

**SHORT COMMUNICATION**

**$(\text{Me}_4\text{N})_2\text{O}_2\text{CSO}_3\text{Sn}(\text{O}_2\text{CSO}_3)_2 \cdot 7\text{Me}_4\text{NSnCl}_5$ :  
SYNTHESIS AND INFRARED STUDY**

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**Abstract:** On allowing  $\text{Me}_4\text{NO}_2\text{CSO}_3\text{H}$  (in water) to react with  $\text{SnCl}_4$  (in ethanol) the studied complex is obtained. The suggested structure is discrete, the anion behaving as a polydentate and chelating ligand, the environment around tin being octahedral.

**Keywords:** *coordinating  $\text{O}_2\text{CSO}_3\text{H}$ , discrete structure, pentagonal environment, supramolecular architecture*

## INTRODUCTION

Several adducts containing  $\text{SnCl}_5^-$  have been reported [1 - 4]. In the framework of our research work on the coordinating ability of oxyanions we have yet published two papers dealing with and containing  $\text{SnCl}_5^-$  complex-anion [5] or  $\text{MX}_2$  [6].

In this dynamic, we have initiated in this work the study of the interactions between  $\text{Me}_4\text{NO}_2\text{CSO}_3\text{H}$  and  $\text{SnCl}_4$  which have yielded the studied octanuclear complex, infrared study of which have been carried out, then a structure suggested on the basis of infrared data.

## EXPERIMENTAL

On reacting imino (amino)methane sulfonic acid with  $\text{Me}_4\text{NOH}$  in water in 1/1 ratio the  $\text{Me}_4\text{NO}_2\text{CSO}_3\text{H}$  salt (white powder) is collected after a solvent evaporation at  $60^\circ\text{C}$ . When a mixed solution (water/ethanol) of  $\text{Me}_4\text{NO}_2\text{CSO}_3\text{H}$  is allowed to react with an ethanolic solution of  $\text{SnCl}_4$  in 1/1 ratio, a clear solution is obtained and stirred during two hours.

A white powder is obtained when this solution is submitted to a slow solvent evaporation. The elemental analyses data: % calculated (% found):

%C: 14.74 (13.97); %H: 3.40 (3.92); %N: 3.97 (4.68) have allowed to suggest  $(\text{Me}_4\text{N})_2\text{O}_2\text{CSO}_3\text{Sn}(\text{O}_2\text{CSO}_3)_2 \cdot 7\text{Me}_4\text{NSnCl}_5$  formula.

The elemental analyses were performed by Department of Chemistry, University of Bath (UK).

The infrared spectra were recorded by a FTIR-Nicolet ( $4000\text{--}400\text{ cm}^{-1}$ ) spectrometer at the University of Addis Ababa (Ethiopia), the sample being as Nujol mulls, using CsI windows.

Infrared data are given in  $\text{cm}^{-1}$ . IR abbreviations: (br) broad, (vs) very strong, (s) strong, (m) medium, (sh) shoulder, (vw) very weak.

All the chemicals were purchased from Aldrich and used without any further purification.

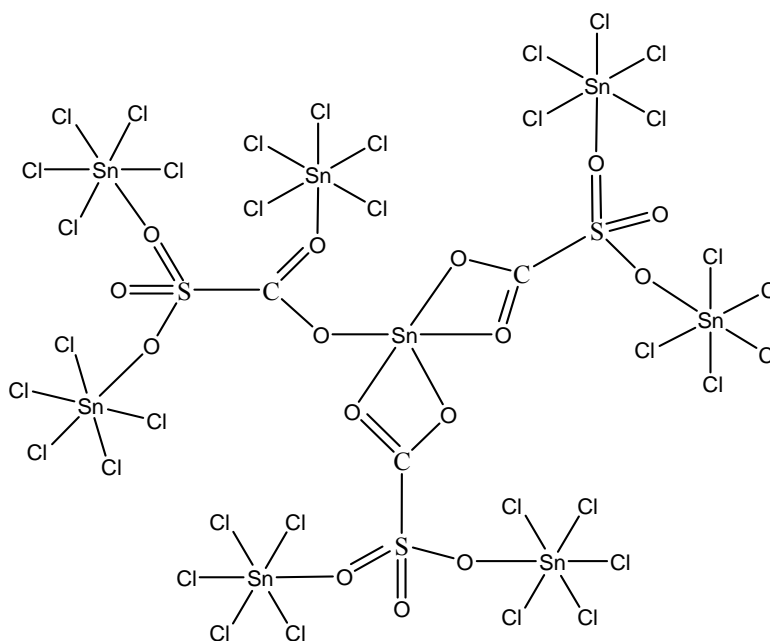
## RESULTS AND DISCUSSION

The spectroscopic infrared data ( $\text{cm}^{-1}$ ):  $\nu\text{SO}_3^-$ : 1146 (m);  $\nu_{\text{as}}\text{CO}_2^-$ : 1655 (m);  $\nu_{\text{s}}\text{CO}_2^-$ : 1292 (m).

The environment around the central tin centre is pentagonal. Two oxyanions bearing each one two  $[\text{SnCl}_5]^-$  are symmetrically chelating the central tin. The third one bearing three  $[\text{SnCl}_5]^-$  is monocoordinated to the central tin. The proposed structure of  $(\text{Me}_4\text{N})_2\text{O}_2\text{CSO}_3\text{Sn}(\text{O}_2\text{CSO}_3)_2 \cdot 7\text{Me}_4\text{NSnCl}_5$  are presented in Figure 1.

## CONCLUSIONS

The studied complex has a discrete structure with tridentate or monochelating and bidentate anion, the environment around the tin centre being octahedral or pentagonal.



**Figure 1.** Proposed structure for the compound  
(Me<sub>4</sub>N)<sub>2</sub>O<sub>2</sub>CSO<sub>3</sub>Sn(O<sub>2</sub>CSO<sub>3</sub>)<sub>2</sub>·7Me<sub>4</sub>NSnCl<sub>5</sub>

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