A polymorph of dipeptide halide Glycyl-L-Alanine Iodide Hydrate: crystal structure, optical second harmonic generation and pyroelectricity

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**Supplementary Information**

**SI1. Crystal Structure**

**Table SI1.** Bond lengths [Å] and angles [°] for Gly-L-Ala.HI.H2O (Poly2).

|  |  |
| --- | --- |
| *Bond lengths* | *Bond angles* |
| C(1)-N(1) 1.481(4)C(1)-C(2) 1.532(6)C(1)-H(1A) 0.9700C(1)-H(1B) 0.9700C(2)-O(1) 1.243(6)C(2)-N(2) 1.323(7)C(3)-N(2) 1.459(7)C(3)-C(5) 1.521(8)C(3)-C(4) 1.537(9)C(3)-H(3) 0.9800C(4)-H(4A) 0.9600C(4)-H(4B) 0.9600C(4)-H(4C) 0.9600C(5)-O(2) 1.221(7)C(5)-O(3) 1.316(7)N(1)-H(10A) 0.88(3)N(1)-H(10B) 0.85(3)N(1)-H(10C) 0.88(3)N(2)-H(20) 0.89(10)O(3)-H(30) 0.91(10)O(4)-H(40) 0.8159(5)O(4)-H(41) 0.8283(5) | N(1)-C(1)-C(2) 109.9(4)N(1)-C(1)-H(1A) 109.7C(2)-C(1)-H(1A) 109.7N(1)-C(1)-H(1B) 109.7C(2)-C(1)-H(1B) 109.7H(1A)-C(1)-H(1B) 108.2O(1)-C(2)-N(2) 124.5(5)O(1)-C(2)-C(1) 120.9(4)N(2)-C(2)-C(1) 114.6(4)N(2)-C(3)-C(5) 109.7(5)N(2)-C(3)-C(4) 111.7(5)C(5)-C(3)-C(4) 110.4(5)N(2)-C(3)-H(3) 108.3C(5)-C(3)-H(3) 108.3C(4)-C(3)-H(3) 108.3C(3)-C(4)-H(4A) 109.5C(3)-C(4)-H(4B) 109.5H(4A)-C(4)-H(4B) 109.5C(3)-C(4)-H(4C) 109.5H(4A)-C(4)-H(4C) 109.5H(4B)-C(4)-H(4C) 109.5O(2)-C(5)-O(3) 123.9(5)O(2)-C(5)-C(3) 124.1(5)O(3)-C(5)-C(3) 111.9(5)C(1)-N(1)-H(10A) 106(4)C(1)-N(1)-H(10B) 109(4)H(10A)-N(1)-H(10B) 100(7)C(1)-N(1)-H(10C) 124(9)H(10A)-N(1)-H(10C) 106(8)H(10B)-N(1)-H(10C) 110(10)C(2)-N(2)-C(3) 124.4(4)C(2)-N(2)-H(20) 118(6)C(3)-N(2)-H(20) 117(6)C(5)-O(3)-H(30) 106(6)H(40)-O(4)-H(41) 99.17(2) |

**Table SI2.** Torsion angles [°] for Gly-L-Ala.HI.H2O (Poly2), Gly-L-Ala.HI.H2O (Poly1) and Gly-L-Ala.

|  |  |  |  |
| --- | --- | --- | --- |
|  | ***Poly 2*** | ***Poly 1* [**[**1**](#_ENREF_1)**]** | ***Gly-L-Ala* [**[**2**](#_ENREF_2)**]** |
| N(1)-C(1)-C(2)-O(1) | -14.2(6) |  |  |
| N(1)-C(1)-C(2)-N(2) | 167.9(5) | 168.13 | -163.54 |
| N(2)-C(3)-C(5)-O(2) | -12.7(7) | -12.94 | -35.88 |
| C(4)-C(3)-C(5)-O(2) | 110.8(7) |  |  |
| N(2)-C(3)-C(5)-O(3) | 167.9(5) | 167.78 | 149.99 |
| C(4)-C(3)-C(5)-O(3) | -68.5(6) |  |  |
| O(1)-C(2)-N(2)-C(3) | -0.2(9) |  |  |
| C(1)-C(2)-N(2)-C(3) | 177.7(5) | 177.88 | -173.94 |
| C(5)-C(3)-N(2)-C(2) | -126.2(5) | -126.27 | -78.36 |
| C(4)-C(3)-N(2)-C(2) | 111.0(6) |  |  |



