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Using Raman spectroscopy to understand the origin of the phase transition observed in the crystalline sulfur based amino acid L-methionine

José A. Lima Jr.^a, Paulo T.C. Freire^{a,*}, Francisco E.A. Melo^a, J. Mendes Filho^a, Jennifer Fischer^b, Remco W.A. Havenith^c, Ria Broer^c, Heloisa N. Bordallo^d

^a Departamento de Física, Universidade Federal do Ceará, C.P. 6030 Fortaleza-CE 60455-760, Brazil

^b Forschungszentrum Jülich, Institut of Complex Systems, 52425 Jülich, Germany

^c Theoretical Chemistry, Zernike Institute for Advanced Materials, University of Groningen, 9747 AG Groningen, The Netherlands

^d Niels Bohr Institute, University of Copenhagen, 2100 Copenhagen, Denmark

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ABSTRACT

We present the Raman spectra of L-methionine ($C_5H_{11}NO_2S$) monocrystals obtained in the spectral region ranging from 3200 to 50 cm^{-1} at temperatures from 20 to 375 K. We investigated the dynamics of the different functional groups in L-methionine and related their behaviour to the structural transition previously reported at about 307 K. Additionally, on cooling, changes in the intensities of some Raman bands were associated with conformational changes of at least one of the two L-methionine conformers in the monoclinic unit cell in the interval 160–140 K. Thermal analysis and DFT calculations provide further support to the interpretation of the Raman results.

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1. Introduction

In the past fifteen years, many studies have focused on the vibrational and structural properties of amino acids [1,2]. This is consequence of the fact that these studies can shed light on the dynamics of the functional groups constituting amino acids, the principal building blocks of proteins and enzymes, as well as provides information about H-bonds, which are of fundamental interest in biochemical processes and drug design, among others. Additionally, some research has pointed out the possibility of using crystalline amino acids in technological applications, including nonlinear optics and in the food industry [3,4]. Very recently, even one of the most impressive physical properties of solids – ferroelectricity – was reported in γ -glycine [5].

Among the proteic amino acids, two of them have sulfur in their structure: cysteine, $C_3H_7NO_2S$, and methionine, $C_5H_{11}NO_2S$. Various studies related to the vibrational and structural properties of both orthorhombic and monoclinic L-cysteine have been reported in the literature [6–13]. From these investigations, several aspects of the material were revealed, including the observation of changes in the orientation of the side group $-\text{CH}-\text{CH}_2\text{SH}$, associated with the extended phase transition observed at low temperatures in the orthorhombic polymorph [8,10], the re-orientation of the side

groups related to the dynamical transition detected in its monoclinic polymorph [12], as well as different responses to hydrostatic pressure on each polymorph [13]. On the other hand, the physical properties of L-methionine, which has a sulfur atom between two carbons implying that the H-bonds allowed in cysteine are forbidden in methionine, have seen little study. To the best of our knowledge, very few Raman studies have been reported to date for the solid crystalline form of L-methionine. Using unpolarized light, one compared the solid-state spectra at 273 and 323 K [14], and the other the spectra observed from solid-states samples and water solutions to theoretical spectra of the single-molecules obtained by quantum chemical calculations [15]. Combining IR and Raman spectroscopy to semi-empirical MNDO conformational analysis [14], it was shown that the phase transition observed by means of calorimetric measurements [16,17] on powder samples at about 307 K is related to the existence of a slightly modified H-bonded double layer giving rise to similar lattice vibrations in both phases. In addition, L-methionine and its N-deuterated isotopomer were characterized by infrared spectroscopy at room temperature in the spectral range $4000\text{--}400\text{ cm}^{-1}$ [18] and a Raman spectroscopic study of L-methionine crystal under high pressure (up to 5 GPa) [19] was also carried out. The latter showed pronounced changes in the Raman spectra, which were associated with a structural phase transition at about 2.2 GPa. On the structural side, studies were reported by three groups, one on powder samples [20] and two others using single crystals obtained from slow evaporation of ethanol–water solutions [21] and 2-propanol in

* Corresponding author. Tel.: +55 85 3366 9906; fax: +55 85 336 9450.

E-mail addresses: tarso@fisica.ufc.br, tarsofreire@gmail.com (P.T.C. Freire).

