Dirhenium Coordination Complex Endowed with an Intrinsically Chiral Helical-Shaped Diphosphine Oxide

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ABSTRACT: A one-pot, multicomponent strategy was used to synthesize the first example of the dirhenium carbonyl coordination complex 2, in which the two metal atoms are connected through a chiral helical-shaped diphosphine oxide. Thanks to the flexibility of the helix of helicene 1, complex 2 was isolated in quite a good yield as a stable compound. It was characterized by analytical and spectroscopic techniques as well as by single-crystal X-ray analysis, which confirmed the chemical structure and the peculiar architecture of 2. In addition, computational studies were in agreement with the transitions observed in the experimental UV–vis spectrum, revealing the presence of two bands with maxima at about 520 (metal-to-ligand charge transfer) and 400 nm (IL).

INTRODUCTION

The helicene-based transition metal complexes are an extremely appealing class of chiral molecules, in which the coordination of metals to π-helical ligand bearing appropriate coordinating functionalities allows the construction of uncommon chiral architectures. Indeed, helicenes are ortho-annulated polycyclic aromatic compounds endowed with an inherently chiral π-conjugated system which have attracted considerable attention because of their unique physicochemical properties and potential applications in different areas of science. On the other hand, reacting π-helicene scaffolds with metals can significantly modify specific properties, obtaining a structural variety that however retains the defining features of the helical structure. In this context, tetrathia[7]helicenes (7-TH), in which the thiophene rings are fused to alternating benzene rings to form a configurationally stable helix, are emerging as the most promising systems, also thanks to the reliable functionalization of the terminal thiophene rings of the 7-TH scaffold that allows the introduction of a diversity of functional groups, including those with an efficient coordinating ability (e.g., cyano and phosphane). Thus, the 7-TH nitrile derivatives were used as monodentate ligands in the synthesis of the corresponding Ru(II) and Fe(II) complexes that have been shown to be interesting candidates for optoelectronic purposes. More recently, Rh(I) and Au(I) complexes based on 7-TH phosphanes or phosphinohelicenes have been successfully employed in the asymmetric organometallic catalysis. It should be noted that the unique flexibility of the 7-TH skeleton ensures the coordination with different metal centers, affording chelated Rh(I) complexes or dinuclear Au(I) complexes formed through the coordination of two phosphorus atoms of 7-TH diphosphanes to a rhodium center or two gold centers, respectively. Among several helicene-based coordination complexes reported so far, only one rhenium complex has been described in the literature. Moreover, enantiopure chiral rhenium complexes are very scarce, and some of them are chiral at the rhenium center, whereas in other cases, a chiral ligand is present. The synthesis of some of these complexes is based on metal-direct self-assembly, which affords supramolecular coordination complexes, whose functionalities such as enantioselective recognition, catalysis, and sensing are improved with respect to the organic counterpart. In this framework, the assembly of fac-Re(CO)₅ units with oxygen anionic donors, or chalcogenide anions, and neutral bridging nitrogen ligand through the direct orthogonal bonding approach represents an example of the synthesis of rhenium-based supramolecular coordination complexes. This one-pot multicomponent approach has also been used for the synthesis of dinuclear rhenium complexes containing the hard ditopic O=P donors as the bridging ligand. However, the rhenium-based polynuclear complexes containing O=P ligands are very scarce, based on tetraphenylidichalcogenimidodiphosphinate anions as ligands, and, more recently, flexible alkyl or benzyl diphosphine oxides as neutral ditopic O=P ligands.

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along with a rigid bischelating O donor (e.g., chloranilic acid and 2,5-dihydroxy-1,4-benzoquinone).32,33 In our continuing studies on the synthesis and characterization of organometallic complexes based on the 7-TH derivatives, we focused on the study of the coordination ability of an uncommon and more rigid helical-based phosphine oxide as a neutral ditopic ligand toward dinuclear rhenium complexes containing chloranilic acid as the anionic donor. Herein, we report the first example of a rhenium(I)-based coordination complex in which 7-TH phosphate oxide (±)-19, used as a racemic mixture for this first study, has been selected as a model helical bis(P=O) donor ligand.

## RESULTS AND DISCUSSION

The dinuclear rhenium(I) complex 2 was synthesized by refluxing Re2(CO)10, the racemic mixture of the /π/-helical phosphate oxide (±)-1, and chloranilic acid (H2-CA), as anionic donor, in mesitylene for 24 h (Scheme 1).

