1.21 x 10⁴ M⁻¹ cm⁻¹) that is typical for a bpy π-π* transition (Figure 1, Table S6).

From these results, [1] is assigned to a mononuclear platinum(II) complex, [Pt(H₄tg-κS)₂(bpy)], which contains two monodentate-S H₄tg⁻ ligands. Compatible with the presence of H₄tg⁻ ligands, [1] is optically active, showing positive CD bands in the LL’CT and π-π* transition regions (Figure 1).
Synthesis and Characterization of $[\text{Pt}_2(\mu_2-\text{H}_4\text{tg-}\kappa_1^1\text{S};\kappa^1_1\text{S})_2(\text{bpy})_2]^{2+}$ ([2]$^{2+}$). To investigate the functionality of [1] as an S-donating metalloligand, the reaction of [1] with [PtCl$_2$(bpy)] was carried out. When [1] was treated with 1 molar equiv of [PtCl$_2$(bpy)] in water, an orange suspension turned to a yellow-brown solution, from which yellow crystals ([2](ClO$_4$)$_2$) were isolated in a satisfactory yield by the addition of an aqueous solution of NaClO$_4$. The elemental analytical data of [2](ClO$_4$)$_2$ were in agreement with the formula for [Pt$_2$(H$_4$tg$_2$(bpy)$_2$(ClO$_4$)$_2$, and the divalent charge of its complex-cation was implied by the molar conductivity in water (191.0 S cm$^2$ mol$^{-1}$). The presence of ClO$_4^-$ as a counter anion was indicated by its IR spectrum (Figure S2), which shows a strong band at 623 cm$^{-1}$ characteristic for ClO$_4^-$.$^{23}$ As in the case of [1], the $^1$H NMR spectrum of [2](ClO$_4$)$_2$ in D$_2$O exhibits aliphatic proton signals in a region $\delta$ 3.2-5.0 and aromatic proton signals in a region $\delta$ 7.8-8.9 (Figure S1). However, the integration ratio of the aliphatic and aromatic proton signals is 7:8, indicating that [2]$^{2+}$ contains H$_4$tg$^-$ and
bpy ligands in a 1:1 ratio. Based on these results, \([\text{2}]^{2+}\) is assigned to a diplatinum(II) complex, 
\([\text{Pt}_2(\mu_2-\text{H}_4\text{tg}\text{-}\kappa\text{1}S:\kappa\text{1}S)_{2}(\text{bpy})_{2})^{2+}\), in which two \([\text{Pt( bpy)}]^{2+}\) units are spanned by two \(\text{H}_4\text{tg}^-\) ligands through sulfur-bridges.\(^{24,25}\) This assignment was confirmed by single-crystal X-ray analysis for 
\([\text{2}](\text{ClO}_4)_2\cdot 9\text{H}_2\text{O}\).