Fig. 1. Crystal structure of L-methionine at 120 K from X-ray diffraction [22]. Two different molecules (A and B) with distinct conformation are observed and only some atoms are labeled for clarification. Hydrogen-atoms (white) are on idealized positions.

tetramethoxysilane gel [22], at 300 and 120 K, respectively. All the studies show that L-methionine crystallizes in the monoclinic $P2_1$ space group with four molecules in the unit cell ($Z=4$), building up a double-layered structure having a hydrophilic and hydrophobic layer. In the hydrophilic layer the NH_3^+ and CO_2^- head groups are linked through H-bonds, while in the hydrophobic layer the residuals are held together by van-der-Waals forces, as shown in Fig. 1.

The L-form of methionine is considered an essential amino acid, being very important in the process of methylation and also constituting a precursor to other amino acids, such as L-cysteine and homocysteine, which are involved in numerous processes of methyl group transfer and which play fundamental roles in the biochemistry of the human body [23]. It has also been used as supportive supplement in immune-deficient patients [24], and its oxidation has been associated with some neurological disease such as Alzheimer's [25]. Recently L-methionine has attracted interest due its role on the inactivation of Pt complexes and in the treatment with Pt drugs [26]. Interestingly, it has been shown that L-methionine side chains in proteins possess unique structural properties. For instance, the flexibility of the L-methionine side chains provides for a malleable nonpolar surface that can easily adapt to a variety of peptide partners [27]. Moreover, in contrast to other hydrophobic residues, the side chains of L-methionine are unbranched allowing for conformational flexibility [28].

Measuring dynamic properties as a function of temperature constitutes a powerful means of obtaining insight on the strength of atomic interactions. On the experimental side, spectroscopy techniques such as nuclear magnetic resonance, Raman scattering, infrared and inelastic neutron spectroscopies are appropriate probes for studies of H-vibrational dynamics because subtle structural changes that arise from changes in packing density due to variations in intermolecular distances in the H-bond network can be followed without difficulty. In reality, since the first systematic Raman scattering study of amino acids by Edsall [29], Raman scattering has been increasingly used in structural biology investigations, as for instance in the study of conformational characterization of peptides and proteins [30] and protein binding constants [31]. In addition, it has been shown that the combined spectroscopic/crystallographic/thermal analysis approach is a powerful new tool to investigate intermolecular interactions in a variety of hydrogenated systems [32,33], as well as in the simpler amino acids residues [34,35]. In the particular case of nonpolar

Fig. 2. The B3LYP/6-31G** optimized geometry of L-methionine.

crystalline amino acids, by using such type of approach it was demonstrated that α -glycine and L-isoleucine are stable between 20 and 300 K [36–38], partially deuterated L-alanine undergoes a solid–solid phase transition around 260 K followed by conformational change at lower temperatures [39], L-valine undergoes a solid–solid phase transition about 120 K [40], L-leucine undergoes three solid–solid phase transitions at 150, 275 and 353 K [41], and interestingly the sulfur containing amino acid L-cysteine undergoes a subtle dynamical transition around 150 K in its monoclinic form [12] while for the orthorhombic polymorph an extended transition around 70 K was observed [10]. It is also worth mentioning that many other studies in amino acid, mainly using Raman spectroscopy, have been published in the past years [42–62].

In this work we investigate the Raman spectra of crystalline L-methionine at low and high temperatures. From this study we were able to demonstrate that the phase transition observed in L-methionine powder samples at 307 K is a result of the flexibility of the residue side chains. Moreover, through the careful inspection of the evolution of the Raman bands in the high wavenumber region, we were able to identify a conformational change of the molecule at low temperatures. DFT studies enabled us to model the observed Raman vibrations of L-methionine allowing for the assignment of most vibrations at the Γ point, while thermal analysis confirmed that the malleability of the L-methionine side chain residues gives rise to an intricate behavior of the H-bond network in this system.

2. Experimental

L-Methionine single crystals were grown by slow evaporation of an aqueous solution of a commercial reagent (Sigma) at a typical temperature of 275 K. After 3–4 weeks, it is possible to obtain crystals as thin elongated plates of diverse sizes. In the Raman experiment the backscattered light was analyzed through a Jobin Yvon Triplemate 64000 micro-Raman system equipped with an N_2 -cooled CCD detector coupled to a microscope Olympus BX40. The slits give a typical 2 cm^{-1} spectral resolution. Excitation ($\lambda=514.5\text{ nm}$) was supplied by an argon ion laser. For the low-temperature Raman measurements, the samples were placed in a closed-cycle helium refrigerator that controlled the temperature with a precision of $\sim 0.2\text{ K}$.