### Scheme 1. Synthesis of Dinuclear Rhenium(I) Complex 2

![Scheme 1: Synthesis of Dinuclear Rhenium(I) Complex 2](image)

This one-pot procedure involves the oxidative addition of the chloranilate anion to [Re2(CO)10], with the reduction of H+ to H2, and dissociation of one CO ligand for each metal center. Then, the further substitution of other two CO groups by the bridging phosphate-oxide ligand 1 affords complex 2. Chromatographic purification of the reaction mixture provided complex 2 in a 79% yield as an air-stable brown-red solid. It is important to note that 7-TH are known to be configurationally stable heterohelicenes, with a racemization barrier, for example, for the unsubstituted 7-TH of 38.7 kcal/mol at 227 °C.36 Because the temperature of the reaction used for the synthesis of complex 2 is quite lower, this could open the possibility to synthesize the optically active rhenium complex 2, starting from the corresponding enantiopure phosphate oxide 1. In fact, the racemization barrier of ligand 1 should be significantly higher than that of the unsubstituted 7-TH because of the presence of two bulky diphenylphosphine oxide groups on the two terminal thieryl rings.

Contrary to other dinuclear rhenium complexes reported by some of us,37 complex 2 displayed, in the (CO) region of the infrared (IR) spectrum, only three bands at 2020, 1912, and 1890 cm⁻¹ instead of the typical four-band pattern. This feature has been already observed for the analogous complexes containing the same chloranilate anion as the ancillary ligand,32,33 in agreement with the modulation of the position of the carbonyl stretching bands by the nature of the ancillary ligand. The 31P{1H} NMR spectrum of 2 (Figure S3 in the Supporting Information) shows only one singlet at δ +42.2 ppm, indicating a symmetrical bridging coordination of 1. The symmetrical and dinuclear structure of complex 2 was further confirmed by X-ray analysis. Single crystals suitable for the X-ray structural determination of 2 were obtained from the slow diffusion of hexane into toluene solution of complex 2. An Oak Ridge thermal ellipsoid plot (ORTEP) view of the complex is reported in Figure 1.

![Figure 1: ORTEP view of 2 with the atomic numbering scheme](image)

In complex 2, the ditopic 7-TH phosphate oxide ligand coordinates two Re(I) cations through its O1 and O2 oxygen atoms. The chloranilic (CA2⁻) unit acts as the bischelating ancillary ligand bridging the Re cations through its O3, O4 and O5, O6 oxygen atoms. The coordination geometry around each Re center is completed by three CO ligands and can be described as distorted octahedron.

Although the atoms of the C2O2 basal plane show very small tetrahedral distortion, the Re1−O1 and Re2−O2 bonds deviate significantly (by 12.1° and 15.2°, respectively) from the normal to the C2O2 plane. Moreover, the {[Re(CO)3][μ-CA][Re(CO)3]} unit deviates from planarity toward a boatlike conformation, with Re1 and Re2 placed at 0.18(1) and 0.39(1) Å, respectively, from the least-squares plane through the CA2⁻ ligand. Such distortions are to be related to the geometrical constraint imposed by the 7-TH ligand, as confirmed by comparison with the analogous dinuclear Re(I) complexes bearing the O=P(CH2)nP=O (n = 335 and 431) ditopic ligands in place of 1. The distance between the Re cations observed in 2, 8.059(11) Å, is slightly shorter than that reported for the complex with the butyl bridge, 8.08 Å,32 but significantly longer than that of the complex with the propyl bridge, 7.84 Å.32 Although in the former complex the {[Re(CO)3][μ-CA][Re(CO)3]} unit is almost planar, in the latter, it assumes a boatlike conformation similar to that observed in 2 but with the Re ions further away from the CA2⁻ plane. On the other side, the 7-TH ligand is as well expected to undergo a slight distortion after coordination to the metal centers. This is supported by the comparison of the P1···P2 distance in 2, 6.609(6) Å, with that of the previously reported 7-TH phosphonate derivative bearing two EtO groups at the phosphorus atoms in place of the two phenyl groups, measuring 5.879 Å. These results further demonstrate the peculiar flexible nature of the helical system of 7-TH.

The absorption spectra of complex 2 and of the 7-TH ligand 1, recorded in a dilute, air-equilibrated dichloromethane solution at room temperature, are shown in Figure 2.

