

Electronic Supplementary Information for MS:

Reversible Solid State Structural Transformation of a Polyhapto Lead(II) Polymeric Chain to a Tetrahapto Lead(II) Two-Dimensional Network by Thermal Dehydration with no Change in Nano-plate Morphology

Kamran Akhbari, Ali Morsali*

^aDepartment of Chemistry, Faculty of Sciences, Tarbiat Modares University, P.O. Box 14115-175, Tehran, Islamic Republic of Iran

Table S1 Bond lengths /Å and angles /° for [Pb(MPOAc)₂(H₂O)]_n (**1**).

Pb1-O3	2.450(3)
Pb1- O1	2.498(3)
Pb1- O4	2.516(3)
Pb1- O4 ⁱ	2.534(3)
Pb1-O2	2.545(4)
Pb1-O1W	2.724(4)
Pb1-C16	2.838(5)
Pb1 ⁱ - O4	2.534(3)
O3-Pb1-O1	80.07(11)
O3-Pb1-O4	52.17(11)
O1-Pb1-O4	111.78(11)
O3-Pb1-O4 ⁱ	70.87(11)
O1-Pb1-O4 ⁱ	73.26(11)
O4-Pb1-O4 ⁱ	118.94(7)
O3-Pb1-O2	86.22(11)
O1-Pb1-O2	51.94(11)
O4-Pb1-O2	76.47(11)
O4 ⁱ -Pb1-O2	123.61(11)
O3-Pb1-O1W	73.83(11)
O1-Pb1-O1W	140.19(10)
O4-Pb1-O1W	74.40(11)
O4 ⁱ -Pb1-O1W	70.07(10)
O2-Pb1-O1W	150.75(11)
O3-Pb1-C16	26.47(12)
O1-Pb1-C16	98.96(13)
O4-Pb1-C16	26.06(11)
O4 ⁱ -Pb1-C16	93.98(12)
O2-Pb1-C16	83.47(12)
O1W-Pb1-C16	69.09(12)
C8-O1-Pb1	94.0(3)
C8-O2-Pb1	91.7(3)
C16-O3-Pb1	94.1(3)
C16-O4-Pb1	91.6(3)
C16-O4-Pb1 ⁱⁱ	144.5(3)
Pb1-O4-Pb1 ⁱⁱ	116.12(13)
O4-C16-Pb1	62.4(2)
O3-C16-Pb1	59.5(2)
C15-C16-Pb1	167.8(3)
Pb1-O1W-H1W	119.6
Pb1-O1W-H2W	114.1

Symmetry operations: i: -x+1/2, y-1/2, -z+1/2 and ii: -x+1/2, y+1/2, -z+1/2.

Table S2 Bond lengths /Å and angles /° for [Pb(MPOAc)₂]_n (**2**).

Pb1-O1	2.409(3)
Pb1-O2	2.552(3)
Pb1-O1 ⁱ	2.694(3)
O1-Pb1-O1 ⁱⁱ	84.45(13)
O ⁱⁱ -Pb1-O2 ⁱⁱ	52.13(8)
O1 ⁱⁱ -Pb1-O2 ⁱⁱ	87.73(10)
O2-Pb1-O2 ⁱⁱ	127.63(15)
O1 ⁱ -Pb1-O1 ⁱⁱⁱ	66.91(10)
O1-Pb1-O1 ⁱⁱⁱ	123.10(8)
O2 ⁱⁱ -Pb1-O1 ⁱⁱⁱ	107.85(8)
O2 ⁱⁱ -Pb1-O1	77.60(8)
O1-Pb1-O1 ⁱ	66.91(10)
O1 ⁱⁱ -Pb1-O1 ⁱⁱⁱ	168.07(12)

Symmetry operations: i: -x, y, -z+1/2; ii: -x, -y, -z and iii: x, -y, z+1/2.

