

SHORT COMMUNICATION

AMINOPHENYL ARSENIATO ADDUCTS AND DERIVATIVES OF SnR_2Cl_2 (R = Ph, Bu): SYNTHESIS AND INFRARED STUDY

Bocar Traoré, Libasse Diop*, Mamadou Sidibé

*Université Cheikh Anta Diop, Faculté des Sciences et Techniques,
Laboratoire de Chimie Minérale et Analytique (LACHIMIA),
Dakar, Senegal*

*Corresponding author: dlibasse@gmail.com

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Abstract: Four new phenylarseniato adducts and derivatives have been synthesized on allowing SnR_2Cl_2 (R = Ph, Bu) to react with $\text{Cy}_2\text{NH}_2\cdot\text{NH}_2\text{PhAsO}_3\text{H}\cdot 3/2\text{H}_2\text{O}$ in specific ratios. The obtained new tin (IV) compounds were studied by infrared, infinite chain and discrete structures were suggested on the basis of spectroscopic data, the oxyanion behaving as a bridging bidentate or a tridentate ligand. When secondary interactions are involved supramolecular architectures were obtained.

Keywords: *coordinating phenylarseniato, SnR_2 residue, infrared, infinite chain and discrete structures, supramolecular architectures*

INTRODUCTION

Evans and Karpel have summarized the main applications found in organotin (IV) family (medicine, agriculture, antifouling paints, and wood preservatives) [1]. This explains the focus of many teams involving our [2-18] in seeking new molecules for widening the family, expecting some applications, and for structural aspects too. Our group initiates here the study of the interactions between $\text{Cy}_2\text{NH}_2\cdot\text{NH}_2\text{AsO}_3\text{H}$ and SnPh_2Cl_2 and SnBu_2Cl_2 which has yielded four new adducts and derivatives, infrared study of which has been carried out then structures suggested on the basis of spectroscopic data.

EXPERIMENTAL

$\text{Cy}_2\text{NH}_2\cdot\text{NH}_2\text{PhAsO}_3\text{H}\cdot 3/2\text{H}_2\text{O}$ (**A**) has been obtained on partially neutralizing $\text{NH}_2\text{PhAsO}_3\text{H}_2$ with Cy_2NH in water and allowing the water to evaporate at 60 °C. On mixing an ethanolic solution of (**A**) with SnPh_2Cl_2 or SnBu_2Cl_2 in specific ratios, precipitation occurs. The precipitate were stirred around two hours and filtered. The analytical data % Calculated (% Found) reported below have allowed to suggest the following formula.

Table 1. Results of the elemental analyses

Compound	Chemical formula	Elemental analysis (%)					
		C		H		N	
		calc.	found	calc.	found	calc.	found
B₁	$\text{NH}_2\text{PhAsO}_3\text{H}\cdot\text{SnPh}_2\text{Cl}_2\cdot 2\text{EtOH}$ (2/1)	40.05	39.32	5.42	6.06	2.12	1.88
B₂	$(\text{Cy}_2\text{NH}_2)_2\text{NH}_2\text{PhAsO}_3\cdot 3\text{SnBu}_2\text{Cl}_2\cdot \text{H}_2\text{O}$ (1/2)	42.72	42.62	7.26	7.03	2.77	3.04
B₃	$(\text{Cy}_2\text{NH}_2)_2\text{NH}_2\text{PhAsO}_3\cdot 2\text{SnPh}_2\text{Cl}_2\cdot \text{Cy}_2\text{NH}_2\text{Cl}\cdot 2(\text{EtOH})$ (1/1)	53.57	54.62	7.01	8.12	3.53	3.85
B₄	$\text{NH}_2\text{PhAsO}_3(\text{SnPh}_2\text{Cl}_2)_2\cdot 2\text{H}_2\text{O}$ (1/2)	41.50	42.00	3.45	4.73	1.61	1.56

The elemental analyses have been performed by the Laboratory of Microanalyses-University of Montpellier II-France The infrared spectra have been recorded at the University of Padova-Italy- by means of a Perkin Elmer spectrometer using CsI, the sample being as Nujol mulls. Infrared data are given in cm^{-1} (IR abbreviations: (vs) very strong, (s) strong, (m) medium, (w) weak, (br) broad). All the chemicals were purchased from Aldrich-Germany- and used as such.