The absorption spectrum of 2 shows two main absorption features. The broad and structureless low-energy absorption band at 514 nm (ca. 1.25 × 10⁻⁴ M⁻¹ cm⁻¹) has been attributed to
the metal-to-ligand $d(\text{Re}) \rightarrow \pi^* (\text{CA}^2\text{−})$ charge-transfer transition (MLCT). Actually, both the energy and the molar absorption coefficients closely resemble those reported for analogous dinuclear rhenium complexes containing phosphine oxide and the chloranilate anion as the ancillary ligand.\cite{32,33} The spectrum also shows another absorption band at a higher energy (maximum in the range 390−409 nm), with a slightly higher intensity ($ε$ ca. $2.10 \times 10^4$ M$^{-1}$ cm$^{-1}$). This band is very similar to that observed for the 7-TH ligand, and it has been attributed to the spin-allowed $\pi-\pi^*$ excitations of the helicene skeleton. In comparison to the 7-TH ligand, this band is only slightly red-shifted, confirming that this electronic transition does not involve the metal centers. Moreover, it displays a less pronounced vibronic structure. This feature could be attributed to the less coplanar geometry adopted by the helicene ligand inside the complex. The band assignment has been confirmed by time-dependent density functional theory (TDDFT) calculations on the DFT optimized complex. The computed spectrum of 2 in CH$_2$Cl$_2$ (see Figure S4 in the Supporting Information) well reproduces the observed one, revealing the presence of two bands with maxima at about 520 and 400 nm. The former is mainly due to the excitations at 527 nm (oscillator strength $f = 0.52$) and 480 nm ($f = 0.16$), corresponding to the highest occupied molecular orbital (HOMO) − 3 − lowest unoccupied molecular orbital (LUMO) (82%) and HOMO − 6 − LUMO (88%) transitions, respectively, where HOMO − 3 is delocalized on the $\{[\text{Re} (\text{CO})_3]\}$ units, and LUMO on the CA$^2$− moiety. The band at higher energy is determined by excitations at 412 ($f = 0.22$) and 396 nm ($f = 0.30$), mainly due to HOMO− LUMO + 1 (96%) and HOMO − 1 − LUMO + 1 (90%), respectively, where HOMO, HOMO − 1, and LUMO + 1 are all orbitals delocalized on the 7-TH ligand (see Figure S5 in the Supporting Information for a plot of the molecular orbitals mainly involved in the electronic transitions).

**CONCLUSIONS**

In conclusion, we have described the synthesis of the first dinuclear rhenium complex in which the two Re centers are connected by an unusual helical diphosphine oxide. This peculiar coordination architecture is possible thanks to the flexibility of the phosphorus-based helical ligand that, also in this case, is able to exhibit different dihedral angles of the helix depending on the steric demand of the surrounding portion of the molecule. Having successfully demonstrated the formation of complex 2, in view of the potential applications of this procedure to synthesize the enantiopure rhenium complexes based on the 7-TH phosphine oxides, some experiments could be performed to assess the configurational stability of these chiral bis(PO=O) donor ligands.

Noteworthy, the spectroscopic characterization of 2 points out a strong absorption maximum in the red portion of the visible spectrum, in line with that observed for the analogous complexes previously reported. This means that the helical system does not perturb the spectroscopic properties of the complex, and this could open the way to investigate the photophysical and chiroptical properties of the corresponding enantiopure complexes.

**EXPERIMENTAL SECTION**

**Materials and Methods.** All the reagents, including Re$_2$(CO)$_{10}$ and chloranilic acid (H$_2$-CA), were purchased from Aldrich and used as received. Phosphine oxide (±)-1 was synthesized as previously reported.\cite{31} All the reactions were performed under an inert N$_2$ atmosphere. All the solvents were deoxygenated and dried using standard methods before use; mesitylene was distilled on Na(s), whereas CH$_2$Cl$_2$ on P$_2$O$_5$. Commercial deuterated solvents were used as received. Column chromatography was performed using Alfa Aesar silica gel 60 (0.032−0.063 mm).$^1$H, $^{13}$C($^1$H), and $^{31}$P($^1$H) NMR spectra were recorded on a Bruker DRX-400 MHz instrument, equipped with a Bruker 5 mm BBI Z gradient probe head with a maximum gradient strength of 5.3 S G$^{-1}$ cm$^{-1}$. Chemical shifts are given as $δ$ (ppm) relative to the residual protonated solvent resonance ($^1$H: $δ = 5.32$ ppm, $^{13}$C: $δ = 53.84$ ppm for CD$_2$Cl$_2$). Multiplicities are given as singlet (s), doublet (d), or multiplet (m). The IR spectra in the solution were acquired on a Bruker Vector 22 FT spectrophotometer. The electronic absorption spectra were recorded on an Agilent model 8543 spectrophotometer at room temperature and using quartz cells with a 1.0 cm path length.