**Figure SI1.** Unit cell of Gly-L-Ala.HI.H2O (Poly2) showing the (010) plane, in yellow, perpendicular to the *b* axis.



**Figure SI2**. Gly-L-Ala.HI.H2O (Poly2) morphology with (*hkl*) labels.



**Figure** **SI3**. The main chain of the dipeptide molecule presents an almost planar *trans* conformation between atoms N1 and C3.

**SI2.** **Thermogravimetric analysis (TGA)**

The TGA spectrum shows a small initial mass loss of 2% happening at 388 K that corresponding to the starting of water molecules evaporation. Above 473 K there is a considerable loss of mass (about 50%), correspondent to the degradation of the crystalline compound. The peak at 533 K corresponds to the fusion temperature of the dipeptide glycyl-l-alanine, is very close to the melting temperature reported for cyclo-glycyl-L-alanine which was 531 K. Therefore, the peak at 648 K, corresponds to the complete degradation of crystalline compound Gly-L-AlaHI.H2O (Poly2).



**Figure SI4.** TGA spectra of synthesized glycyl-l-alanine iodide hydrate crystals.

**SI3. DSC analysis**

In figure SI5, the measured DSC of synthesized glycyl-l-alanine iodide hydrate crystals, shows one first peak at 363 K, area 5.55 J/g that indicates the starting of water loss in the structure. A second double peak shows at 388 K is followed by the maximum at 393 K with area 12.3 J/g corresponding to full loss of water. The cooling curve does not reproduce the heating curve, indicating that above 393 K the compound is not any more the initial one, according to what has just been stated before.



**Figure SI5**. DSC spectra of synthesized glycyl-l-alanine iodide hydrate crystals.

**SI4. FTIR-ATR analysis**

In Figure SI6 the measured FTIR-ATR between 550-3800 cm-1 is shown for a glycyl-l-alanine iodide hydrate powder sample.

The intense and sharp band at 3475 cm-1 is assigned to the strong hydrogen-bonding established between internal water molecules and the oxygen atoms from the dipeptide bond. The N–H stretching bands are relatively weak and appear at 3232 cm-1 and 3188 cm-1, while the C–H stretching bands are observed from 2960 to 3099 cm-1 region. Carbonyl stretching modes appear as sharp singlets at 1716 cm-1 and 1649 cm-1. These results are in accordance to those reported in the literature for the glycyl-l-alanine molecule [[3](#_ENREF_3)]. Deviation is observed in some bands, namely those related to asymmetric stretches in the terminal amine group, which can be explained by the fact that the dipeptide is in a different chemical environment due to its binding to the iodide ion and water molecule.



**Figure SI6.** FTIR-ATR spectra of glycyl-l-alanine iodide hydrate powder between 550 – 3800 cm-1.

**SI5. RAMAN Spectra**

In Figure SI7, the measured Raman modes between 50 – 800 cm-1, 800 – 1800 cm-1 and 2700 – 3500 cm-1 are shown for glycyl-l-alanine iodide hydrate polycrystalline sample. The measurements were made under no polarized light (black), vertical (VV, red) and horizontal (VH, blue) polarized light. In the spectra, the low frequency modes under 500 cm-1 are due to crystal lattice vibrations. In the region 800 – 1800 cm-1 there are 3 peaks, for which the relative intensities are for VV polarization smaller (935 cm-1 and 1276 cm-1) or higher for VH polarization (1090 cm-1). These indicates that although the sample was polycrystalline there was some preferential orientation on the crystallite’s distribution. It was not possible to compare the vibrations of the Raman modes with those of glycyl-l-alanine molecule because there is no information available in the literature.