Calculations on molecular L-methionine were performed at the B3LYP/6-31G** level of theory with the GAMESS-UK package [63]. The geometry was optimized and confirmed to be a genuine minimum by a Hessian calculation. The rotational profiles around the various bonds were determined by varying the corresponding dihedral angle from 0° to 360° in steps of 10° , and this dihedral was kept fixed during optimization of the remaining structural parameters. Fig. 2 depicts the optimized geometry of the free molecule together with its optimized bond lengths. Note that in the gas-phase the molecule is not in the zwitterionic form. Raman intensities were calculated at the B3LYP/6-31G** level of theory with the Dalton [64] program. Additional calculations on the crystalline L-methionine were performed with the CRYSTAL [65,66] package at the

Fig. 3. Calculated structure of the free L-methionine molecule indicating the formation of two distinct C=O bond lengths of the carbonyl group. Note that the hydrogen of the COOH is transferred to the NH₂ group in the crystal.

B3LYP/6-21G level of theory. The k -points were selected according to the procedure of Monkhorst and Pack [67] for an $8 \times 8 \times 8$ mesh. The experimental crystal structure [21,22] was the starting point for the ion positions optimization (Fig. 3). The Raman frequencies were determined at the Γ point.

Differential scanning calorimetry measurements, DSC, were performed using a Netzsch device between 190 and 360 K on powder samples mounted into a standard aluminum pan, using different scan speeds on cooling and heating.

Heat capacity measurements were performed using a Quantum Design PPMS equipped with the Quantum Design Heat Capacity option between 5 K $< T < 390$ K. Grains of the powder sample of L-methionine were mixed with the contact agent Apiezon H on the sample platform. To confirm the reproducibility of the data, three measurements were performed with each sample as follows: on heating starting from room temperature to 390 K, then after the sample was cooled down to 5 K, a second set of data was collected from 5 to 390 K, and finally after cooling the sample to 100 K, data were collected between 100 and 390 K.

3. Results

3.1. Revealing molecular motions and proving conformational changes in the L-methionine molecules by correlating polarized Raman to DFT calculations

As shown in Fig. 4, computed Raman spectra and the experimental results compare favorably in terms of the overall spectral profile. The differences between the calculated geometrical features and vibrational frequencies of the free L-methionine molecule compared to those in the crystal are caused by formation of the H-bonds in the latter. In the crystal the hydrogen of the COOH is transferred to the NH₂ group, and in this zwitterionic form the C=O bond lengths of the carbonyl group are almost identical with a bond length around 1.3 Å, see Fig. 3. In contrast, for the free molecule in the gas-phase, two distinct bond lengths of 1.21 Å and 1.36 Å (see Fig. 2). All other geometrical features are found to be rather similar. As a consequence of the formation of the zwitterion in the crystal that possesses COO[−] and NH₃⁺ groups, and forms hydrogen bonded networks, the differences in the calculated vibrational frequencies

of the free molecule compared to those of the crystal are mostly concerned with the vibrations in which the COOH and NH₂ groups participate. In particular the NH stretch and NH₂ bending modes are affected. The COO–H stretch and NH stretch modes are shifted from 3800 to 3500 cm^{−1} region for the free molecule to 3100–2800 cm^{−1} region in the crystal. Also, in the crystal, additional NH₂ bending modes appear in 2000–1500 cm^{−1} region. Other small differences are found due to differences in basis set.

Now we turn to the detailed analysis of the different spectral regions of the Raman scattering data.

Fig. 5 presents the Raman spectra of L-methionine monocrystal in the Z(Y)Z scattering geometry for selected temperatures (a) from 17 to 295 K and (b) from 300 to 375 K. For amino acid crystals peaks appearing at $\omega < 200$ cm^{−1} are, in general terms, assigned as lattice modes, except for bands associated with torsion of the CO₂[−] group, $\tau(\text{CO}_2)$ [40]. For L-methionine the lowest wavenumber band, at ~ 129 cm^{−1}, which is also the most intense peak in the spectrum taken at 17 K, is assigned to an external (lattice) mode and the band

Fig. 4. Comparison of calculated and measured Raman spectra of L-methionine.

Fig. 5. Raman spectra of L-methionine crystal in the region between 600 and 100 cm^{-1} for 17–295 K (a) and for the 300–375 K (b) temperature ranges.

at $\sim 190 \text{ cm}^{-1}$ to the $\tau(\text{CO}_2^-)$. It is possible that for $\omega < 100 \text{ cm}^{-1}$ other bands associated with external modes exit, however due to the shoulder of the laser line, occurring just below 90 cm^{-1} , all opportunity to observe Raman peaks in this region was eliminated. In spite of this, and as described below, our understanding of the already reported phase transition at 307 K was not jeopardized, and in addition we were able to evidence changes in the molecular conformation at lower temperatures.