The asymmetric unit of \([\text{2}](\text{ClO}_4)_2\cdot 9\text{H}_2\text{O}\) contains a complex-cation and two perchlorate anions, besides solvated water molecules. The number of perchlorate anions implies that the complexcation is divalent, consistent with the molar conductivity. As shown in Figure 2, the complexcation \([\text{2}]^{2+}\) consists of a \([\text{Pt( H}_4\text{tg}_{2}(\text{bpy})]\) unit that chelates to a \([\text{Pt( bpy)}]^{2+}\) moiety through thiolato S atoms, forming an S-bridged dinuclear structure in 
\([\text{Pt}_2(\mu_2-\text{H}_4\text{tg}\text{-}\kappa\text{1}S:\kappa\text{1}S)_{2}(\text{bpy})_{2})^{2+}\) with a \(\text{Pt}^{\text{II}}_2\text{S}_2\) diamond core (av. S-Pt-S = 82.02(2)\(^{\circ}\), Pt-S-Pt = 96.8(6)\(^{\circ}\)). This implies that \([\text{Pt( H}_4\text{tg}\text{-}\kappa\text{S})_{2}(\text{bpy})]\) (1) acts as a bidentate-S,S’ chelating metalloligand toward a Pt\(^{\text{II}}\) center. Alternatively, it is considered that this dinuclear structure in \([\text{2}]^{2+}\) is constructed by the connection of two \([\text{Pt( bpy)}]^{2+}\) units with use of two bridging \(\text{H}_4\text{tg}^-\) ligands. Indeed, \([\text{2}](\text{ClO}_4)_2\) was also obtained by the direct reaction of \([\text{PtCl}_2(\text{bpy})]\) with \(\text{NaH}_4\text{tg}\) in a 1:1 ratio in water (Scheme 1). Each Pt\(^{\text{II}}\) atom in \([\text{2}]^{2+}\) has a square-planar \textit{cis}-\(N_2S_2\) geometry (av. N-Pt-N = 79.78(14)\(^{\circ}\), N-Pt-S = 99.1(15)\(^{\circ}\)), coordinated by a bidentate-N,N’ bpy and two monodentate-S \(\text{H}_4\text{tg}\text{-}\) ligands. The Pt-N (av. 2.046(5) Å) and Pt-S (av. 2.296(7) Å) bond distances are comparable with the normal Pt-N (av. 2.054 Å) and Pt-S (av. 2.308 Å) distances for \([\text{Pt}^{\text{II}}(N)_2(S)_2]\)-type complexes.\(^{26}\) The bond distances and angles about each \(\text{H}_4\text{tg}\text{-}\) ligand are very similar to those found in \(\text{KH}_4\text{tg},^{27}\) having the \(\beta\)-D-pyranose ring with the usual chair conformation. Considering the relative arrangement of two \(\beta\)-D-pyranose moieties, two geometrical configurations, \textit{syn} and \textit{anti}, are possible for this type of dinuclear complexes (Figure 3). As shown in Figure 2, \([\text{2}]^{2+}\) adopts the \textit{syn} configuration that appears to be sterically
crowded. To avoid the steric repulsion, the two PtN$_2$S$_2$ planes are deviated from co-planarity with a dihedral angle of 164.36(4)$^\circ$. In packing structure, each diplatinum(II) complex-cation is connected to 4 adjacent complex-cations through intermolecular $\pi$-$\pi$ interactions between bpy ligands (av. C···C = 3.34(6) Å), constructing a 2D layer structure with a hydrophilic surface covered with $\beta$-D-pyranose moieties and a hydrophobic surface covered with bpy ligands (Figure S4). These layers are stacked in an (AB)$_n$ fashion such that wide hydrophilic and narrow hydrophobic 2D coordination spaces are alternately formed. There exist interlayer $\pi$-$\pi$ interactions between bpy ligands in the hydrophobic spaces (av. C···C = 3.30(3) Å), while hydroxyl groups of $\beta$-D-pyranose moieties form hydrogen bonding networks with ClO$_4^-$ anions and water molecules in the hydrophilic spaces (av. O$_\text{OH}$···O$_\text{anion}$ = 2.871(6) Å, O$_\text{OH}$···O$_\text{water}$ = 2.8(2) Å). It seems that these multiple $\pi$-$\pi$ and hydrogen-bonding interactions are responsible for the formation of the syn configuration in [2]$^{2+}$, rather than the anti configuration.

![Figure 2. Perspective view of [2]$^{2+}$. Pt: green, S: yellow, O: pink, N: blue, C: gray, H: light gray.](image)
The absorption spectrum of $[2](\text{ClO}_4)_2$ in water in the near-UV region resembles that of $[1]$, showing a bpy $\pi-\pi^*$ transition band at a similar wavelength (Figure 1). In the visible region, however, no clear LL’CT band is observed for $[2](\text{ClO}_4)_2$. This is most likely due to the formation of a $\mu_2$-thiolato structure in $[2]^{2+}$, which stabilizes the HOMO of the thiolato donor and causes the blue shift of the LL’CT band. In parallel with the absorption spectral feature, $[2](\text{ClO}_4)_2$ does not show any detectable CD bands in the visible region, although several CD bands are observed in the near-UV region (Figure 1).

**Synthesis and Characterization of** $[\text{PtCu}(\mu_2-H_4\text{tg-}\kappa^1\text{S}:\kappa^2\text{O},\text{S})_2(\text{bpy})_2]^{2+}$ ($[3]^{2+}$) and $[\text{PtNi}(\mu_2-H_4\text{tg-}\kappa^1\text{S}:\kappa^2\text{O},\text{S})_2(\text{bpy})_2]^{2+}$ ($[4]^{2+}$). To clarify the possible coordination mode of $[1]$ toward transition metal ions that can take both square-planar and octahedral geometries, the reactions of $[1]$ with Cu$^{2+}$ and Ni$^{2+}$ in the presence of bpy were studied. Treatment of an aqueous suspension of $[1]$ with $[\text{CuCl}_2(\text{bpy})]$ in a 1:1 ratio gave a green solution, from which green crystals ($[3](\text{NO}_3)_2$) were isolated in a moderate yield by the addition of an aqueous solution of NaNO$_3$. Similar treatment of $[\text{Pt}(H_4\text{tg-}\kappa^S)_2(\text{bpy})]$ with a 1:1 mixture of Ni(NO$_3$)$_2$·6H$_2$O and bpy in water, followed by the addition of NaNO$_3$, afforded pale green crystals ($[4](\text{NO}_3)_2$). X-ray fluorescence spectrometry indicated the presence of Cu and Ni atoms in $[3](\text{NO}_3)_2$ and $[4](\text{NO}_3)_2$, respectively, besides Pt atoms, and the elemental analytical data of $[3](\text{NO}_3)_2$ and $[4](\text{NO}_3)_2$ were

