Electrophilic Addition of Polyaromatic o–Quinones to an Iron(II) Diaminoclahrochelate Leading to the Macrobicyclic Iron(II) tris–Dioximates with Extended Annulated Ribbed Substituents

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Macrobicyclic iron(II) tris-dioximate with two inherent amino groups undergoes electrophilic addition with 1,10-phenanthroline-5,6-dione and phenanthrene-9,10-dione to give a cage complexes with extended ribbed substituents. These clathrochelates have been characterized using single crystal XRD, multinuclear NMR and other spectral techniques. They have been found to undergo the ligand-centered redox processes; the obtained phenanthroline-containing iron(II) cage complex is proposed as a potent chelating clathrochelate ligand.

Keywords: Macrocyclic compounds, clathrochelates, ligand reactivity, electrophilic addition.

Электрофильное присоединение полиароматических o–хинонов к диаминоклатрохелату железа(II), приводящее к макробициклическим трис–диоксиматам железа(II) с расширенными аннелированными реберными заместителями

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Трис-диоксиматный клатрохелат железа(II) с двумя амино-группами вступает в реакцию электрофильного присоединения с 1,10-фенантролин-5,6-дионом и фенантрен-9,10-дионом, приводящую к клеточным комплексам с расширенными реберными заместителями. Эти клатрохелаты были охарактеризованы методами РСА, мультиядерной ЯМР-спектроскопии и другими спектральными методами. Установлено, что полученные клеточные комплексы претерпевают лиганд-центрированные редокс-процессы; фенантролин-содержащий клеточный комплекс железа(II) предложен в качестве перспективного хелатирующего клатрохелатного лиганда.

Ключевые слова: Макроциклические соединения, клатрохелаты, реакционная способность лигандов, электрофильное присоединение.
Introduction

Boron-capped metal tris-dioximates (cage complexes, clathrochelates), possessing rather high chemical stability combined with their ability to be functionalized with various substituents, are promising molecular platforms for the design of polytopic multifunctional molecular and supramolecular systems, including those prospective as molecular electronic devices.[1–4] Recently we described an iron(II) clathrochelate with annulated ribbed quinoxaline fragment that exhibited the reversible ligand-centered redox transformations coupled with electrochromism. [5] Its annulated azaheterocyclic moiety has been constructed using nucleophilic substitution of the reactive chlorine atoms in a dichloroclatrochelate precursor FeBd2(Cl2Gm)(BF2), (where Bd2- and Cl2Gm2- are α-benzildioxime and dichloroglyoxime dianions) (Scheme 1).

An alternative pathway to such annulated azaheterocycles is based on the reactions of the appropriate carbonyl compounds as electrophilic agents towards the anionic derivatives of a diaminoclathrochelate FeBd2((NH2)2Gm)(BF2) (where Gm is glyoxime residue) as a reactive macrobicyclic precursor (Scheme 2).[6]

The same synthetic approach (i.e. electrophilic addition) was successfully used to extend the family of the iron(II) clathrochelates with redox-non-innocent azaheterocyclic fragments via condensation of the deprotonated form of this diaminoclatrochelate precursor with polyaromatic α-quiones, such as 1,10-phenanthroline-5,6-dione and phenanthrene-9,10-dione. The presence of donor phenanthroline moiety in a clathrochelate molecule allows the synthesis of heterometallic coordination compounds with clathrochelate ligands.

Experimental

Materials and Methods

Diaminoclatrochelate FeBd2(Gm(NH2)2)(BF2) was obtained as described elsewhere. [5] DMSO was dried over NaOH pellets and distilled in vacuo prior to use. Acetonitrile was purified according to the standard procedure. [8] The commercial reagents, 1,10-phenanthroline-5,6-dione, phenanthrene-9,10-dione and 1,8-diazabicyclo[5.4.0]undecene-7 (DBU) were purchased from Sigma-Aldrich® and used without their further purification. The solution UV-Vis spectra were recorded on a Specord UV-VIS and a Shimadzu UV-3101 PC spectrophotometers. 1H, 13C, 11B and 19F NMR spectra were recorded on a Bruker Avance III 500 spectrometer (working frequencies 500.03 (1H), 125.73 (13C), 160.33 (11B) and 470.49 MHz (19F)) in CDCl3 solutions. The 1H and 13C NMR chemical shifts are reported in ppm of the δ scale.

Scheme 1.