In the low temperature range, the positions of the band associated with $\tau(\text{CO}_2^-)$ as well as the nearby peak located at 196 cm^{-1} (at room temperature) change significantly from 17 to 295 K. Moreover, the low intensity band at 238 cm^{-1} , assigned as torsion of CC, $\tau(\text{CC})$, and the CSC bending, $\delta(\text{CSC})$, located at 281 cm^{-1} , can be observed. Bands of low intensity observed at $\sim 350 \text{ cm}^{-1}$ are assigned as skeleton bending, $\delta(\text{skel})$, and the peak observed at $\sim 448 \text{ cm}^{-1}$ as the NH_3^+ torsion, $\tau(\text{NH}_3)$ [14]. In some cases the study of this band is interesting because it gives information on the H-bonds of the system. For instance, a comparative investigation of the behavior of $\tau(\text{NH}_3)$ with pressure showed that for L-threonine and taurine the wavenumber increases with pressure, while for L-alanine the wavenumber of $\tau(\text{NH}_3)$ decreases on compression [68]. In L-alanine, this behavior was understood as a deformation of the H-bond network under applied pressure. On the other hand, the observation of wavenumber change of the $\tau(\text{NH}_3)$ on DL-serine was associated with molecular rearrangements in the unit cell through changes in the H-bonding interactions [6]. In the present study, however, no great change is observed on the wavenumber of the $\tau(\text{NH}_3)$ mode, indicating subtle, if any, changes related to the NH_3^+ -group in the 17–300 K interval. Such behavior, in fact, is very similar to that observed for α -glycine, where the wavenumber of the $\tau(\text{NH}_3)$ mode is almost constant in a 300 K interval, indicating no significant change in H-bond length [69].

In Fig. 5 the peak observed at 543 cm^{-1} is associated to the CO_2 rocking vibration, $r(\text{CO}_2)$. For this band a linear behavior of the wavenumber with temperature is also observed, although the variation of wavenumber in the temperature interval investigated is greater than the variation of wavenumber of $\tau(\text{NH}_3)$

mode. As mentioned previously, in the Z(YY)Z scattering geometry, the band at 129 cm^{-1} , marked by a down arrow in Fig. 5, is the most intense in the spectrum of L-methionine in the spectral range from 600 to 100 cm^{-1} and, on heating, it is still visible up to $\sim 307 \text{ K}$. For temperatures higher than 309 K, however, this band is no longer observed. Note that the other bands, although lower in intensity, remain clearly visible up to $T = 375 \text{ K}$ – compare also, the intensity of the band assigned to the lattice vibration and the band associated with $r(\text{CO}_2)$. Therefore we can conclude that the intensity decrease leading to the disappearance of the peak at 129 cm^{-1} is intrinsic to this band. So, this particular band, associated with a lattice mode of the crystal, vanishes when $T \rightarrow 307 \text{ K}$, thus implying that the symmetry of the unit cell is changing. In other words, the phase transition observed in the previous calorimetric measurements on powder samples can be related to changes of this particular lattice mode in the crystal.

At lower temperatures, no further changes in the Raman spectra that could be associated with structural or even conformational modification in L-methionine crystal were observed in the spectral region depicted in Fig. 5. Therefore, in order to analyze the behavior of the material at low temperatures, i.e. for $T < 300 \text{ K}$, we have measured the Raman spectra at Z(YY)Z and Z(YX)Z scattering geometries for other spectral regions. Fig. 6(a) and (b) presents the Raman spectra of L-methionine monocrystal in the spectral range $1050\text{--}600 \text{ cm}^{-1}$ for selected temperatures in the Z(YY)Z and Z(YX)Z scattering geometries, respectively. This region is characterized by vibrations assigned to CS stretching, $\nu(\text{CS})$, CO_2^- wagging and bending, $\omega(\text{CO}_2^-)$ and $\delta(\text{CO}_2^-)$, and to CH_3 and NH_3^+ , rocking, $\rho(\text{CH}_3)$ and $\rho(\text{NH}_3^+)$ [16,18]. As expected the Z(YY)Z spectra arising from a totally symmetric representation have a signal-to-noise relationship higher than the spectra from the other scattering geometry. Besides, an intensity increase is observed for both set of spectra on cooling, including the bands at 721 and 766 cm^{-1} , which are associated with the $\nu(\text{CS})$, and the bands observed at 806 and 877 cm^{-1} , associated with the $\nu(\text{CC})$. No further notable changes are observed in this spectral region.