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**Figure 3.** *Syn* and *anti* configurations for a dinuclear complex.
consistent with the formulas for \([\text{PtCu(H}_{4}\text{tg})_{2}(\text{bpy})_{2}](\text{NO}_3)_{2}\) and \([\text{PtNi(H}_{4}\text{tg})_{2}(\text{bpy})_{2}](\text{NO}_3)_{2}\), respectively. The divalent nature of the complex-cations in \([3](\text{NO}_3)_{2}\) and \([4](\text{NO}_3)_{2}\) was implied by their molar conductivities in water (192.7 S cm\(^2\) mol\(^{-1}\) for \([3](\text{NO}_3)_{2}\), 210.2 S cm\(^2\) mol\(^{-1}\) for \([4](\text{NO}_3)_{2}\)), and the presence of NO\(_3^-\) as counter anions was indicated by the appearance of an IR band at 1382 cm\(^{-1}\) characteristic for NO\(_3^-\) (Figure S2).\(^{23}\) The Pt\(^{II}\)Cu\(^{II}\) and the Pt\(^{II}\)Ni\(^{II}\) dinuclear structures in \([3]\)^\(^{2+}\) and \([4]\)^\(^{2+}\) were established by single-crystal X-ray crystallography for \([3](\text{ClO}_4)_{2}\cdot2\text{H}_2\text{O}\) and \([4](\text{NO}_4)_{2}\cdot2\text{H}_2\text{O}\), respectively.

