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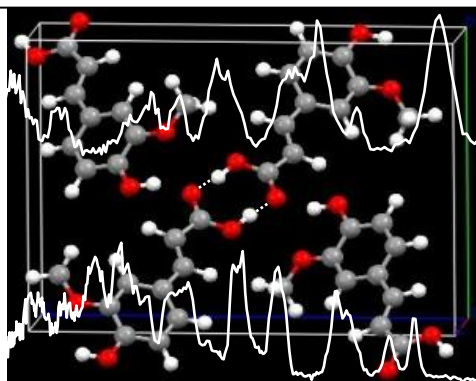
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**An Inelastic Neutron Scattering Study  
of Dietary Phenolic Acids**

M.P. M. Marques, L.A.E. Batista de Carvalho  
R. Valero, N.F.L. Machado and S.F. Parker

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# An Inelastic Neutron Scattering Study of Dietary Phenolic Acids

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M. Paula M. Marques,<sup>\*,a,b</sup> Luís A. E. Batista de Carvalho,<sup>a</sup> Rosendo Valero,<sup>a</sup>

Nelson F. L. Machado,<sup>a</sup> and Stewart F. Parker<sup>c</sup>

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The conformational preferences and hydrogen-bonding motifs of several potential chemopreventive hydroxycinnamic derivatives were determined by inelastic neutron scattering spectroscopy. The aim is to understand their recognized beneficial activity and establish reliable structure-activity relationships for this type of dietary phytochemicals. A series of phenolic acids with different hydroxyl/methoxyl ring substitution patterns were studied: *trans*-cinnamic, *p*-coumaric, *m*-coumaric, *trans*-caffeic and ferulic acids. Their INS spectra were completely assigned by theoretical calculations performed at the Density Functional Theory level, both for the isolated molecule, dimeric centrosymmetric species and the solid (using plane-wave expansion approaches). Access to the low energy vibrational region of the spectra enabled the identification of particular modes associated with intermolecular hydrogen-bonding interactions, which are the determinant of the main conformational preferences and antioxidant capacity in these systems.

## 1. Introduction

Phenolic acid derivatives constitute one of the most ubiquitous groups of plant metabolites, present in the human diet in significant amounts, which have long been known to display antioxidant properties, in particular due to the availability of phenolic hydrogens that allows them to act as H-donating radical scavengers. It is generally assumed that this radical scavenging capacity is closely related with the compounds' hydrogen or electron donating ability, as well as with the stability of the resulting phenoxyl radicals. Since oxidative damage caused by reactive oxidant species to vital biomolecules, such as DNA, lipids and proteins, is responsible for numerous pathological processes including inflammation, atherosclerosis, cancer and neurodegenerative disorders,<sup>1-4</sup> these compounds have lately been the object of intense research in Health Sciences.<sup>1,3,5-11</sup> While extensively used as antioxidant additives in both the food and pharmaceutical industries,<sup>12</sup> these types of molecules display a large variety of biological functions, from anti-inflammatory and antimutagenic to antitumoral and carcinogenesis modulation activities,<sup>2,13</sup> due to their ability to inhibit deleterious oxidative processes. In particular, cinnamic acid and its derivatives (e.g. caffeic acid phenethyl ester (CAPE)<sup>14</sup>) are known to display interesting antioxidant and selective antitumor properties,<sup>1,3,6,9,15-18</sup> and are widely distributed in plant tissues, thus being regularly consumed in the daily diet.<sup>19,20</sup> Ferulic acid, for instance, a methoxy-containing hydroxycinnamic acid, is abundant in plant cell wall components (e.g. cereals and seeds of coffee, apple, orange and peanut). Plant-derived polyphenolic compounds (phenolic acids, esters or amides) are therefore promising

candidates as preventive agents against oxidative stress-induced diseases, apart from being widely exploited as model systems for drug development. The evaluation of the antioxidant properties of phenolic derivatives of natural or synthetic origin, aiming at the prediction of their biochemical and pharmacological activity, is nowadays a key area of research in Medicinal Chemistry.

The prospective health benefits arising from the antioxidant capacity of phenolic derivatives are ruled by strict structure-activity relationships (SAR's), namely the number and relative position of the ring hydroxyl substituents, or the chemical nature and spatial orientation of the linker between the ring and the carboxylate/ester moiety. These, apart from determining their biological action, modulate their lipophilicity, systemic distribution and bioavailability in sites of oxidation within the cell. Also, conformational preferences such as flexibility, formation of hydrogen bonds – either intra- or intermolecular – and planar or skewed relative orientations of the substituent groups in the aromatic ring determine the biological properties of the systems. Within these polyphenolic molecules, the conformational preferences and relative stability are mostly determined by an extensive  $\pi$ -electron delocalization, as well as by the formation of medium to strong (O)H $\cdots$ O and/or (C)H $\cdots$ O intra- and intermolecular hydrogen bonds, which are long known to be relevant for the stability and function of numerous systems. These close contacts also account for the formation of phenolic dimeric species, shown to occur both in the solid state and in solution for this type of compounds.<sup>21,22</sup>

An accurate conformational analysis of this class of molecules is therefore essential in order to clearly understand their mode of action and to establish reliable SARs, which is an

essential step in the rational design of new phenolic-based chemopreventive/therapeutic agents, coupling an optimized (and perhaps selective) efficacy to a minimal toxicity. Such a goal can be attained through the combined use of spectroscopic techniques (*e.g.* vibrational spectroscopy) and theoretical approaches, in order to determine the compounds' most stable geometries, as well as their main conformational preferences and possible interactions with biological targets and receptors.