Scheme 2.
Electrophilic Addition of Polyaromatic o-Quinones to an Iron(II) Diaminocloathrochelate

and referred to signals of the solvent (7.26 ppm for residual protons in 1H NMR spectra and 77.23 ppm in 13C{1H} NMR spectra). 1B and 19F NMR chemical shifts are referred to the external standards BF3·O(C2H5)2 (δ0 H=6 ppm) and C6H6CF3 (δ0 H= –63.72 ppm), respectively. Heteronuclear C–H correlations (HMBC and 1H–1H couplings) are not exact values but only illustrate the shape of the observed multiplets.

X-Ray Crystallography

Intensity data for the complexes FeBd2((Phn)Gm)(BF)2·nCHCl3 and FeBd2((Phen)Gm)(BF)2·nCHCl3 were collected on a Bruker Apex DUO CCD diffractometer equipped with a graphite-monochromated MoKα (λ=0.71073 Å) radiation source at 150(2) K. The SMART software was used for data collection and also for indexing the reflections and determining the unit cell parameters; the collected data were integrated using SAINT software and corrected with SADABS for absorption. The structures were solved using SHELXTL software. Heteronuclear C–H correlations (HMBC and 1H–1H couplings) were used for the signal assignment. NMR spectra of the clathrochelates FeBd2((Phen)Gm)(BF)2 were not complete due to its low solubility.

Table 1. Crystal data and structure refinement for the iron(II) clathrochelates FeBd2((Phn)Gm)(BF)2 and FeBd2((Phen)Gm)(BF)2.

<table>
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<th>Parameter</th>
<th>FeBd2((Phn)Gm)(BF)2</th>
<th>FeBd2((Phen)Gm)(BF)2</th>
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<tr>
<td>Empirical formula</td>
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<td>C42H26B2F2FeN10O6</td>
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<td>12.0729(14)</td>
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<tr>
<td>b(Å)</td>
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<td>13.6088(16)</td>
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<td>V(Å³)</td>
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<td>2075.1(4)</td>
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<td>R1 = 0.0405, wR2 = 0.0908</td>
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<td>R indices (all data)</td>
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<td>R1 = 0.0574, wR2 = 0.0954</td>
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<td>Largest diff. peak and hole (e/Å³)</td>
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Synthesis

FeBd2((Phn)Gm)(BF)2, FeBd2((NH2)2Gm)(BF)2 complex (0.103 g, 0.15 mmol) and 1,10-phenanthroline-5,6-dione (0.318 g, 0.15 mmol) were dissolved/suspended in DMSO (15 ml) and DBU was added (0.4 ml). The reaction mixture was left for three weeks at r.t. and then was poured into brine. The precipitate formed was filtered off, dried in air and extracted with chloroform. The extract was chromatographically separated on silica gel (230–400 mesh, 1×20 cm column, eluent: chloroform). The green product was collected, the solvent was evaporated to dryness and the solid residue was dried in vacuo. Yield: 0.100 g (75 %). X-Ray quality single crystals of the complex FeBd2((Phn)Gm)(BF)2·n CHCl3, suitable for X-ray diffraction experiment, were obtained by recrystallization from chloroform–heptane 6:1 mixture. Found (%): C 56.38; H 3.14; N 15.79; F 4.40; B 2.42. Calculated for FeC38H28N8O6B2 (%): C 56.38; H 3.42; N 15.87; F 4.39; B 2.41.

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FeBd2((Phn)Gm)(BF)2. This complex was obtained similar to FeBd2((Phen)Gm)(BF)2 except Fe(Bd)2((NH2)2Gm)(BF)2 (0.112 g, 0.16 mmol), phenanthrene-9,10-dione (0.153 g, 0.17 mmol) and DBU (0.3 ml) were used. Yield: 0.110 g (80 %). X-Ray quality single crystals of the complex FeBd2((Phn)Gm)(BF)2·CHCl3 were obtained by recrystallization from chloroform–heptane 6:1 v/v mixture.
Found (%): C 59.92; H 3.30; N 12.66; B 2.42; F 4.27. Calculated for FeC_{40}H_{30}N_{6}O_{6}B_{2}F_{2} (%): C 60.04; H 3.21; N 12.73; B 2.46; F 4.32. 1H NMR (CDCl_{3}) δ ppm: 7.28 (d, J = 8 Hz, o-Ph), 7.34 (m, 12H (m-Ph+p-Ph), 7.63 (td, 2H, J_{1}=7.6 Hz, J_{2}=0.9 Hz (H-8+H-15)), 7.79 (td, 2H, J_{1}=7.6 Hz, J_{2}=1.2 Hz (H-9+H-14)), 8.36 (br.d, 2H, J=8.0 Hz (H-10+H-13)), 9.23 (dd, 2H, J_{1}=8.0 Hz, J_{2}=1.2 Hz (H-7+H-16)). Cyclic Voltammetry

