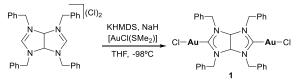
Supramolecular Self-Assembly Structures from a **Di-Au(I)** Complex with a Hinge-Shaped Ligand

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Key Words: Supramolecular • N-heterocyclic • Gold • Hinge • Self-assembly

Among N-heterocyclic carbenes (NHCs), polytopic NHC ligands have emerged as tools in promising the design of supramolecular structures.¹ In this regard, our group of research designed fusedbis(imidazolinylidene) ligands with folded geometry and reported their coordination to Rh and Ir,²⁻³ and Pd.³ More recently, one of these hinge-shaped ligands allowed us to prepare the Au(I)-based dimetallic complex 1 (Scheme 1).

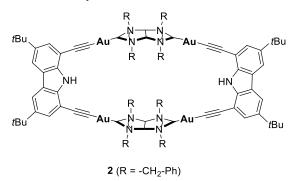


Scheme 1. Synthesis of di-Au(I) complex 1

The combination of the folded geometry of the ligand with a coinage metal, extensively employed in the synthesis of supramolecular assemblies,⁴⁻⁵ makes complex **1** a perfect synthon for the preparation of metallosupramolecular structures by means of the metal-directed self-assembly approach. This strategy allows the generation of sophisticated molecular architectures from relatively easy materials.6

With this in mind and taking into account that Au(I) complexes are prone to form stable linear compounds with acetylides, we reacted complex 1 with 3,6-di(*tert*-butyl)-1,8diethynyl-9H-carbazole in the presence of a base. This reaction allowed the preparation of the supramolecular complex 2 (Scheme 2).

The synthesis of other Au(I) supramolecular complexes with inner cavities of different size by means of the combination of 1 with the appropriate alkyne, is currently underway in our laboratory.



Scheme 2. Tetra-Au(I) complex 2

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