The asymmetric unit of \([3](\text{ClO}_4)_{2}\cdot2\text{H}_2\text{O}\) contains a complex-cation and two perchlorate anions, besides solvated water molecules. The number of perchlorate anions implies that the complex-cation is divalent. As shown in Figure 4, the complex-cation \([3]\)^\(^{2+}\) consists of a square-planar \([\text{Pt(H}_{4}\text{tg})_{2}(\text{bpy})]\) unit that chelates to a \([\text{Cu(bpy)})\]^\(^{2+}\) moiety through thiolato S atoms, forming an S-bridged Pt\(^{II}\)Cu\(^{II}\) dinuclear structure with a Pt\(^{II}\)Cu\(^{II}\)S\(_2\) diamond core (av. S-Pt-S = 84.94(8)\(^{\circ}\), Pt-S-Cu = 95.8(10)\(^{\circ}\), S-Cu-S = 83.14(9)\(^{\circ}\), Pt-S = 2.288(9) Å, Cu-S = 2.33(3) Å). It is interesting to note that thiolato-bridged dinuclear complexes with a Pt\(^{II}\)-S-Cu\(^{II}\) linkage have not been reported to date. The bond distances and angles about the Pt\(^{II}\) center in \([3]\)^\(^{2+}\) (av. N-Pt-N = 79.6(4)\(^{\circ}\), N-Pt-S = 97.8(9), Pt-N = 2.052(11) Å), as well as the chair conformational \(\beta\)-D-pyranose ring of H\(_{4}\)tg\(^{-}\), are very similar to those in \([2]\)^\(^{2+}\). In \([3]\)^\(^{2+}\), however, two hydroxyl O atoms from two H\(_{4}\)tg\(^{-}\)ligands also bind to a Cu\(^{II}\) center at apical positions, with one shorter and the other longer Cu-O distances (Cu1-O2 = 2.386(4) Å, Cu1-O7 = 2.680(6) Å), such that the Cu\(^{II}\) atom is situated in a Jahn-Teller distorted octahedral geometry with a \(trans(O)\)-N\(_2\)O\(_2\)S\(_2\) donor set. This implies that \([\text{Pt(H}_{4}\text{tg-κS})_{2}(\text{bpy})]\) acts as a tetradentate-O,O’,S,S’ metalloligand toward a Cu\(^{II}\) center by using two thiolato and two hydroxyl groups. Unlike in \([2]\)^\(^{2+}\), the two H\(_{4}\)tg\(^{-}\) ligands in \([3]\)^\(^{2+}\) adopt the \(anti\) configuration, which allows the two H\(_{4}\)tg\(^{-}\) ligands to chelate to the Cu\(^{II}\) center through
thiolato and hydroxyl groups. Of two chiral configurations, Δ and Λ, which arise from the skew pair of a N,N’-bpy and two O,S-H₄tg⁻ chelate rings, the CuII center in [3]²⁺ has the Δ configuration. This chiral configuration regulates the two asymmetric bridging S atoms of H₄tg⁻ to have the R configuration. The five-membered O,S-chelate rings of H₄tg⁻ adopt a gauche form with the λ conformation that is thermodynamically stable.²⁸ Molecular model examinations reveal that the two O,S-chelate rings are enforced to have an unfavorable envelop form when the CuII center has the Λ configuration. Moreover, there exists a non-bonding interaction between a C2 proton of H₄tg⁻ and a bpy aromatic ring of [Pt(bpy)]²⁺ in the Λ configuration. The selection of the Δ configuration in [3]²⁺ is most likely ascribed to these steric factors that are caused by the presence of the rigid, bulky H₄tg⁻ ligands with the β-D-pyranose ring. In packing structure, the complex-cations [3]²⁺ are connected with each other through intermolecular π-π interactions (av. C…C = 3.45(10) Å) between bpy ligands, forming a 2D sheet-like structure (Figure S5). The 2D sheets are further connected by intermolecular hydrogen bonds between hydroxyl groups of H₄tg⁻ (av. O…O = 2.748(3) Å), which completes a 3D network structure.
The asymmetric unit of \([4](\text{NO}_3)_2\cdot2\text{H}_2\text{O}\) contains a complex-cation and two nitrate anions, besides solvated water molecules. The number of nitrate anions implies that the complex-cation is divalent. As shown in Figure 5, the complex cation \([4]^{2+}\) consists of a square-planar \([\text{Pt(H}_4\text{tg)}_2(\text{bpy})]\) unit that binds to a \([\text{Ni(bpy)}]^{2+}\) moiety through two thiolato S and two hydroxyl O atoms from two H\(_4\text{tg}^-\) ligands, forming an S-bridged Pt\(^{II}\)Ni\(^{II}\) dinuclear structure in \([\text{PtNi(\mu}_2\text{-H}_4\text{tg-κ}^1\text{S:κ}^2\text{O,S})}_2(\text{bpy})_2]\)\(^{2+}\) with a Pt\(^{II}\)Ni\(^{II}\)S\(_2\) diamond core (av. S-Pt-S = 86.40(4)\(^\circ\), Pt-S-Ni = 96.0(1)\(^\circ\), S-Ni-S = 80.71(4)\(^\circ\), Pt-S = 2.30(2) \(\text{Å}\), Ni-S = 2.43(3) \(\text{Å}\)). The overall dinuclear structure in \([4]^{2+}\) with \textit{anti} configurational H\(_4\text{tg}^-\) ligands corresponds well with that in \([3]^{2+}\), except the presence of Ni\(^{II}\) atom in place of Cu\(^{II}\) atom. In particular, the bond distances and angles about the Pt\(^{II}\) atom in \([4]^{2+}\) (av. N-Pt-N = 79.69(15)\(^\circ\), N-Pt-S = 97(2)\(^\circ\), Pt-N = 2.058(1) \(\text{Å}\)) are very similar to those in \([3]^{2+}\). The Ni-S bond distances in \([4]^{2+}\) are longer than the Cu-S distances in \([3]^{2+}\), while the Ni-O bond distances in \([4]^{2+}\) (2.139(3) \(\text{Å}\), 2.145(3) \(\text{Å}\)) are much shorter than the Cu-O distances in \([3]^{2+}\). In addition, the O-Ni-O angle in \([4]^{2+}\) (176.77(12)\(^\circ\)) is very close to 180\(^\circ\), compared with the O-Cu-O angle in \([3]^{2+}\) (166.83(15)\(^\circ\)). Thus, the Ni\(^{II}\) atom in \([4]^{2+}\) has an octahedral geometry more ideal than the Cu\(^{II}\) atom in \([3]^{2+}\), which is understood by the absence of the Jahn-Teller distortion for a d\(^8\) metal center. Like the Cu\(^{II}\) center in \([3]^{2+}\), the Ni\(^{II}\) center in \([4]^{2+}\) is chelated by a bidentate-N,N’ bpy and two bidentate-O,S H\(_4\text{tg}^-\) ligands to have the Δ configuration, with two bridging S atoms being fixed to the R configuration. A related S-bridged Pt\(^{II}\)Ni\(^{II}\) dinuclear complex with two terminal bpy ligands, \([\text{PtNi(μ}_2\text{-aet-κ}^1\text{S:κ}^2\text{S,N})}_2(\text{bpy})_2]\)\(^{2+}\) (aet = 2-aminoethanethiolate), has been reported, although a pair of enantiomers (Δ\(_{RR}/Λ_{SS}\)) with the Δ/Δ configurational Ni center and the R/S configurational S center is formed in this structure because
of the use of achiral aet ligands. In the packing structure, the complex-cations are connected with each other through intermolecular π-π interactions to form a 1D chain (av. C…C = 3.48(12) Å) in [4](NO3)2·2H2O, and the 1D chains are further connected by intermolecular hydrogen bonds between hydroxyl groups of H4tg− (av. O…O = 3.0(2) Å) to complete a 3D network structure (Figure S6).

