Research Article

Synthesis and Characterization of

$[2\text{-CH}_3\text{CH}_2\text{C}_6\text{H}_4\text{NH}_3]_6\text{P}_6\text{O}_{18} \cdot 4\text{H}_2\text{O}$

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Single crystals of $[2\text{-CH}_3\text{CH}_2\text{C}_6\text{H}_4\text{NH}_3]_6\text{P}_6\text{O}_{18} \cdot 4\text{H}_2\text{O}$ are synthesized in aqueous solution by the interaction of cyclohexaphosphoric acid and 2-ethylaniline. This compound crystallizes in the monoclinic system with $P2_1/c$ space group the unit cell dimensions are:

- $a = 16.220(4) \text{ Å}$,
- $b = 10.220(5) \text{ Å}$,
- $c = 20.328(4) \text{ Å}$,
- $\beta = 113.24(3)^\circ$,
- $Z = 2$, and
- $V = 3096.5(18) \text{ Å}^3$.

The atomic arrangement can be described by layers formed by cyclohexaphosphate anions $\text{P}_6\text{O}_{18}^-$ and water molecules connected by hydrogen bonds $\text{O} \cdots \text{H} \cdots \text{O}$. These inorganic layers are developed around $bc$ planes at $x = 1/2$ and are interconnected by the H-bonds created by ammonium groups of organic cations. All the hydrogen bonds, the van der Waals contacts and electrostatic interactions between the different entities give rise to a three-dimensional network in the structure and add stability to this compound. The thermal behaviour and the IR spectroscopic studies of this new cyclohexaphosphate are discussed.

1. Introduction

Organic-inorganic hybrid compounds have been of great interest in recent years due to their unique opportunity to combine the remarkable features of organic materials with those of inorganic compounds. In particular, hybrid materials based on the phosphates have attracted much attention due to their technological interest in several areas such as biomolecular sciences, catalysts, and optics [1–3].

The cohesion forces in these hybrid compounds are dominated by electrostatic interactions, van der Waals contacts and hydrogen bond (O–H···O, N–H···O and C–H···O). In order to enrich these materials and to investigate the influence of hydrogen bonds on the chemical and structural features, we report here the synthesis, crystal structure, thermal analysis, and IR spectroscopy of a new organic cyclohexaphosphate, $[2\text{-CH}_3\text{CH}_2\text{C}_6\text{H}_4\text{NH}_3]_6\text{P}_6\text{O}_{18} \cdot 4\text{H}_2\text{O}$.

2. Experimental Section

2.1. Synthesis. The title compound was prepared by an acid/base reaction in two steps. In the first one, we prepared Li$_6$P$_6$O$_{18}$·6H$_2$O according to the process described by Schülke and Kayser [4]. From this lithium salt, we prepared an aqueous solution of cyclohexaphosphate acid H$_6$P$_6$O$_{18}$ by passing a solution of Li$_6$P$_6$O$_{18}$·6H$_2$O (1 g, 2.3 mmol) through an ion-exchange resin in its H-state (Amberlite IR 120). In the second step, at 20 mL of the aqueous solution of H$_6$P$_6$O$_{18}$ freshly prepared, we add drop by drop a solution of 2-ethylaniline (1.70 mL, 13.8 mmol, $d = 0.983$) in 20 mL of ethanol under continuous stirring. Schematically, the reaction can be written as follows:

$\text{H}_6\text{P}_6\text{O}_{18} + 6 \cdot (2\text{-CH}_3\text{CH}_2\text{C}_6\text{H}_4\text{NH}_2) + 4\text{H}_2\text{O} \rightarrow [2\text{-CH}_3\text{CH}_2\text{C}_6\text{H}_4\text{NH}_3]_6\text{P}_6\text{O}_{18} \cdot 4\text{H}_2\text{O}$. 