Cyclic voltammetry (CV) was performed using a Metrohm 797 VA Computrace instrument with a glassy carbon (GC) and platinum working electrodes and a saturated Ag/AgCl in 3.5 M KCl as a reference electrode. Tetra-n-butylammonium perchlorate in chloroform (0.1 M) was used as the supporting electrolyte. Solutions of the samples in the electrolyte (1–2 mM) were degassed by purging with argon prior to the CV measurements.

Results and Discussion

Condensation of FeBd_{2}((NH_{2})_{2}Gm)(BF)_{2} with 1,10-phenanthroline-5,6-dione or phenanthrene-9,10-dione in the presence of 1,8-diazabicyclo[5.4.0]undecene-7 (DBU) as a strong organic base in aprotic polar DMSO media afforded the target macrobicyclic complexes FeBd_{2}((Phen)Gm)(BF)_{2}, the derivative of phenanthroline, and FeBd_{2}((Phn)Gm)(BF)_{2}, the derivative of phenanthrene, in high yields (Scheme 4).

These reactions proceeded via electrophilic attack of the corresponding dicarbonyl component (i.e. phenanthro- linedione or phenanthrenedione) at the deprotonated amino groups of the macrobicyclic precursor followed by elimination of water molecules. In the course of this addition – elimination reaction, the initial red reaction mixture changed color to grayish-crimson thus allowing its naked-eye control.

The single crystal X-ray structures of the clathrochelate products FeBd_{2}((Phen)Gm)(BF)_{2} and FeBd_{2}((Phn)Gm)(BF)_{2} are shown in Figure 1. They confirmed that the reaction of FeBd_{2}((NH_{2})_{2}Gm)(BF)_{2} with the above α-quinones gave the target clathrochelates with annulated 1,4,8,9-tetraazatriphenylene or 1,4-diazatriphenylene fragment. The intermediate trigonal prismatic (TP)–trigonal antiprismonic (TAP) geometry of their cage frameworks is only slightly affected by the formation of these annulated fragments. The heights of both TP–TAP FeN_{6}-coordination polyhedra is approximately 2.37 Å; their distortion angles φ are equal to 21.6° and 23.4°, respectively (φ = 0° for an ideal TP, φ = 60° for an ideal TAP). Fe–N distances fall in a narrow range 1.90–1.94 Å for both these cage complexes. The terminal phenanthroline fragment of the annulated tetraazatriphenylene heterocyclic substituent of FeBd_{2}((Phen)Gm)(BF)_{2} is structurally quite similar to 1,4,8,9-tetraazatriphenylene-2,3-dicarbonitrile.[13]

On the other hand, the pyrazine fragment of the molecule FeBd_{2}((Phn)Gm)(BF)_{2} has some touch of quinoid alternation with C_{6}N_{4} bond lengths close to 1.35 Å and N–C het distances of approximately 1.32 Å. Similarly, the carbocyclic (phenanthrene) part of FeBd_{2}((Phn)Gm)(BF)_{2} was found to be structurally very similar to 2,3-dicyanodibenzo(f,h) quinoxaline.[14] Its pyrazine moiety also has a quinoid character and the same C_{6}N_{4} and N–C_{het} bond lengths of 1.35 Å and 1.32 Å, respectively.

The crystals of these cage complexes contain the clathrochelate dimers formed via stacking interactions shown in Figure 2.

In the FeBd_{2}((Phen)Gm)(BF)_{2}, crystal, the heterocylic fragments of their macrobicyclic entities are almost parallel. These macrobicyclic molecules are related by the
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Figure 1. General views of the clathrochelates FeBd₂(Phen)Gm(BF)₂ (top) and FeBd₂(Phn)Gm(BF)₂ (bottom).

Figure 2. Side view of the clathrochelate dimers formed by the complexes FeBd₂((Phen)Gm)(BF)₂ (top) and FeBd₂((Phn)Gm)(BF)₂ (bottom). Semi-transparent spheres denote van der Waals radii of the non-hydrogen atoms of their annulated ribbed fragments. Hydrogen atoms are omitted for clarity.

