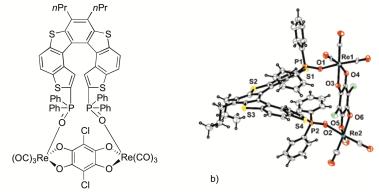
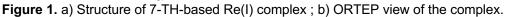
Synthesis and characterization of a tetrathia[7]helicene-based rhenium(I) complex

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Tetrathia[7]helicenes (7-TH), formed by thiophene and benzene rings *ortho*-fused in an alternating fashion, are emerging as one of the most popular class of chiral helical-shaped molecules, thanks to their peculiar electronic and chiroptical properties suitable for manifold applications in different areas of science.¹ In particular, transition metal-based 7-TH systems are an extremely appealing class of complexes, in which the coordination of metals with the π -helical ligand, bearing appropriate coordinating functionalities, provides original chiral architectures. Indeed, the effective functionalization of the α -position(s) of the terminal thiophene ring(s) of the 7-TH scaffold allows the introduction of a variety of substituents, including those with efficient coordinating ability (*e.g.* cyano², phosphane³, phosphine oxide⁴). For example, Rh(I)⁵ and Au(I)⁶ complexes based on 7-TH phosphanes have been successfully used in the homogenous transition metal catalysis. In our ongoing studies on 7-TH-based organometallic complexes, we have focused on a novel field of investigation concerning the development of rhenium-based polynuclear complexes containing 7-TH phosphine oxide ligands. In this communication, we describe the synthesis and the characterization of a novel dinuclear rhenium(I) complex (Figure 1), along with the elucidation of its tridimensional structure by single crystal X-ray diffraction studies.





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