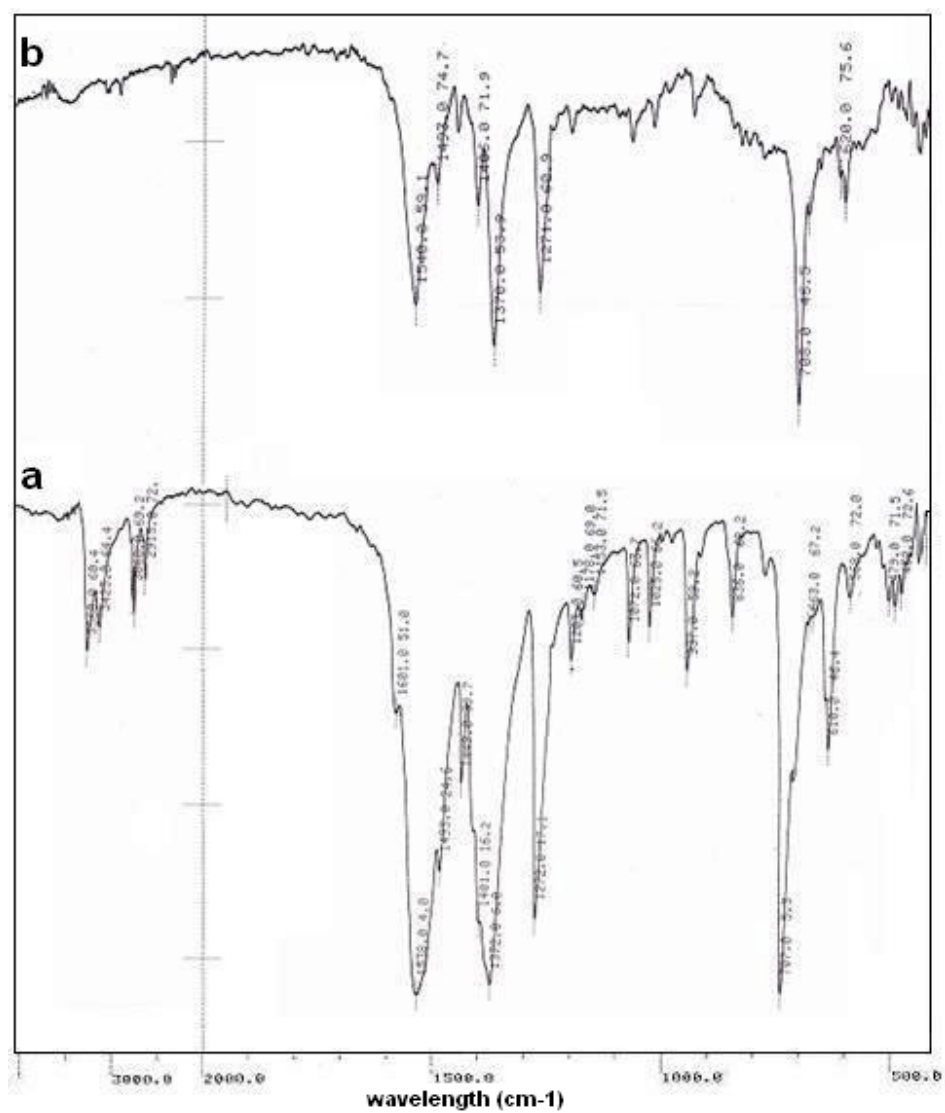


Figure S1. IR spectra of **a**) compound **1** and **b**) bulk materials obtained by heating of compound **1**.

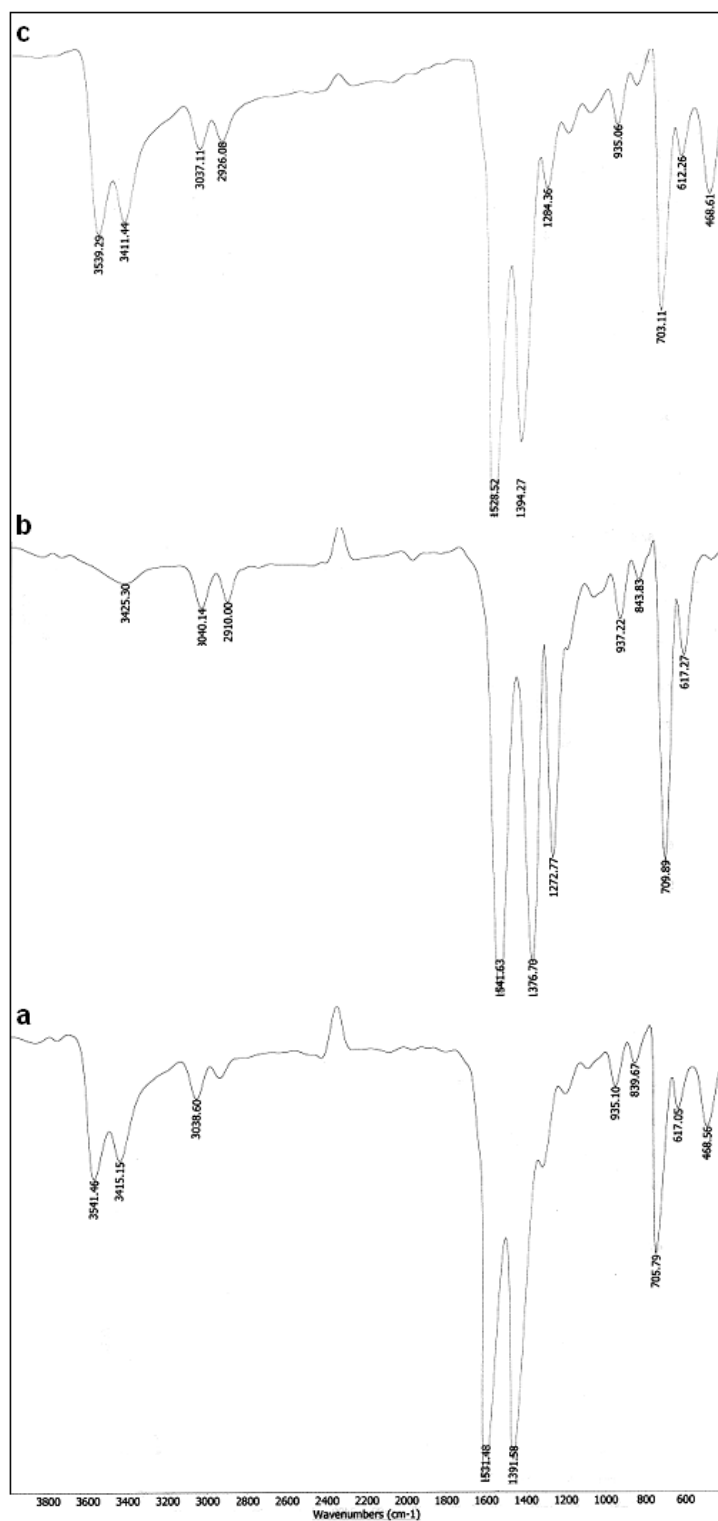


Figure S2. IR spectra of **a)** compound **1** nano-plates prepared by microwave-assisted process, **b)** compound **2** nano-plates obtained by heating of compound **1** nano-plates at 120 °C for 12 hours and **c)** the microcrystalline sample obtained from dehydrated compound **2** nano-plates under hydrothermal conditions at 165 °C for two days.