Inelastic Neutron Scattering (INS) spectroscopy is a technique that is well suited to the study of hydrogenous compounds such as phenolic systems. The neutron scattering cross-section of an element ( $\sigma$ ) is a characteristic of that atom and independent of its chemical environment. Since the value for hydrogen (80 barns) far exceeds that of all other elements (typically *ca.* 5 barns), the modes of significant hydrogen displacement ( $u_i$ ) dominate the INS spectra. For a mode at a given energy  $\nu_i$ , the intensity from a powdered sample obeys the simplified relationship,

$$S_i^*(Q^2, \nu_i) = \frac{(Q^2 u_i^2) \sigma}{3} \exp\left(-\frac{Q^2 \alpha_i^2}{3}\right) \quad (1)$$

where  $Q$  ( $\text{\AA}^{-1}$ ) is the momentum transferred from the neutron to the sample and  $\alpha_i$  ( $\text{\AA}$ ) is related to a weighted sum of all the displacements of the atom. Thus, not only the energies of the vibrational transitions (the eigenvalues,  $\nu_i$ ) but also the atomic displacements (the eigenvectors,  $u_i$ ) are available from experimental observation. This significantly enhances the information obtainable from the vibrational spectrum and adds to that from the complementary Raman and infrared vibrational spectroscopic methods. Additionally, the absence of selection rules and ready access to the region below  $400 \text{ cm}^{-1}$  enable the detection of some low frequency modes unavailable to these optical techniques. Since the spectral intensities can be quantitatively compared with those calculated by theoretical methods, by combining the INS results with quantum mechanical calculations it is possible to link molecular geometry with the experimental spectroscopic features, and produce a consistent conformation for the systems under investigation.

Furthermore, since vibrational INS intensities are strongly dependent upon the motions of hydrogens, INS spectroscopy displays a unique sensitivity to hydrogen interactions, being an excellent technique for studying H-bonding properties in solids. The out-of-plane (O-H $\cdots$ O) mode, for instance, involves almost exclusively the motion of the hydrogen atom relative to the undeformed molecule, thus being easily identified by INS. Additionally, unlike optical methods such as Fourier transform infrared (FTIR) or Raman spectroscopies, INS is known to yield intense features in the low wavenumber region of the spectrum, where intermolecular vibrations associated with hydrogen-bonding mostly occur. Nevertheless, to this date very few studies have been reported to probe the low-energy vibrational modes of phenolic systems, no INS data being found in the literature on this subject to the best of the authors' knowledge. A terahertz time-domain (THz-TD) analysis was performed for some hydroxycinnamic acids (including caffeic and ferulic acids) aimed at the detection and assignment of their low-frequency oscillators, in the solid state.<sup>23</sup>

The present work applies INS spectroscopy, coupled with Density Functional Theory (DFT)/Plane-wave (PW) calculations, to the study of a series of phenolic acids with different hydroxyl substitution patterns: *trans*-cinnamic (3-(phenyl)-2-propenoic, Cin), *p*-coumaric (3-(4-hydroxyphenyl)-2-propenoic or 3-hydroxycinnamic, *p*-C), *m*-coumaric (3-(3-hydroxyphenyl)-2-propenoic or 4-hydroxycinnamic, *m*-C), *trans*-caffeic (*trans*-3-(3,4-dihydroxyphenyl)-2-propenoic, CA) and ferulic (*trans*-3-(4-hydroxy-3-methoxyphenyl)-2-propenoic, FA) acids (Figure 1). These are structurally different regarding the number and localization of the hydroxyl (or methoxyl) ring substituent groups. Their main conformational characteristics were previously determined by Raman spectroscopy, combined with DFT calculations, for the isolated molecule, their dimeric species and the solution species.<sup>3,6,9,16,21,24-28</sup> Particular emphasis is given to the investigation of the hydrogen-bonding profile of this kind of systems, aiming at a better understanding and prediction of their antioxidant activity. Location of low-frequency modes due to H-bonding is envisaged, namely the (O $\cdots$ H) stretching. To this aim, deuteration of some of the compounds was carried out at specific sites (*e.g.* the carboxylic acid group), in order to better assign the oscillators involved in hydrogen interactions by comparison with the undeuterated samples.

