HYDROGENOTUNGSTATO COMPLEXES WITH TIN HALOGENURES (IV): INFRARED AND MÖSSBAUER STUDIES

DINORA DE BARROS¹, MOHAMADOU SEMBENE BOYE*², LIBASSE DIOP¹

¹Department of Chemistry, Faculty of Science and Technology, Laboratory of Mineral and Analytical Chemistry (LACHIMIA), Cheikh Anta DIOP University, Dakar, Senegal

²Department of Physics Chemistry, Faculty of Sciences and Technologies of Education and Training, Cheikh Anta Diop University, Boulevard Habib Bourguiba, 5036, Fann-Dakar, Senegal

Abstract: Six new hydrogenotungstato adducts and complexes have been synthesized and studied by infrared and Mössbauer techniques. The structures are mainly discrete, the anion behaving as a bidentate -one case- or a monodentate ligand in most cases. When extra hydrogen bonds are considered, supramolecular architectures may be obtained.

Keywords: discrete structures, hydrogenotungstato tin (IV) halides adducts, infrared, monodentate hydrogenotungstate ligand, Mössbauer, supramolecular architectures

1. INTRODUCTION

SnX₄ (X = Cl, Br, I) adducts were studied by Beattie and Rule [1, 2] and by Ohkaku and Nakamoto [3]. If the behavior of sulfate, selenate, oxalate and chromate ions [4–8] has already been the subject of several studies in the context of their association with tin (IV) compounds, organo- and halostannic molybdato and tungstato compounds have been the subject of less research [9] to our knowledge. Within the framework of these studies initiated for several years in our laboratory we were interested in the study of interactions between the quaternary ammonium salts of WO₄H₂ and the tin tetrahalides in order to obtain tin (IV) compounds with interesting biological activity. We have thus been able to isolate six new compounds of which infrared and Mössbauer spectroscopic studies have been carried out and presented here.

2. EXPERIMENTAL SETUP

2.1. Synthesis of tungstic acid salts

(Me₄N)₂WO₄Me₄NHWO₄.2H₂O (L₁) and (Et₄N)₂WO₄Et₄NHWO₄.4H₂O (L₂) used for synthesis of the studied complexes have been obtained by reacting H₂WO₄ acid 97 % (Merck product) with tetramethylyl- (10 % water solution Merck product) or tetraethylammonium hydroxide (20 % water solution Merck product) in 1:2 ratio. The solutions have been evaporated in an oven (60 °C) giving powders which are recrystallized from ethanol, washed with ether and then maintained under P₂O₅ in a desiccator.

2.2. Complexes synthesis

On allowing ethanolic solutions of (L₁) and SnX₄ (X = Cl, Br) in 2:3 ratio (A, B, are obtained) while (L₂) in ethanol reacting with SnX₄ (X = Cl, Br) in 2:3 or 1:2 ratio gives (C, D, E and F). The various complexes were obtained as white chloro or yellow bromo precipitates stirred for around two hours, filtered and washed with hot

* Corresponding author, email: mouhasboye@hotmail.com
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ethanol. The elemental analyses are reported in Table 1 with the various mixture stoichiometries (SnX₄/salt) M-L.

The analytical data [\% calculated (\% found)], have allowed to suggest the following formulae (Table 1).

A = Me₂NHWO₄SnBr₄  
B = Me₂NHWO₄SnCl₄NMe₄  
C = Et₃NHWO₄SnCl₄NEt₄  
D = Et₃NHWO₄SnBr₄NEt₄  
E = (Et₃NHWO₄)₂SnCl₄HWO₄  
F = (Et₃NHWO₄)₂SnBr₄HWO₄

| Compounds | M-L | % C | % H | % N | % X |
|-----------|-----|-----|-----|-----|-----|
| (L₂)      | –   | 19.65 (19.90) | 5.19 (5.35) | 5.73 (5.74) | –   |
| (L₄)      | –   | 30.04 (29.82) | 7.09 (7.22) | 4.38 (4.32) | –   |
| A         | 3-2 | 9.71 (9.98)   | 3.28 (3.14) | 1.80 (1.90) | 42.02 (42.44) |
| B         | 3-2 | 13.85 (13.69) | 3.60 (3.64) | 4.04 (4.01) | 25.61 (26.19) |
| C         | 3-2 | 23.85 (23.50) | 4.96 (4.91) | 3.47 (3.40) | 22.05 (22.10) |
| D         | 3-2 | 18.68 (18.41) | 3.85 (3.67) | 3.72 (3.60) | 38.93 (38.50) |
| E         | 1-2 | 16.84 (16.51) | 3.66 (3.71) | 2.46 (2.51) | 10.67 (11.16) |
| F         | 1-2 | 14.06 (13.95) | 3.15 (3.05) | 2.05 (2.01) | 17.56 (18.21) |

M-L = metal-ligand

The elemental analyses have been performed at the Service Central d’Analyses-CNRS- Vernaison-France or at the Microanalyses Laboratory of the University of Padua (Italy). The infrared spectra were recorded at the University of Padova (Italy) at room temperature using a Perkin Elmer spectrophotometer (4000-200 cm⁻¹) and a Nicolet 5300 type Fourier transform infrared spectrophotometer (600-50 cm⁻¹) as a suspension of the powders in Nujol. The faces used are cesium iodide or polyethylene for low frequencies. Infrared abbreviations: (vs) very strong; (s) strong; (m) medium; (vw) very weak; (w) weak; (sh) shoulder. Mössbauer spectra were recorded using a source of Ca¹¹⁹SnO₂ in constant acceleration; the sample being maintained between 80 K and 100 K. An accurate program based on the least square’s method allows to find by calculation the experimental spectrum and to make a deduction of the different parameters (δ: isomeric shift; ΔE: quadrupole splitting; Γ: full width at half-height).