RESULTS AND DISCUSSION

Let us consider the IR data of **B₁-B₄**:

(**B₁**) = νOH :429(br); νNH_2 :2425(m); δNH_2 :1212(s); ρNH_2 :1212(s); νAsO :868(vs); δAsO :445(vs); νSnPh_2 :272(m);

(**B₂**) = νOH :3400(s); $\nu\text{NH}_{2\text{cat}}$:2525(s); $\nu\text{NH}_{2\text{an}}$:2425(m); $\delta\text{NH}_{2\text{cat}}$:1622(m); $\delta\text{NH}_{2\text{an}}$:1596(s); ρNH_2 :1223(w); νAsO :849(vs); νasSnBu_2 :743(s); νSnBu_2 :609(m);

(**B₃**) = νOH :3400(br); $\nu\text{NH}_{2\text{cat}}$:2525(s); $\nu\text{NH}_{2\text{an}}$:2425(m); $\delta\text{NH}_{2\text{cat}}$:1622(m); $\delta\text{NH}_{2\text{an}}$:1596(s); ρNH_2 :1223(w); νAsO :890(vs); νSnPh_2 :273(s);

(**B₄**) = νOH :3300(br); νNH_2 :2400(m); δNH_2 :1650(s); ρNH_2 :1220(s); νAsO :880(vs); δAsO :440(vs); νSnPh_2 :270(m);

SnBu_2 appears as a weak bond allowing us to conclude to linearity for this compound. Therefore we can suggest the discrete structures reported on Figure 1, 2, 3 and 4.

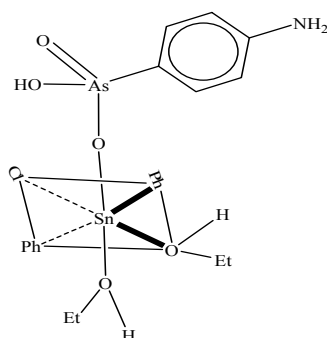


Figure 1. (B₁)

This structure is discrete with a monocoordinating anion, the environment around the tin (IV) centre being octahedral. The ethanol molecules are coordinating ones.

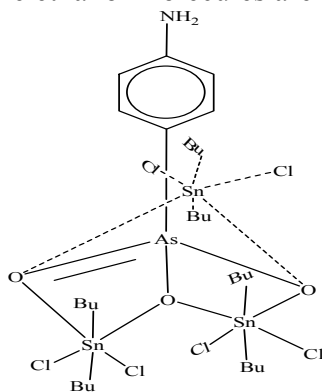


Figure 2. (B₂)

This structure is discrete with a trichelating anion, the environment around the tin (IV) centers being octahedral with the butyl groups in *trans* positions.

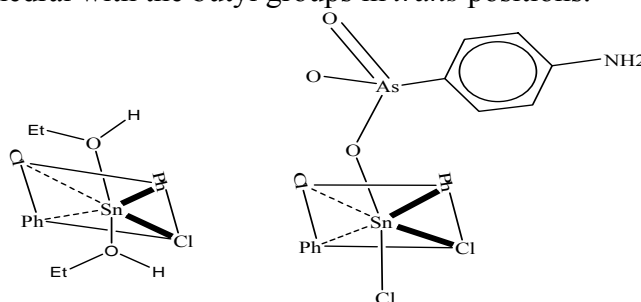


Figure 3. (B₃)

This compound is a 1/1 adduct between $\text{SnPh}_2\text{Cl}_2 \cdot 2\text{H}_2\text{O}$ and $(\text{Cy}_2\text{NH}_2)_3\text{NH}_2\text{PhAsO}_3 \cdot \text{SnPh}_2\text{Cl}_3$; it has an ionic structure with two metallic components (an anionic one and a neutral component), the tin (IV) centers being in an octahedral environment. The anion is a monocoordinating ligand, the water molecules being coordinated to the tin (IV) centre.

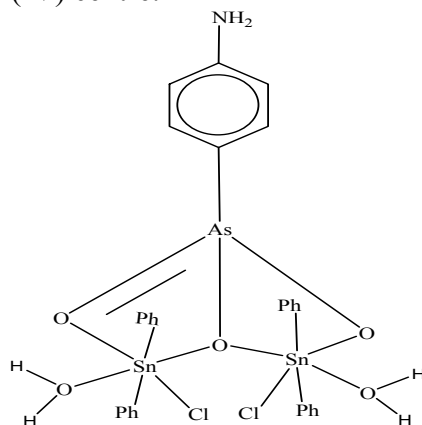


Figure 4. (B_4)

CONCLUSIONS

This compound has a discrete structure with a bichelating oxoanion, the environment around the tin centres being octahedral. The water molecules are coordinated ones. When cations or OH groups are involved in $\text{NH} \cdots \text{O}$, $\text{NH} \cdots \text{Cl}$ or $\text{OH} \cdots \text{Cl}$ hydrogen bonds, supramolecular architectures are obtained for all the structures.

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