Fig. 6. Low temperature Raman spectra of L-methionine crystal in the region between 1050 and 600 cm^{-1} for the Z(YY)Z (a) and Z(YX)Z (b) scattering geometries.

Fig. 7 shows the Raman spectra of L-methionine monocrystal for selected temperatures in the two selected scattering geometries for the spectral range 1550–1200 cm^{-1} , where, among others, the bands associated with the CH_2 , CH and CH_3 symmetric bending, $\delta(\text{CH}_2)$, $\delta(\text{CH})$ and $\delta_s(\text{CH}_3)$ are expected to be observed [16,18]. Although an apparent peak splitting between 1300 and 1400 cm^{-1}

seems to occur on cooling, at 295 K this spectral region can be already fitted by four peaks at 1321, 1338, 1354 and 1363 cm^{-1} in the Z(YY)Z scattering geometry and at 1323, 1341, 1356 and 1365 cm^{-1} in the Z(YX)Z scattering geometry. Nevertheless, it is very unambiguous that the peak assigned to the $\delta_s(\text{CH}_3)$, observed at 1355 cm^{-1} , becomes clearly evident at about 240 K in the Z(YY)Z

Fig. 7. Low temperature Raman spectra of L-methionine crystal in the region between 1550 and 1200 cm^{-1} for the Z(YY)Z (a) and Z(YX)Z (b) scattering geometries.

Fig. 8. Low temperature Raman spectra of L-methionine crystal in the region between 3050 and 2800 cm^{-1} for the Z(YY)Z (a) and Z(YX)Z (b) scattering geometries. The inset presents the representation of an L-methionine molecule (hydrogen atoms not shown) with atom numbering – see text.

scattering geometry and at about 220 K in the spectrum taken in the Z(YX)Z geometry, whereas at about 140 K, the peak at 1455 cm^{-1} , assigned as $\delta_{\text{as}}(\text{CH}_3)$, is visibly distinguished (Table 1).

Fig. 8 presents the Raman spectra of L-methionine crystal in the high wavenumber spectral region for both Z(YY)Z and Z(YX)Z scattering geometries for selected temperatures. No band was observed for wavenumbers higher than 3200 cm^{-1} . This region, dominated by the CH and CH_2 stretching vibrations, gives important insight on the conformation of the molecules in the unit cell as already pointed out in Ref. [6]. In our study, changes around 150 K were observed for several bands. In particular, it is observed that a band at $\sim 2900 \text{ cm}^{-1}$ appears on cooling in the Z(YY)Z spectra taken at 140 K (down arrow in Fig. 8(a)). Moreover, in both geometries we observe the splitting of the bands centered around 2925 cm^{-1} . Even if one can argue that these bands are already present at room temperature in both geometries, the increase of their intensity is remarkable. Therefore these observations reveal that conformational changes are triggered at 140 K, and that such changes can be considered as precursors of the phase transition observed at 307 K. In other words, on heating the molecules begin to reorganize and gain motional freedom, and as the temperature approaches the transition temperature, the distance range of this cooperation increases, leading to correlated interactions, and consequently, a sudden change of order at the phase transition temperature.

Finally, in Fig. 9 we present the wavenumber vs. temperature plot for the modes appearing in the Z(YY)Z scattering geometry in the spectral range $550\text{--}100 \text{ cm}^{-1}$ in order to illustrate the discontinuity of a mode at 307 K.

3.2. Thermal behavior: analyzing the structural phase transition and the conformational changes observed in L-methionine

As already mentioned, a phase transition at about 307 K in L-methionine, characterized by a very broad peak in the thermal

analysis in powder samples has been well documented [16,17]. However, neither the origin of such observation has been fully explored, nor has the reproducibility of the data been verified. Therefore, in order to better understand the origin of the conformational change observed in our Raman results, we have performed

Fig. 9. Wavenumber vs. temperature plot for modes of L-methionine in the spectral range $550\text{--}100 \text{ cm}^{-1}$ for the Z(YY)Z scattering geometry showing the disappearance of a band associated with the lattice mode at $\sim 307 \text{ K}$. The dashed line represents the temperature where the crystal undergoes a phase transition.

Fig. 10. Heat capacity, C_p , of L-methionine, showing that in the polycrystalline sample a broad transition is observed between 275 K and 325 K with a maximum at 307.0 ± 0.2 K.

a series of measurements using powder samples on an extended temperature range, $5 \text{ K} < T < 320 \text{ K}$.