In order to avoid the hydrolysis of the ring anion, the above reaction is performed at room temperature. The obtained solution was then slowly evaporated until the formation of colorless prisms of $[2\text{-CH}_3\text{CH}_2\text{C}_6\text{H}_4\text{NH}_3]_6\text{P}_6\text{O}_{18} \cdot 4\text{H}_2\text{O}$ ($m = 1.9$ g, yield = 65%). The title compound was stable for months under normal conditions of temperature and relative humidity.
2.2. X-Ray Diffraction. The intensity data were collected at room temperature using an Enraf-Nonius CAD4 diffractometer with MoKα radiation. The cell parameters were determined from a least square refinement of 25 reflections. Two-standard reflections were periodically measured for every 120 min during data collection. Unique reflections (8617) were measured of which only 3940 had I ≥ 2σ(I) and were used for structure determination and refinement. The structure was solved by direct method using the program SHELXS-97 [5] in the WinGX package [6] and refined by full-matrix least-squares method with the program SHELXL-97 [5]. All nonhydrogen atoms were refined isotropically and then anisotropically by full matrix least-square method. All water H atoms were first located in a difference Fourier map and then refined with distance restraints of O–H = 0.85(1) Å and H⋯H = 1.44(2) Å. The remaining hydrogen atoms were placed geometrically and treated as riding. The parameters used for the X-ray diffraction data collection as well as the strategy used for the crystal structure determination and its final results are reported in Table 1. The main bond distances and bond angles for the title compound are provided in Table 2 and the main geometrical features of the hydrogen-bond scheme in Table 3.

Crystallographic data (CIF) for the structure reported in this paper have been deposited in the Cambridge Crystallographic Data centre as supplementary materials No CCDC 798115. Copies of the data can be obtained, free of charge, on application to the CCDC, 12 Union Road,
2.3. Thermal Analysis. Thermal analysis was performed using the “multimodule 92 Setaram analyser” operating from room temperature up to 673 K at an average heating rate of 5 K min⁻¹.

2.4. Infrared Spectroscopy. IR spectrum is recorded in the range 4000–400 cm⁻¹ with a “PerkinElmer FTIR” spectrometer using a sample dispersed in spectroscopically pure KBr pellet.

3. Results and Discussion

3.1. Crystal Structure. The title compound (Figure 1) is a hybrid of organic and inorganic entities (2-ethylaniline, \( \text{P}_6\text{O}_{18} \) and \( \text{H}_2\text{O} \)). The asymmetric unit of \( [2-\text{CH}_3\text{CH}_2-\text{C}_6\text{H}_4\text{NH}_3]_6\text{P}_6\text{O}_{18} \cdot 4\text{H}_2\text{O} \) contains a half of the anion, three organic cations and two water molecules crystallographically independent.

The atomic arrangement of the title compound is a typical organization in layers. These layers are built up from anions and water molecules that develop parallel to the bc planes at \( x = 1/2 \) (Figure 2). Charge compensation of these layers is achieved by the incorporation of the protonated 2-ethylanilinium cations in the interlayer spaces establishing H-bonds via their ammonium groups with \( \text{P}_6\text{O}_{18} \) ring and water molecules. Adjacent \( \text{P}_6\text{O}_{18} \) groups are linked pairwise with \( \text{O}11 \) water molecules so as to form an infinite chain spreading along the b direction. On the other hand, \( \text{O}10 \) water molecules link these infinite chains to give rise to infinite layers. Each cyclohexaphosphate group is connected to its adjacent neighbours by eight water molecules through O–H···O hydrogen bonds. Inside such a structure, the complete cyclohexaphosphate anion is generated by inversion symmetry located at \((1/2, 1/2, 0)\) and \((1/2, 0, 1/2)\). So, it is built up by only three independent tetrahedral, \( \text{P}(1)\text{O}_4 \), \( \text{P}(2)\text{O}_4 \), and \( \text{P}(3)\text{O}_4 \). The geometry of this ring (Table 2) does not differ from the other known cyclohexaphosphates. The P–O distances range in (1.467(3)–1.610(3) Å) and the O–P–O bond angles in (99.56(15)–120.34(15)°). It is the same for the P–P distances ranging from 2.897 (1) to 2.940 (1) Å, which are comparable to values generally measured [7–9].

The calculated average values of the distortion indices [10] corresponding to the different angles and distances in the \( \text{PO}_4 \) tetrahedra (DI(OPO) = 0.041, DI(PO) = 0.041; and DI(OO) = 0.012), show a pronounced distortion of the
PO distances and OPO angles if compared to OO distances. So, the PO₄ group can be considered as a rigid regular arrangement of oxygen atoms, with the phosphorus atom slightly displaced from the tetrahedron gravity centre.