**Figure 5.** Perspective view of [4]2+. Pt: green, Ni: yellow-green, S: yellow, O: pink, N: blue, C: gray, H: light gray.

The absorption spectrum of [3](NO3)2 in water is characterized by a weak visible band at 616 nm (ε = 4.2 x 10^2 M⁻¹ cm⁻¹), which is typical for a d-d transition band found in CuII species, besides an intense near-UV band at 310 nm (ε = 2.61 x 10^4 M⁻¹ cm⁻¹) due to a bpy π-π* transition (Figure 6). This absorption spectral feature is essentially the same as that of its diffuse reflection spectrum in the solid state (Figure S7). In addition, the CD spectrum of [3](NO3)2 in water gives a negative CD band at 603 nm (Δε = −3.33 M⁻¹ cm⁻¹) in the visible d-d band region, which is also found in its CD spectrum in the solid state. Thus, it is reasonable to assume that the
S-bridged Pt\textsuperscript{II}Cu\textsuperscript{II} dinuclear structure in [3]\textsuperscript{2+} is retained in water, although a partial Cu-O bond cleavage may not be ruled out.

**Figure 6.** Absorption and CD spectra of complexes [3](NO\textsubscript{3})\textsubscript{2} (left) and [4](NO\textsubscript{3})\textsubscript{2} (right) in H\textsubscript{2}O (solid line) and MeOH (dashed line).

When [4](NO\textsubscript{3})\textsubscript{2} is dissolved in water, its absorption and CD spectra does not give any detectable band in the visible region, and their spectral features are very similar to those of [1] ([Pt(H\textsubscript{4}tg-κ\textsubscript{S})(bpy)]) over the whole region (Figure 6). In the solid state, [4](NO\textsubscript{3})\textsubscript{2} exhibits a characteristic weak visible band (574 nm) assignable to a d-d transition for an octahedral Ni\textsuperscript{II} center\textsuperscript{31} and a negative CD band (569 nm) in this d-d band region (Figure S8). Thus, it is considered that the S-bridged Pt\textsuperscript{II}Ni\textsuperscript{II} structure of [4]\textsuperscript{2+} found in the crystal is decomposed into the mononuclear [1] and aqua Ni\textsuperscript{II} species in water. This seems to be related to the longer Ni-S bonds in [4]\textsuperscript{2+}, which facilitates the cleavage of Ni-S bonds in water. On the other hand, the absorption spectrum of [4]\textsuperscript{2+} in methanol shows a weak d-d band at 571 nm ($\varepsilon = 29 \text{ M}^{-1} \text{ cm}^{-1}$),
besides a bpy $\pi$-$\pi^*$ transition band at 305 nm ($\varepsilon = 2.7 \times 10^4 \text{ M}^{-1} \text{ cm}^{-1}$) (Figure 6). In addition, a negative CD band is observed in the d-d band region (568 nm), as in the case of the CD spectrum in the solid state. This is suggestive of the retention of the S-bridged Pt$^\text{II}$Ni$^\text{II}$ dinuclear structure in methanol, which is understood by a weaker coordination ability of MeOH compared with that of H$_2$O.