Heat capacity (C_p) measurements on L-methionine powder samples are reported in Fig. 10; in agreement with the previous work [16], the broad and strong peak indicating a phase transition extended over a wide temperature ranging from about 250 to 325 K with a maximum at 306.9 ± 0.2 K was reproduced. However, unlike Ref. [16], a second anomaly was observed at lower temperatures about 220 K. This change in the C_p coincides to the temperature where the Raman peak assigned to the $\delta(\text{CH}_3)$ becomes noticeable in the Raman spectra. Therefore we can argue that due to the malleability of the CH_3 -group, structural relaxation can occur and that at ambient conditions L-methionine is either in a metastable state or shows differences in the local structure, yielding to some disorder in the form of a distribution of energy barriers that are altered on cooling. From this line of argument, we deduce that the changes observed in the thermodynamical response as well as in the vibrational modes of L-methionine at lower temperatures are related to variations of the interactions of the L-methionine molecules with the crystalline environment, similarly to the response previously observed in L-cysteine [12] and L-alanine [39].

The DSC data, shown in Fig. 11, were collected as follows. We started by heating the sample from 295 K up to 360 K (red curve) at 2 K/min and, subsequently, performing consecutive cooling and heating cycles with scan speeds of 2, 4, 8 and 16 K/min. The peak observed in the C_p -measurements was also clearly observed in the temperature range from 260 K to 320 K. By extrapolating to 0 K/min scan rate [70,71], a transition temperature of $T_0 = 307.0 \pm 0.3$ K was determined. By calculating the changes in enthalpy and entropy [70], further information on the mechanisms involved in the energetic change as well as the characterization of the transition as a first order phase-transition was obtained. All parameters are listed in Table 2. In Table 3 we furnish a summary of results obtained in the present work, as well as the type of material (crystal or powder) utilized in the experiments and the temperature of the phase transition observed by the several procedures.

4. Discussion

The modifications verified in the Raman spectra of L-methionine at low temperatures are very different from those observed through high pressure experiments [72]. For example, in the spectral region $650\text{--}700 \text{ cm}^{-1}$, where the bands associated with the bending of CCO [14] and/or rocking of the CH_2 [18] are expected, it was observed the appearance of a very intense band, up to now, one of

Fig. 11. DSC-curves of a 24.708 ± 0.001 mg of polycrystalline L-methionine at heating rates of 2, 4, 8 and 16 K/min between 260 and 360 K. A phase transition is observed around 300 K. Since the heating/cooling system needs some time for stabilization, the features observed at the beginning and at the end of the heating and cooling cycles, and represented with light gray color, are due to unstable oven temperatures only. (For interpretation of the references to color in the text, the reader is referred to the web version of the article.)

the most impressive phenomena reported in the literature related to proteic amino acids. Such an exceptional effect, however, is not seen here in the whole temperature range exploited. Therefore, the modification verified by Raman spectroscopy at high pressure is very particular to the change of that thermodynamic parameter.

On the other hand, even if interactions leading to conformational changes are small, the variation in the system energy are most likely observable in the C_p results, but expected to clearly manifest themselves in the dynamical measurements over a broad energy range. This is indeed what was demonstrated here by combining Raman scattering to thermal analysis. In other words, our low temperature measurements, presenting few and slight modifications in the Raman spectra of L-methionine crystal, do not indicate that this crystalline amino acid is undergoing a structural phase transition such as the one clearly observed under applied hydrostatic pressure conditions or even at $T = 307 \text{ K}$, when lattice modes changes are observed. Although no clear modification has been observed in the low wavenumber region where we expect to find lattice vibrations, and although no impressive modification has been detected in the $650\text{--}700 \text{ cm}^{-1}$ region, some changes were revealed in certain internal modes. This result can be indeed fully understood as follows. At room temperature, two conformers of L-methionine, A and B, are present in the crystalline unit cell [14]. In the A conformer the torsion of lateral chains is given by $\chi^1 = \tau[\text{N}-\text{C}^2-\text{C}^3-\text{C}^4] = 166^\circ$; $\chi^2 = \tau[\text{C}^2-\text{C}^3-\text{C}^4-\text{S}] = 174^\circ$; $\chi^3 = \tau[\text{C}^3-\text{C}^4-\text{S}-\text{C}^5] = 180^\circ$ (where C^2 is the α carbon, C^3 and C^4 are between the α carbon and sulfur atom, and C^5 is the carbon of the CH_3 group, see Fig. 1(b)). On the other hand, for conformer B the corresponding torsional angles are $\chi^1 = 166^\circ$; $\chi^2 = 174^\circ$; $\chi^3 = 74^\circ$. When the crystal undergoes the high temperature phase transition, at about 307 K, the monoclinic structure is conserved but the conformations of the molecules A and B are changed. Therefore the variation of intensities of the low frequency modes can be associated with such conformational modification. Additionally, as shown in Fig. 8, on cooling the band at $\sim 2925 \text{ cm}^{-1}$ gains in intensity between room temperature and 140 K, and from previously calculated vibrational spectra [18] of seven possible (stable) configurations for L-methionine, only one configuration allows for the observation of a band at 2926 cm^{-1} . This particular configuration is characterized by the torsional angles: $\chi^1 = -60^\circ$; $\chi^2 = 180^\circ$ and $\chi^3 = 180^\circ$. Therefore our results imply that on cooling at least one of the two conformers

