

## **Bis(phosphane)copper(I) and silver(I) dithiocarbamates: crystallography and anti-microbial assay**

### **ABSTRACT**

The crystal and molecular structures of  $(\text{Ph}_3\text{P})_2\text{M}[\text{S}_2\text{CN}(\text{Me})\text{CH}_2\text{CH}_2\text{OH}]$ ,  $\text{M}=\text{Cu}$ , isolated as a 1:1 dichloromethane solvate ( $1 \cdot \text{CH}_2\text{Cl}_2$ ), and  $\text{M}=\text{Ag}$  (**4**) show the central metal atom to be coordinated by a symmetrically ( $1 \cdot \text{CH}_2\text{Cl}_2$ ) and asymmetrically chelating (**4**) dithiocarbamate ligand. The distorted tetrahedral geometries are completed by two  $\text{PPh}_3$  ligands. The presence of hydroxyl— $\cdots\text{S}$ (dithiocarbamate) hydrogen bonds leads to centrosymmetric dimeric aggregates in each crystal structure. In the molecular packing of  $1 \cdot \text{CH}_2\text{Cl}_2$ , channels comprising **1** are formed via aryl-C—H $\cdots$ O interactions with the solvent molecules associated with the walls of the channels via methylene-C—H $\cdots$ S,  $\pi$ (aryl) interactions. For **4**, the dimeric aggregates are connected via a network of aryl-C—H $\cdots$  $\pi$ (aryl) interactions. Preliminary screening for anti-microbial activity was conducted. The compounds were only potent against Gram-positive bacteria. Some further selectivity in activity was noted. Most notably, all compounds were active against methicillin resistant *Staphylococcus aureus*.

**Keyword:** Copper(I); Crystal structure analysis; Dithiocarbamate; Silver(I); X-ray diffraction