Three independent 2-ethylanilinium cations are identified in the title compound. Intertatomic bond lengths and angles of these groups are as expected [11, 12]. The aromatic rings are planar with an average deviation of 0.000349 Å and form dihedral angles of 2.56°, 8.50° and 8.59° between them. The interplanar distance between nearby phenyl rings is in the vicinity of 3.972 Å, which is slightly longer than 3.80 Å for the π–π interaction [13].

The 2-ethylanilinium cations occupy the interlayer spaces and establish with the anionic framework N–H···O hydrogen bonds. The N(1)H3 groups produce the internal P₅O₁₈ ring cohesion through hydrogen bonds involving external oxygen atom of each PO₄ tetrahedron. The N(3)H₃ groups link two different rings so as to form infinite chain extending along the b direction at z = 1/2. While the N(2)H₃ groups are bounded to two different water molecules and one ring situated at z = 0 and so contribute to the layer cohesion of this compound. The organic groups are also interconnected through weak C–H···O hydrogen bonds with the C···O separation of 3.281(6)–3.422(8) Å. The C–H···O bonds have already been evidenced by several authors in molecular crystals [14].

There are two crystallographically independent water molecules O(10) and O(11) in this atomic arrangement. They interconnect the cyclohexaphosphate rings and the organic groups by establishing the hydrogen bonds; thus
the water molecules act as proton donors and acceptors while the protonated amine groups are exclusively acceptors (Figure 3, Table 3). The cohesion forces in this compound are assured by electrostatic interactions, van der Waals contacts and hydrogen bonds (O–H···O, N–H···O and C–H···O).

3.2. Thermal Analysis. The two curves corresponding to DTA and TGA analysis carried out in an argon atmosphere in the temperature range [298 K–673 K] are given in Figure 4. The DTA curve shows that the cyclohexaphosphate undertakes a succession endothermic phenomenon. The first two peaks correspond to the dehydration, which occurs at about 361 K, and 377 K are accompanied with a weight loss obviously observed in the TGA curve. From this latter, we deduce a departure of four water molecules (% experimental: 5.59%, calculated: 5.63%). At about 419 K, another endothermic peak appears, corresponding to the degradation of the 2-ethylanilinium, accompanied with a significant weight loss clearly observed on the TGA curve. This degradation gives a viscous substance of polyphosphoric acids contaminated with a carbon black deposit.
3.3. IR-Absorption Spectroscopy. The infrared absorption spectrum is reported in Figure 5. It exhibits

(i) broad bands between 3600–3300 cm$^{-1}$ and 1600–1370 cm$^{-1}$ corresponding, respectively, to the stretching and bending modes of NH$_3$. They are indicative of the presence of the 2-ethylaniline molecules in its protonated form [15]. The broadness of the IR bands is probably due to the existence of hydrogen bonds of varying strengths in the crystal;

(ii) various valency vibration bands whose positions, between 1350 and 660 cm$^{-1}$, are characteristic of cyclohexaphosphate anion [16]. In this type of anions, the stretching vibrations $\nu_{as}(\text{OPO})^{-}$ and $\nu_{s}(\text{OPO})^{-}$ are observed, respectively, between 1350–1180 cm$^{-1}$ and 1180–1060 cm$^{-1}$, while the stretching vibration bands originating from both asymmetric $\nu_{as}(\text{POP})$ and symmetric $\nu_{s}(\text{POP})$ modes are, respectively, observed in the range (1060–960) and (850–660) cm$^{-1}$ region, and those below 660 cm$^{-1}$ correspond to bending, translation, and rotation of the P$_6$O$_{18}$ ring.

4. Conclusions

[2-CH$_3$CH$_2$C$_6$H$_4$NH$_3$]$_6$P$_6$O$_{18}$·4H$_2$O is prepared as single crystals at room temperature and characterized by X-ray diffraction, thermal behaviour, and IR spectroscopic. Crystal structure of this compound was found to be built by infinite layers of P$_6$O$_{18}^{-}$ and water molecules parallel to the bc planes around $\chi$ = 1/2. Between these layers, the 2-ethylanilinium cations are located. Both inorganic and organic components perform different interactions (electrostatic, H-bonds and van der Waals) to stabilize the three-dimensional network. When heated, the compound loses the four coordinated water molecules at 361 K and 377 K. By heating further, the compound exhibits a degradation of the organic entities confirmed by the obtained carbon black residue at the end of the experiment.

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