**Synthesis of Complex 2.** Re$_2$(CO)$_{10}$ (10.0 mg, 0.0153 mmol), H$_2$-CA (1.6 mg, 0.0077 mmol), and phosphine oxide (±)-1 (13.6 mg, 0.0153 mmol) were dissolved in 3 mL of anhydrous mesitylene and the reaction mixture, which was stirred under reflux for one day, slowly turning dark brown. The solution was then evaporated to dryness under a reduced pressure, and the dark red product was purified through column chromatography (toluene/ethyl acetate 8:2) yielding 10 mg (0.0061 mmol) of microcrystalline powder (yield 79%). IR (CH$_2$Cl$_2$) (CO): 2020 (s), 1912 (s), and 1905 (s) cm$^{-1}$. $^1$H NMR (CD$_2$Cl$_2$): 300 K, 400 MHz: δ 7.88−7.83 (m, 4H, Ar-H 7-TH), 7.59−7.43 (m, 8H, Ar-H –POPh$_2$), 7.26−7.04 (m, 12 H, Ar-H –POPh$_2$), 6.59 (d, 2H, $J = 9.2$ Hz, Th-H 7-TH), 3.15−2.99 (m, 4H, CH$_2$CH$_2$CH$_3$), 1.84−1.72 (m, 4H, CH$_2$CH$_2$CH$_3$), 1.10 (t, 6H, $J = 7.24$ Hz, CH$_3$CH$_2$CH$_3$), $^{13}$C NMR (CD$_2$Cl$_2$): 300 K, 100 MHz): δ 141.5 (Cq), 140.3 (Cq), 136.7 (Cq), 134.4 (Cq), 133.7 (d, $JC_P = 11.2$ Hz, 2CH$_3$thien), 131.5 (2 Cq), 131.2 (d, $JC_P = 10.5$ Hz 2CH), 130.5 (CH), 129.2 (d, $JC_P = 10.5$ Hz 1CH$_3$meta), 128.7 (d, $JC_P = 12.0$ Hz 2CH$_3$meta), 126.8 (Cq), 121.8 (CH), 121.0 (CH), 34.2 (CH$_3$), 23.2 (CH$_2$), 14.2 (CH$_3$). $^{31}$P NMR (CD$_2$Cl$_2$): 300 K, 161 MHz): δ 42.2 (s). Elemental Anal. Calcd for Cs$_6$H$_{10}$Cl$_2$O$_{12}$P$_2$Re$_2$: C, 47.03; H, 2.47. Found: C, 47.15; H, 2.53.

**Single-Crystal X-ray Diffraction Studies on Complex 2.** Very thin dark red single crystals of 2 suitable for X-ray diffraction have been obtained by recrystallization from toluene. Repeated recrystallization attempts did not allow to improve the
quality of the crystals, which was in all cases very poor. The intensity of the crystals were collected on a Bruker Smart Apex CCD area detector (Bruker AXS Inc., Madison, WI, US) using the graphite-monochromated Mo Kα radiation ($\lambda = 0.71073$ Å). Data reduction was made with SAINT, and the absorption corrections based on multiscan were obtained by SADABS.\textsuperscript{38} The structure was solved by SHELXS-97\textsuperscript{39} and refined on F2 by full-matrix least-squares using SHELXL-14.\textsuperscript{40} Data collection was performed at low temperature (120 K) to enhance the diffracted intensities and increase the extent of the observed data. A quite low data/parameters ratio prevented to get a meaningful description of the thermal parameters, and, for this reason, several restraints (DELU and ISOR) have been applied on the heavy atoms of the more disordered groups. Moreover, the large void present in the crystal structure suggests the presence of disordered co-crystallized solvent molecule(s), whose structure, however, could not be determined owing to the low resolution of the X-ray data.

Hydrogen atoms were placed in the calculated positions and included in the refinement as “riding”. The H isotropic thermal parameters were fixed at 1.2 (1.5 for methyl groups) times the equivalent thermal parameter of the atoms to which they are bonded. The program ORTEPIII was used for graphics.\textsuperscript{47} The crystallographic data were deposited with the Cambridge Crystallographic Data Centre under the Cambridge Crystallographic Data Centre Centre under the supplementary publication no. CCDC 1844855. These data can be obtained free of charge via www.ccdc.cam.ac.uk/conts/retrieving.html (or from CCDC, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44-1223-336-033; e-mail: deposit@ccdc.cam.ac.uk).

**Theoretical Studies.** Geometry optimization of isolated, gas-phase complex 2 was performed by the Gaussian16 program package,\textsuperscript{42} using the def2-SVP\textsuperscript{43,44} basis set and the M06 functional\textsuperscript{45} because of its specific parametrization on organometallic complexes. Rhenium has been treated as a 15-electron system, with relativistic effective core potential taken from the literature.\textsuperscript{46} The X-ray molecular structure of 2 was used as the starting point for geometry optimization. The electronic absorption spectrum was computed in dichloromethane on the optimized structure using the same basis set and the PBE0 functional,\textsuperscript{47} owing to its optimal performance in reproducing the absorption spectra of organic\textsuperscript{48} and organometallic compounds.\textsuperscript{49} Solvent effects (CH$_2$Cl$_2$) have been taken into account in all calculations by the polarizable continuum model using the integral equation formalism variant.\textsuperscript{50}

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