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**Figure SI7.** Raman spectra for glycyl-l-alanine iodide hydrate powder, expansions between 50 – 800 cm-1, 800 – 1800 cm-1 and 2700 – 3500 cm-1.

**SI6. Second Harmonic Measurements**

Figure SI8 shows a representative fit to the acquired second harmonic spectra that we expect to be approximately Gaussian as a function of frequency, i.e.

 

For these data the fit yields a background B = -3.4 ± 0.3 Counts, an amplitude A = 828 ± 2 Counts, a central frequency 0 = 753.31±0.01 THz and a full width half maximum frequency of = 8.11 ± 0.02 THz. For reference, 759 THz corresponds to a wavelength of approximately 400nm:



**Figure SI8**. An example of the Gaussian fit to the obtained spectra. Blue circles represent the data points while the back line is the best least squares fit to the data.

For crystals belonging to point group 2, the nonlinear ssuceptibility tensor takes the form:

  (1)

Here, 1, 2 and 3 (or x , y and z respectively) are the electric field components of the incident laser beam in the frame of the crystal’s principal dielectric axes with the crystallographic b axis taken to coincide with the direction of the dielectric y axis. If one assumes Kleinman sysmetry then d21 = d16; d25=d36=d14 and d34=d23.

For these biaxial crystals the two eigen polarizations for the slow ($\hat{e}\_{s}$) and fast ($\hat{e}\_{f}$) waves of a beam propagating at an aribitary direction described by the polar angles  relative to the crystal’s dielectric z-axis are respectively:

  (2a)

and

  (2b)

Here  is the angle between the polarization direction of the slow wave and the plane of propagation containing the z-axis [[4](#_ENREF_4)]. Using these expressions it is possible to demonstrate that a fundamental beam propagating as an ordinary wave (extra-ordinary wave) incident along the dielectric y  axis will not generate any extraordinary (ordinary) second harmonic light [[4](#_ENREF_4)]. This precludes any chance of phase matching for beams travelling along the y dielectric axis due the dispersion between the fundamental and second harmonic waves. For beams that do propagate along or close to the y axis, appreciable second harmonic generation will in general only be observed when the incident beam waist is close to the exit surface.

To carry out the fit shown in Figure 15 of the main text, we followed the “double integral method” developed by Wang and Weiner [[5](#_ENREF_5)] to describe second harmonic generation by ultrashort pulses in the simultaneous presence of spatial and temporal walk-off as well as phase mismatch. Assuming Gaussian temporal and spatial profiles with a transverse circular symmetry, Wang and Weiner arrive at the following expression for the energy of the second harmonic wave generated by a crystal of length L:



Here the energies of the fundamental and second harmonic pulses are  and  respectively,  is the full width half maximum temporal pulse width,  and  are the fundamental beam’s Rayleigh distance and beam waist position in the crystal,  is the vaccum fundamental wavelength, while  and  are the refracitve indices of the fundamental and second harmonic beams respectively. The parameter  represents the combined effects of spatial and temporal walk-off and is given by,



with  the spatial walk-off angle in radians,  characterizes the extent to which a linear chirp has broadened the pulse beyond the Fourier transform limit (), , and  is the inverse group velocity mismatch between the fundamental and second harmonic beams. A nonlinear least squares minimization routine [[6](#_ENREF_6)] was used to estimate the values of and . The results corresponding to the fit shown in figure 15 are, ,, with  being so large (above 500 m) that it has a negligible effect.

We used a 2 mm Beta-Barium Borate (BBO) to calibrate our second harmonic microscope. BBO has a walk-off angle of 68.4 mrad, an inverse group velocity mismatch of  and we estimate that the linear chirp after passing through the microscope objective corresponds to a value of . Assuming the same value for the Rayleigh distance as given by the above fit for the glycyl-L-alanine dipeptide crystal, numerical integration of the double integral for BBO gives a peak value of 1560. In contrast, the peak value for the glycyl-L-alanine dipeptide crystal using the above estimated values is only 0.53 due to the large phase mismatch.

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