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Biomimetic polyimidazole complexes: A thermoanalytical study of Co(II)-, Ni(II)- and Cu(II)-bis(imidazol-2-yl)methane complexes

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Abstract

Imidazole rings of histidine residues often form part of the metal-binding site in metalloproteins, so ligands with the 2-imidazolyl units are good models for mimicking biological metal-binding sites because the histidine imidazole ring is often attached via the 2-position to the side chain when the imidazole is coordinated to a metal. Thermoanalytical studies on biomimetic molecules are useful to complete the informations on the mechanism of action of the metal-binding sites of metalloproteins and to relate their chemico-physical properties. The synthesis, the spectroscopic characterization and the thermoanalytical study of bis(imidazol-2-yl)methane (BIM) complexes with divalent cobalt, nickel and copper, with a general formula M(BIM)₂Cl₂, are reported: the thermal stability and the decomposition steps were determined by thermogravimetry (TG), derivative thermogravimetry (DTG) and differential scanning calorimetry (DSC). The released products, due to the thermal decomposition, were analysed by on-line coupling a FTIR spectrometer to the thermobalance; the so obtained evolved gas analysis (EGA) allowed to prove the proposed decomposition steps.

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1. Introduction

Copper metalloproteins are involved in various biological functions such as electron transfer, oxygen transport, or substrate oxidation [1-5]. These functions are the consequence of the redox properties of the copper ion modulated by the protein ligands and of the selection of substrates by the active site.

Imidazole rings of histidine residues form part of the metal-binding site in metalloproteins [6–8]. The ubiquitous histidine ligation in metalloenzymes has stimulated syntheses of imidazole-containing multidentate ligands for biomimetic studies. Many bi-, tri- and tetradentate metal chelating ligands containing substituted imidazole have been developed [9,10]: ligands with the 2-imidazolyl units are good models for mimicking biological metal-binding sites because the histidine imidazole ring is often attached via the 2-position to the side chain when the imidazole is coordinated to a metal.

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In this view, thermoanalytical studies on biomimetic molecules are useful to complete the information on the mechanism of action of the metal-binding sites of metalloproteins and to relate their physico-chemical properties [11–17].

Bis(imidazol-2-yl)methane (BIM) is the simplest polyimidazole ligand useful to model multihistidine coordination, as also reported by Place et al. [18].



Bis(imidazol-2-yl)Methane

(BIM)

In this work, the synthesis of the Cu(BIM)₂Cl₂ complex is reported. Its thermal stability and the decomposition mechanisms were studied by thermogravimetric analysis (TGA), derivative thermogravimetry (DTG) and differential scanning calorimetry (DSC) and compared to

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cobalt and nickel complexes synthesized with the same procedure.

For each TG process, the released products, due to the thermal decomposition, were analysed by on-line coupling a FTIR spectrometer to the thermobalance; the so obtained evolved gas analysis (EGA) [19,20] allowed to prove the proposed decomposition steps.

The results are discussed on the basis of the different instrumental characterizations reported in the literature.

2. Experimental

2.1. Syntheses of the complexes

All the complexes of general formula $M(BIM)_2Cl_2$ (where M = Co(II), Ni(II) or Cu(II)) were synthesized by following the procedure reported in literature [18] with few modifications.

BIM was obtained as reported by Collman et al. [21]. The metal chloride salts were purchased from Sigma–Aldrich.

In detail, a 20 mL methanol solution of 1058 g of BIM (7.14 mmol) was added to 3.57 mmol methanol solutions of cobalt(II)-, nickel(II)- or copper(II)-chloride. After few minutes of stirring, the complexes spontaneously precipitated to give powders of $M(BIM)_2Cl_2$ (where M = Co(II), Ni(II) or Cu(II)).

2.1.1. Elemental analysis (metal by ICP-OES)

Co(BIM)₂Cl₂: C 39.4% (39.4), H 3.5% (3.7), N 26.3% (26.2), Co 13.90% (13.8); Ni(BIM)₂Cl₂: C 39.5% (39.4), H 3.7% (3.7), N 26.9% (26.2), Ni 14.10% (13.8); Cu(BIM)₂Cl₂ C 39.0% (39.0), H 3.8% (3.7), N 26.3% (26.0), Cu 15.00% (14.7).

2.2. Instrumental

UV-vis spectra were recorded by using a Perkin Elmer Lambda 11 spectrophotometer.

The thermoanalytical curves were obtained by using a Perkin Elmer TGA7 thermobalance (range 20-1000 °C) and a Perkin Elmer DSC7 calorimeter; the atmosphere was either pure nitrogen or air, at a flow rate of 100 mL min^{-1} ; the heating rate was varied between 5 and 40 °C min^{-1} , with the best resolution achieved at a scanning rate of 10 °C min^{-1} .

To obtain the IR spectra of the gases evolved during the thermogravimetric analysis, the thermobalance was coupled with a Perkin Elmer FTIR spectrometer, model 1760X. The TGA7 was linked to the heated gas cell of the FTIR instrument by means of a heated transfer line, the temperatures of the cell and of the transfer line being independently selected.

3. Results and discussion

The synthesis of BIM is reported in two articles by Joseph et al. [22] and Collman et al. [21]. The complexes were initially synthesized by both the procedures to determine the differences: since the yield resulted very similar, the second approach was preferred and is so cited in Section 2.

The complexes present a weak absorption in the visible region, with a large Gaussian shape centred at 600–615 nm.

A similar absorption is also reported for the complex $Cu(BIM)_2(ClO_4)_2$ [18].

To verify the structure of the precipitated new $Cu(BIM)_2Cl_2$, it was solved by X-ray crystal structure to be compared with $Cu(BIM)_2(CIO_4)_2$ compound [18]. To this end, a diluted methanol solution was slowly evaporated (usually more than 2 days) resulting in square crystals. Both the chloride and the perchlorate complexes show the copper(II) ion in a square plane



Fig. 1. TG (—) and DTG (-----) curves of Co(BIM)₂Cl₂ (a), Ni(BIM)₂Cl₂ (b) and Cu(BIM)₂Cl₂ (c) complexes. Heating rate: $10 \,^{\circ}\text{C}\,\text{min}^{-1}$. Air flow at 100 mL min⁻¹ rate.

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