**Synthesis and Characterization of** $[\text{Cd}\{\text{Pt}(\mu_2\text{-H}_4\text{tg-}\kappa_1\text{S};\kappa_2\text{O},\text{S})(\mu_2\text{-H}_4\text{tg-}\kappa_1\text{S};\kappa_1\text{S})(\text{bpy})\}_2]^2^+ ([5]^2^+)$. We also investigated the coordination ability of [1] toward thiophilic Cd$^{2+}$ that can take various coordination geometries, including tetrahedron and octahedron. When $[\text{Pt(H}_4\text{tg-}\kappa\text{S})_2(\text{bpy})]$ was treated with a 1:1 mixture of CdCl$_2$·2.5H$_2$O and bpy in water, we obtained a yellow reaction solution, from which yellow block crystals ($[5](\text{ClO}_4)_2$) were isolated by the addition of NaClO$_4$. Compound $[5](\text{ClO}_4)_2$ is almost insoluble in any common solvents, including water and alcohol, and thus, its characterization in solution could not be carried out. The presence of bpy and H$_4$tg$^-$ ligands in $[5](\text{ClO}_4)_2$, besides ClO$_4^-$ ions, was confirmed by the IR spectrum that is similar to the spectrum of $[2](\text{ClO}_4)_2$ (Figure S2), the diffuse reflection spectrum that shows characteristic $\pi$-$\pi^*$ transition bands centered at 325 nm, and the solid-state CD spectrum that gives a negative band at 341 nm (Figure S9). However, the elemental analytical data for $[5](\text{ClO}_4)_2$ is not in agreement with a formula for an expected Pt$^\text{II}$Cd$^\text{II}$ dinuclear complex but for a Pt$^\text{II}$Cd$^\text{II}$Pt$^\text{II}$ trinuclear complex, $[\text{Cd}\{\text{Pt(H}_4\text{tg)}_2(\text{bpy})\}_2](\text{ClO}_4)_2$. In addition, X-ray fluorescence spectrometry implies the presence of Pt and Cd atoms in a 2:1 ratio in $[5](\text{ClO}_4)_2$, rather than in a 1:1 ratio. These results indicate that $[5]^{2+}$ contains $[\text{Pt(H}_4\text{tg)}_2(\text{bpy})]$ and Cd$^{\text{II}}$ in a 2:1, with the lack of an additional bpy ligand bound to a Cd$^{\text{II}}$ center. Consistent with this, yellow block crystals of $[5](\text{ClO}_4)_2$ were obtained in a reasonable yield by the 2:1 reaction of [1]·2H$_2$O with CdBr$_2$ in water, followed by the addition of NaClO$_4$, without the
addition of bpy to the reaction mixture. The Pt\textsuperscript{II}Cd\textsuperscript{II}Pt\textsuperscript{II} trinuclear structure in \([5]\)\textsuperscript{2+} was unambiguously determined by single-crystal X-ray analysis for \([5](\text{ClO}_4)\)\textsubscript{2}6H\textsubscript{2}O.