Figure 3. Perpendicular views of the clathrochelate dimers, formed by the cage complexes FeBd₂((Phen)Gm)(BF)₂ (top) and FeBd₂((Phn)Gm)(BF)₂ (bottom). Hydrogen atoms are omitted for clarity.

Figure 4. CV of the chloroform solutions of FeBd₂((Phn)Gm)(BF)₂ (1 GC, 2 Pt) and FeBd₂((Phen)Gm)(BF)₂ (3 GC, 4 Pt) at scan rate 50 mV·s⁻¹.

Crystallographic inversion center; the deviations of their atoms from the corresponding mean plane do not exceed 0.0025 Å. The distance between the mean planes of these interacting fragments is approximately 3.34 Å. In the crystal of the FeBd₂((Phn)Gm)(BF)₂ complex, its annulated fragment is more distorted with the deviations reaching 0.1 Å, and the distance between the above mean planes in its clathrochelate dimer being even slightly smaller (3.32 Å); overlapping between the heterocyclic fragments of the corresponding macrobicyclic entities is shown in Figure 3. Such a short interplane separation and rather small slippage (the distances between the centroids of their annulated
heterocyclic systems fall in the range 3.5–3.6 Å) results in the substantial overlapping causing a strong stabilization of these supramolecular assemblies. Theoretical estimations of dimerization energy for triphenylene (approximately 10 kcal/mol) suggest the same order of magnitude for that of the above clathrochelate dimers.

Such dimerization may be a reason of the observed poor solubility of these cage complexes in most common organic solvents estimated to be in the range 10⁻⁵–10⁻⁶ mol/L, while the typical values for other cage iron(II) tris-dioximates of this type are higher by approximately two orders of magnitude. Only in the case of chloroform as a solvent, the solubility of the clathrochelates obtained approaches to 10⁻³ mol/L.

CVs of FeBd₂(Phn)(Gm)(BF₂) and FeBd₂(Phen)(Gm)(BF₂) (Figure 4) show one reduction wave with $E_{1/2}$ of 70 mV and 190 mV, respectively. The CV curves registered of Pt and GC are practically identical in shape and position. The symmetric shapes of this curves and the little drift of the peak potentials with scan rate (within 50–250 mV·s⁻¹) suggest the equilibrium character of these electrochemical redox processes.

The green-bluish color of chloroform solutions of the clathrochelates FeBd₂(Phen)(Gm)(BF₂) and FeBd₂(Phn)(Gm)(BF₂), is caused by the presence of more intensive band with the maximum at approximately 635 nm (for both of these complexes) and less intensive band with maxima at approximately 430 and 440 nm, respectively, in their UV-Vis spectra (Figure 5).

We also succeeded in the registration of UV-Vis spectrum of the solution of FeBd₂(Phen)(Gm)(BF₂) in electrochemically inert acetonitrile; its poor solubility in this solvent was compensated both by high molar extinction coefficient of this cage complex and by long optical path (10 cm cuvette). The initial UV-Vis spectrum of the clathrochelate FeBd₂(Phen)(Gm)(BF₂), and its changes during reduction of this cage complex with metallic lithium are shown in Figure 6. The green-bluish color of this acetonitrile solution is determined by two broad bands with maxima at approximately 410 and 640 nm. The above reduction caused the appearance of new band at approximately 500 nm and the disappearance of the absorption bands of the initial cage complex. So, the reduction resulted in the formation of the red solution. The absence of proper isosbestic points can be explained by the low concentration of the initial complex comparable with that of the trace contaminants remaining in the solvent. The spectrum of the reduced macrocyclic derivative of FeBd₂(Phen)(Gm)(BF₂) is quite similar to that of the anion-radical derivative of the quinoxaline-containing clathrochelate analog, thus suggesting chemical one-electron ligand-centered reduction of FeBd₂(Phen)(Gm)(BF₂).

The solubility of FeBd₂(Phen)(Gm)(BF₂) in acetonitrile was found to be even lower than that of FeBd₂(Phen)(Gm)(BF₂), and we were unable to register its UV-Vis spectrum in proper quality.

Conclusions

Thus, we have demonstrated that electrophilic addition can be successfully used for preparation of the iron(II) tris-dioximate clathrochelates bearing extended redox-active annulated ribbed substituents. The study of donor properties of the phenanthroline-terminated clathrochelate FeBd₂(Phen)(Gm)(BF₂), as a potent chelating macrocyclic ligand is in progress in our group.

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