3. RESULTS AND DISCUSSION

In Tables 2 and 3, the main infrared and Mössbauer data of the six compounds are reported (the spectral quality did not allow to locate ν OH which appears very weakly in infrared when the –OH group is not involved in a hydrogen bond).

| Attributions → | νWO₄ | δWO₄ | νₛM-X | νM-O |
|----------------|------|------|-------|------|
| Compounds ↓    |      |      |       |      |
| A              | 982 m| 890 sh| 815 s | 586 m| 444 m| 355 m| 210 vs| 280 w |
| B              | 980 m| 890 m| 815 s | 650 sh| 580 m| 445 w| 395 sh| 350 sh| 300 vs| –   |
| C              | 983 s| 891 w| 814 vs| 584 m| 445 m| 295 vs| –   |       |
| D              | 983 s| 880 w| 812 vs| 584 m| 445 m| 370 w| 350 w| 280 w | 210 vs| 280 w |
Table 3. Mössbauer parameters of complexes

| Compounds | A | B | C | D | E | F |
|-----------|---|---|---|---|---|---|
| δ (mm s\(^{-1}\)) | 0.52 | 0.37 | 0.37 | 0.58 | 0.13 | 0.21 |
| ΔE (mm s\(^{-1}\)) | 0.66 | 0.44 | 0.44 | 0.54 | 0.56 | 0.60 |
| Γ (mm s\(^{-1}\)) | 0.99 | 0.97 | 0.97 | 0.99 | 0.94 | 1.05 |

For Me\(_4\)NHWO\(_4\)SnBr\(_4\) (A), the important quadrupole splitting of SnBr\(_4\) and the sharpness of the band assigned to νSnBr\(_4\) (vibration Eu type according to the Group Theory) allow to conclude to a SnBr\(_4\) molecule of D\(_{4h}\) symmetry (transcoordination and centrosymmetry according to Group Theory): the proposed structure is an infinite chain with transcoordinated SnBr\(_4\) molecules, the hydrogenotungstate being a bi-unidentate ligand (Figure 1).

For R\(_4\)NHWO\(_4\)SnX\(_5\)NR\(_4\) (B, C and D); (R = Me, Et and X = Cl, Br), the isomeric shifts values are similar to those of the literature- the increase of the isomeric shift values when going from chlorine to bromine is respected [10]. The proposed structure is a discrete structure with a monodentate hydrogenotungstate (Figure 2).

The formulae difference between Me\(_4\)NHWO\(_4\)SnBr\(_4\) and Me\(_3\)NHWO\(_4\)SnCl\(_3\)NMe\(_4\) compounds can be due to the size of the halides since [HWO\(_4\)SnCl\(_4\)]\(^{-}\) and [HWO\(_4\)SnCl\(_3\)]\(^{2-}\) complexes anions could be obtained and not [HWO\(_4\)SnBr\(_5\)]\(^{2-}\). In fact, we know that the ratio (cation radius/anion radius) determines the type of crystal arrangement and the global environment around the metallic center. So, we see that with the tetraethylammonium ion, we find in the chloro complexes, as in the bromo complex, [HWO\(_4\)SnX\(_3\)]\(^{2-}\) complex anion whereas with smaller cation (tetramethylammonium ion) it was not possible to obtain Me\(_4\)NHWO\(_4\)SnBr\(_5\)NMe\(_4\).
Fig. 2. Proposed structure for compounds B, C and D.

For $\text{(Et}_4\text{NHWO}_4)_2\text{SnX}_3\text{HWO}_4$ (E and F) ($X = \text{Cl, Br}$), with an excess of ligand it was possible to substitute a halogen of SnX$_3$ by HWO$_4^-$ to obtain (SnX$_3$HWO$_4$) which then reacts with two hydrogenotungstate ions to give the complexes E and F.

The values of the isomeric shift confirm the expected increase when going from chlorine to bromine [10]. The proposed structure is discrete, the environment around the tin being octahedral (Figure 3) (we note that [SnX$_3$(HWO$_4$)$_3$]$^2-$ complex anion could only be obtained with the tetraethylammonium ion and not with the tetramethylammonium ion).

N.B.: In all the proposed structures, the $\text{–OH}$ groups, when considered involved in hydrogen bonding, supramolecular architectures may be obtained.

4. CONCLUSION

With the hydrogenotungstate, tin (IV) halides form mononuclear complexes of mostly discrete structure (five out of six), the anion generally behaving as a monodentate. As in the case of organotin, the hydrogenotungstate was found to be in the starting acidic salt, the chemical entity having the strongest coordinating power. The size of the antagonist cation appears to be decisive in the stoichiometry of the complex ion obtained.
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