As shown in Figure 7, the complex cation consists of two square-planar \([\text{Pt}(\text{H}_4\text{tg})_2(bpy)]\) units and one Cd\textsuperscript{II} atom. Each of two \([\text{Pt}(\text{H}_4\text{tg})_2(bpy)]\) units chelates to the Cd\textsuperscript{II} atom through two thiolato S atoms (av. Cd-S = 2.65(4) Å, S1-Cd-S1 = 93.42(8)°, S1-Cd-S2 = 75.00(5)°, S2-Cd-S2 = 118.82(8)°), forming an S-bridged Pt\textsuperscript{II}Cd\textsuperscript{II}Pt\textsuperscript{II} trinuclear structure with a crystallographic \(C_2\) symmetry. An interesting structural feature of \([5]\)\textsuperscript{2+} is the coordination of a hydroxyl group from one of two H\textsubscript{4}tg\textsuperscript{-} ligands in each \([\text{Pt}(\text{H}_4\text{tg})_2(bpy)]\) unit (Cd-O = 2.446(5) Å), such that the Cd\textsuperscript{II} center is situated in a highly distorted octahedral geometry with a trans(O)-O\textsubscript{2}S\textsubscript{4} donor set (O-Cd-O = 145.0(2)°, trans S-Cd-S = 163.08(6)°). Thus, \([5]\)\textsuperscript{2+} is formulated as \([\text{Cd}\{\text{Pt}(\mu_2-\text{H}_4\text{tg}-\kappa^1S: \kappa^2O,S)(\mu_2-\text{H}_4\text{tg}-\kappa^1S: \kappa^1S)(bpy)\}_2\}]^{2+}\), in which each \([\text{Pt}(\text{H}_4\text{tg})_2(bpy)]\) unit acts as a tridentate-O,S,S’ metalloligand. This coordination mode of \([\text{Pt}(\text{H}_4\text{tg})_2(bpy)]\) is distinct from the modes found in \([2]\)\textsuperscript{2+}, \([3]\)\textsuperscript{2+}, and \([4]\)\textsuperscript{2+}, although the bond distances and angles about the \([\text{Pt}(\text{H}_4\text{tg})_2(bpy)]\) unit (av. S-Pt-S = 89.56(7)°, N-Pt-N = 78.9(2)°, trans N-Pt-S = 173(2)°, Pt-S = 2.293(4) Å, Pt-N = 2.076(6) Å) are very similar. Like in \([3]\)\textsuperscript{2+} and \([4]\)\textsuperscript{2+}, the four bridging S atoms in \([5]\)\textsuperscript{2+} adopt the \(R\) configuration, and the two O,S-chelate rings of H\textsubscript{4}tg\textsuperscript{-} have a gauche form with the \(\lambda\) conformation. However, the Cd center in \([5]\)\textsuperscript{2+} is regulated to have the \(\Lambda\) configuration, which is opposite to the \(\Delta\) configurational Cu\textsuperscript{II} and Ni\textsuperscript{II} centers in \([3]\)\textsuperscript{2+} and \([4]\)\textsuperscript{2+}. Thus, the bis(O,S-chelation) mode to an octahedral metal center from two \([\text{Pt}(\text{H}_4\text{tg})_2(bpy)]\) units leads to the chiral configuration opposite to that formed by the bis(O,S-chelation) mode from one \([\text{Pt}(\text{H}_4\text{tg})_2(bpy)]\) unit. Molecular model examinations indicate that the O,S-chelate rings are enforced to have an unfavorable envelop form when the Cd\textsuperscript{II} center adopts the \(\Delta\) configuration. Moreover, there exists a non-bonding interaction between a C2 proton of H\textsubscript{4}tg\textsuperscript{-} and a lone-pair on bridging S
atom from different [Pt(H₄tg)₂(bpy)] unit in the Δ configuration. Notable are the existing intramolecular hydrogen bonds between hydroxyl groups of H₄tg⁻ from different [Pt(H₄tg)₂(bpy)] units in [5]²⁺ (av. O···O = 2.68(5) Å), which appear to sustain the PtIICdIIPtII trinuclear structure with the Δ configurational CdII center. In packing structure, each complexation [5]²⁺ is hydrogen-bonded with 6 adjacent complex-cations through hydroxyl groups of H₄tg⁻ (av. O···O = 2.78(5) Å), constructing a 3D network structure (Figure S10). The 3D network structure is reinforced by hydrogen bonds between hydroxyl groups of H₄tg⁻ and ClO₄⁻ ions (av. O···O = 2.91(3) Å). It is considered that these multiple hydrogen-binding interactions are responsible for the insolubility of [5](ClO₄)₂ in any common solvents.