Table 1Experimental values for $\omega(295\text{ K})$, $\omega(17\text{ K})$ and coefficients ω_0 and α for the linear fitting $\omega = \omega_0 + \alpha \cdot T$ for L-methionine crystal at low temperatures.

Z(YY)Z geometry				Z(YX)Z geometry				Assignments
$\omega(295\text{ K}) (\text{cm}^{-1})$	$\omega(17\text{ K}) (\text{cm}^{-1})$	$\omega_0 (\text{cm}^{-1})$	$\alpha (\text{cm}^{-1}/\text{K})$	$\omega(295\text{ K}) (\text{cm}^{-1})$	$\omega(17\text{ K}) (\text{cm}^{-1})$	$\omega_0 (\text{cm}^{-1})$	$\alpha (\text{cm}^{-1}/\text{K})$	
127	129	28.5	−0.0034	121	127	125.4	−0.0218	Lattice
					167	167.3	−0.0164	Lattice
178	190	191.7	−0.0424	171	186	187.7	−0.0514	$\tau(\text{CO}_2^-)$
196	216	218.1	−0.0724	192	215	216.6	−0.0795	Lattice
238	248	250.1	−0.0372					$\tau(\text{CC})$
281	282	282.1	−0.0055	282	279	279.0	0.0106	$\delta(\text{CSC})$
338	340	339.8	−0.0004	337	341	339.4	−0.0058	$\delta(\text{skel})$
				358	368	368.7	−0.0382	$\delta(\text{skel})$
354	362	362.9	−0.0262					$\delta(\text{skel})$
				419	421	420.1	−0.0026	$\tau(\text{NH}_3^+)$
447	451	451.4	−0.0117	448	451	452.5	−0.0136	$\tau(\text{NH}_3^+)$
543	549	549.1	−0.0178	546	551	551.2	−0.0176	$r(\text{CO}_2^-)$
645	645	645.8	−0.0025	645	646	645.8	−0.002	$\nu(\text{CS})$
659	660	660.0	−0.0014	660	660	660.6	−0.003	$\nu(\text{CS})$
682	685	685.2	−0.0111	683	685	685.3	−0.0098	$\omega(\text{CO}_2^-)$
721	725	725.9	−0.0151	722	725	726.2	−0.0142	$\delta(\text{CO}_2^-)$
766	770	771.2	−0.0159	766	771	771.5	−0.0171	$\nu(\text{CS})$
806	808	808.0	−0.0059	806	808	808.7	−0.0066	$\nu(\text{CC})$
877	880	880.2	−0.0115	877	880	880.6	−0.0117	$\nu(\text{CC})$
954	954	954.2	0.0005	950	953	953.6	−0.0118	$\rho(\text{CH}_3)$
989	996	997.4	−0.0278	989	997	998.7	−0.0269	$\rho(\text{NH}_3^+)$
1029	1035	1035.3	−0.0178	1032	1035	1035.6	−0.0162	$\nu_s(\text{CN})$
1245	1249	1250.4	−0.0137	1244	1249	1250.4	−0.02	$\delta(\text{CH}_2)$
1268	1269	1269.3	−0.0047	1270	1270	1570.3	−0.0023	$\delta(\text{CH})$
1321	1322	1321.4	0.0004	1323	1322	1321.8	0.0026	$\omega(\text{CH}_2)$
1338	1340	1340.6	−0.0105	1341	1342	1340.8	−0.001	$\omega(\text{CH}_2)$
1354	1355	1355.8	−0.0049	1356	1356	1355.9	0.0014	$\delta_s(\text{CH}_3)$
1363	1369	1370.7	−0.0266	1365	1368	1369.0	−0.0089	$\delta_s(\text{CH}_3)$
1412	1413	1413.0	−0.0014	1413	1412	1412.6	0.0033	$\nu(\text{CC})$
1425	1429	1429.8	−0.0191	1428	1428	1429.4	−0.0034	$\delta_{\text{as}}(\text{CH}_3)$
1444	1448	1448.1	−0.017	1444	1448	1448.1	−0.0161	$\delta_{\text{as}}(\text{CH}_3)/\delta_s(\text{CH}_2)$
1451	1456	1456.5	−0.0174	1452	1456	1456.4	−0.0147	$\delta_{\text{as}}(\text{CH}_3)/\delta_s(\text{CH}_2)$
1515	1520	1520.6	−0.0154	1517	1523	1524.6	−0.0253	$\nu_{\text{as}}(\text{CN})$
	1535	1535.0	−0.0198					
2837	2833	2833.5	0.0136					$\nu_s(\text{CH}_2)$
2858	2855	2855.2	0.0107	2858	2855	2854.9	0.0106	$\nu_s(\text{CH}_2)$
2917	2916	2915.5	0.0042	2917	2915	2914.8	0.0061	$\nu_{\text{as}}(\text{CH}_2)$
2921	2925	2925.2	0.0008	2923	2926	2925.4	0.0041	$\nu(\text{CH})$
		2929.5	−0.029			2929.0	−0.0216	
2939	2941	2941.4	−0.0102	2938	2941	2941.3	−0.0119	$\nu_{\text{as}}(\text{CH}_2)$
2961		2960.8	0.0013	2961	2966	2962.0	−0.0047	$\nu_{\text{as}}(\text{CH}_3)$
2971	2967	2966.9	0.0151	2869	2985	2965.1	0.0383	$\nu_{\text{as}}(\text{CH}_3)$
						2968.6	0.002	
2987	2985	2984.8	0.0065	2984	2992	2985.8	−0.0039	$\nu_{\text{as}}(\text{CH}_3)$
	2991	2990.9	0.0206					