**CONCLUDING REMARKS**

In this study, we showed that Cl⁻ ions in [PtCl₂(bpy)] are readily replaced by H₄tg⁻ in water to form the mononuclear [Pt(H₄tg-κS)₂(bpy)] ([1]). Compound [1] further reacted with [PtCl₂(bpy)] to give the diplatinum(II) complex, [Pt₂(μ₂-H₄tg-κ²:S;κ¹S)₂(bpy)₂]²⁺ ([2]²⁺), in which two [Pt(bpy)]²⁺ moieties are spanned by two H₄tg⁻ ligands through thiolato S atoms. This result
indicates that [1] functions as a bidentate-S,S’ metalloligand toward a square-planar metal center due to the high nucleophilicity of H₄tg⁻ thiolato groups bound to a Pt²⁺ center. Interestingly, [1] was found to serve as a tetradequate-O,O’,S,S’ metalloligand to M = Cu²⁺, Ni³⁺ in the presence of bpy, affording the heterometallic dinuclear structures in [PtCu(μ₂-H₄tg-κ¹:S:κ²O,S)₂(bpy)₂]²⁺ ([3]²⁺) and [PtNi(μ₂-H₄tg-κ¹:S:κ²O,S)₂(bpy)₂]²⁺ ([4]²⁺), which contain an octahedral metal center bound by two thiolato and two hydroxyl groups, besides a bpy ligand. While several metal complexes with H₄tg⁻ or H₃tg²⁻ have been reported, coordination of a hydroxyl group toward a metal center has not been recognized. A similar reaction with Cd²⁺ led to the formation of the Pt³⁺Cd³⁺Pt³⁺ trinuclear complex, [Cd{Pt(μ₂-H₄tg-κ¹:S:κ²O,S)(μ₂-H₄tg-κ¹:S:κ¹S)(bpy)}₂]²⁺ ([5]²⁺), in which each of two molecules of [1] binds to an octahedral Cd³⁺ center in a tridentate-O,S,S’ mode. Thus, the present study showed that [1] functions as a versatile multidentate metalloligand, the coordination modes of which are varied with response to the coordination environment of a metal center. Of two geometrical configurations, syn and anti, [2]²⁺ adopted the syn configuration so as to form both of the intermolecular O-H···O hydrogen bonds between hydroxyl groups of H₄tg⁻ and the intermolecular π-π interactions between bpy ligands. On the other hand, [3]²⁺, [4]²⁺, and [5]²⁺ selected the anti configuration, which permits H₄tg⁻ in [1] to chelate to a metal center through thiolato and hydroxyl groups to complete an octahedral geometry. The most remarkable structural feature in the present system is the formation of the Δ configurational metal center in [3]²⁺ and [4]²⁺ and the Λ configurational metal center in [5]²⁺, indicative of the availability of [1] as the chiral-selective construction of S-bridged multinuclear structures. This is explained by the steric demand around an octahedral metal center, which is regulated by the coordination of chiral H₄tg⁻ with a rigid, bulky β-D-pyranose ring. It is worth to mentioning that the Pt³⁺Cu³⁺ structure in [3]²⁺ is retained in water, while the Pt³⁺Ni³⁺ structure in
[4]$^{2+}$ is decomposed into [1] in water, although [4]$^{2+}$ is fairly stable in methanol. The difference in stability between [3]$^{2+}$ and [4]$^{2+}$ in water is most likely ascribed to the stronger Cu-S bonds in [3]$^{2+}$, which is induced by the Jahn-Teller elongation of axial Cu$^{II}$-O bonds. Finally, it should be noted that only a few transition metal complexes with H$_4$tg$^-$ or H$_3$tg$^{2-}$ have been isolated and crystallographically characterized to date.$^{9,10}$ The successful preparation and structural determination of all the multinuclear complexes obtained in this work are thanks to the use of the chiral metalloligand with H$_4$tg$^-$, which leads to the stereoselective formation of a single isomer for each complex, as well as thanks to the introduction of a hydrophobic co-ligand of bpy in the hydrophilic H$_4$tg$^-$ system. A similar strategy could be applicable for the isolation and structural characterization of other monosaccharide coordination systems.

ASSOCIATED CONTENT

**Supporting Information.** X-ray crystallographic files in CIF format for all complexes, crystallographic data, selected bond distances and angles, Figures of IR, solid-state diffuse reflection, CD, ESI-mass and NMR spectra. This material is available free of charge via the Internet at http://pubs.acs.org.

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The retention of the $C_2$ symmetrical diplatinum(II) structure in [2]$^{2+}$ in solution is supported by its $^{13}$C NMR spectrum in D$_2$O, which shows only a single set of 16 carbon signals (Figure S1). The $^{13}$C NMR spectrum of the other diamagnetic [1], which is neutral species, could not be measured because of its poor solubility.

In the ESI mass spectrum in MeOH, [2](ClO$_4$)$_2$ dominantly showed a cluster of signals at $m/z = 546.1$ corresponding to [Pt$_2$(H$_4$tg)$_2$(bpy)$_2$]$^{2+}$ (Figure S3). The ESI mass spectra of [3](ClO$_4$)$_2$ and [4](NO$_3$)$_2$ displayed main signals at $m/z = 764.1$ corresponding to {[Pt(H$_4$tg)$_2$(bpy)]$^+$}+Na$^+$, suggestive of the dissociation of [3]$^{2+}$ and [4]$^{2+}$ into the mononuclear [1] under a very dilute MeOH solution. The ESI mass spectra of [1] and [5](ClO$_4$)$_2$ could not be measured because of their insolubility in MeOH.

The averaged Pt–N and Pt–S coordination bonds were evaluated based on 275 crystal structures with $R < 0.075$ from the December 2012 version of the Cambridge Crystallographic Database.


(32) A ruthenium(II) complex with $\text{H}_4\text{tg}^-$ has recently been reported. However, its structural characterization has not been made, and the coordination of hydroxyl group to a metal center has not been established. See the reference 8c.