Table 2Change in enthalpy (ΔH) and entropy (ΔS) derived by calculation of the area under the transition peaks for L-methionine crystals. According to [73], transitions with $\Delta S > 1\text{ J/mol K}$, can be considered as first order phase transitions.

Sample state	$\Delta T (\text{K/min})$	$T_{\text{onset}} (\text{K})$	$T_{\text{final}} (\text{K})$	$T_{\text{peak}} (\text{K})$	$\Delta S (\text{J/mol K})$	$\Delta H (\text{kJ/mol})$	Type of transition
Powder	2	243	332	307	2.4	7.7	First order
	4	238	332	307	2.4	8.0	
	8	253	335	307	2.5	8.0	
	16	263	340	307	2.4	7.8	

of L-methionine is changing conformation. Furthermore, the intensity changes observed in the $\delta(\text{CH}_3)$ around 220–240 K in the Raman data can be related to the transition observed by C_p measurements, which are extremely sensitive to the way atoms and molecules are

bond in a solid. The jump observed around 220 K resembles the one observed in glassy systems, further confirming that on cooling, the crystalline structure of L-methionine evolves from an arrangement possessing a large distribution of H-bonds to a more stable system.

Table 3

Summary of the results on L-methionine, showing the sample state (crystal or powder), the details of the procedure and the temperature of transition observed with the three different experimental techniques.

Technique	Sample state	Details of procedure	Temperature of transition (K)
Raman spectroscopy	Crystal	Heating	307.0 ± 0.2
Heat capacity	Powder	Heating	306.9 ± 0.2
DSC	Powder	Heating and consecutive cooling/heating cycles	307.0 ± 0.3

5. Conclusion

We can summarize our results as follows. In this report we have shown the Raman spectra of L-methionine crystal in the interval 100–3000 cm⁻¹ for two scattering geometries at several temperatures, explaining the origin of a phase transition previously reported at $T=307\text{ K}$ and most importantly evidencing a conformational change of the molecule which is triggered around 240–220 K and finally happens in the interval 160–140 K. Change in the intensities of some Raman bands and a jump in the C_p were associated with conformational changes of at least one of the two L-methionine conformers in the monoclinic unit cell. The high-temperature phase transition was related to a broad peak observed during the thermal analysis and to the disappearance of a lattice mode located at 129 cm⁻¹ and thus implying that the symmetry of the unit cell in the high temperature phase changes. Finally, all vibrations could be assigned from comparison with the DFT calculated spectrum, showing that such studies are adequate to model the observed Raman vibrations of the amino acid L-methionine.

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