In the analysis of the data particular attention was paid to the low wavenumber region of the spectra, where the most relevant changes are expected to occur. The theoretical INS transition intensities were obtained from the calculated normal mode eigenvectors, and the corresponding spectra were simulated using the aCLIMAX program.<sup>29</sup> Thus, the combined INS and DFT data for both the condensed state and the isolated molecules allow us to assign the spectral changes due to both intra- and intermolecular H-bonds, and to detect the characteristic (O-H $\cdots$ O) modes. This constitutes preliminary work for a future study of the dynamics of the proton transfer along hydrogen bonds within this type of phenolic systems, which is of the utmost relevance for numerous chemical and biochemical processes.<sup>30</sup>

## 2. Experimental

### 2.A. Reagents

*Trans*-Cinnamic, *p*-coumaric, *m*-coumaric, caffeic and ferulic acids (*p.a.* grade) were purchased from Sigma-Aldrich Química S.A. (Sintra, Portugal).

O-deuterated ferulic acid (at the carboxylic moiety) was prepared by solubilizing the solid compound in D<sub>2</sub>O (*ca.* 10% excess) and evaporating to dryness under vacuum (this process being repeated three times).

### 2.B. INS Spectroscopy

The INS spectra were obtained at the ISIS Pulsed Neutron and Muon Source of the Rutherford Appleton Laboratory (United Kingdom), on the TOSCA spectrometer. This is an indirect geometry time-of-flight, high resolution ( $(\Delta E/E)$  *ca.* 1.25 %), broad range spectrometer.<sup>31</sup> The samples (2-3 g) were wrapped in 4×4 cm aluminium foil sachets, which filled the beam, and placed

in thin walled aluminium cans. To reduce the impact of the Debye-Waller factor (the exponential term in equation (1)) on the observed spectral intensity, the samples were cooled to *ca.* 10 K. Data were recorded in the energy range -24 to 4000  $\text{cm}^{-1}$  and converted to the conventional scattering law,  $S(Q, \nu)$  vs energy transfer (in  $\text{cm}^{-1}$ ) through standard programs.

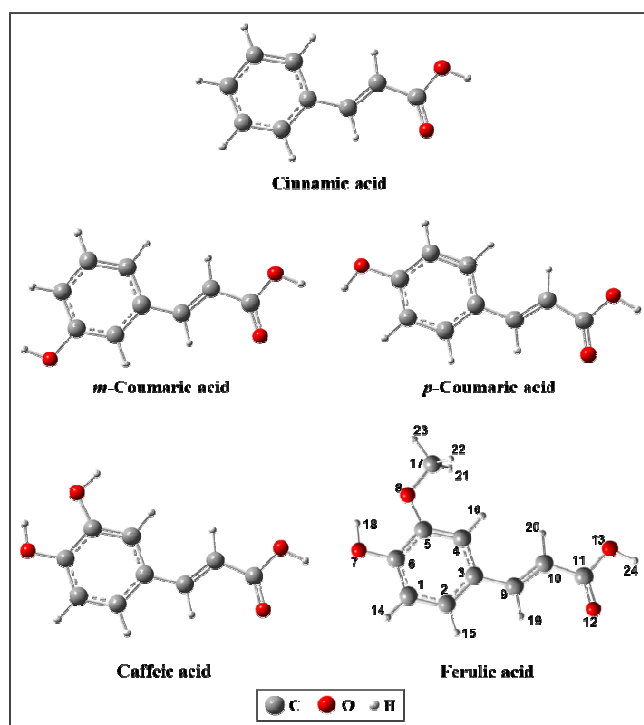
## 2.C. Quantum Mechanical Calculations

The quantum mechanical calculations were performed using the GAUSSIAN 03W program<sup>32</sup> (G03W) within the Density Functional Theory (DFT) approach, in order to properly account for the electron correlation effects (particularly important in this kind of conjugated systems). The widely employed hybrid method denoted by B3LYP, which includes a mixture of HF and DFT exchange terms and the gradient-corrected correlation functional of Lee, Yang and Parr,<sup>33,34</sup> as proposed and parameterized by Becke,<sup>35,36</sup> was used, along with the double-zeta split valence basis set 6-31G\*\*.<sup>37</sup> Molecular geometries were fully optimised by the Berny algorithm, using redundant internal coordinates:<sup>38</sup> the bond lengths to within *ca.* 0.1 pm and the bond angles to within *ca.* 0.1°. The final root-mean-square (rms) gradients were always less than  $3 \times 10^{-4}$  hartree.bohr<sup>-1</sup> or hartree.radian<sup>-1</sup>. No geometrical constraints were imposed on the molecules under study. The basis set superposition error (BSSE) correction for the dimerization energies was estimated by counterpoise (CP) calculations.

Natural bond orbital (NBO) analyses,<sup>39-41</sup> as implemented in G03W,<sup>42</sup> were carried out for the optimised geometries aiming at obtaining information on the nature of the interactions that determine conformer stability. Special attention was given to the Wiberg bond orders and energetics of the main interactions between "filled" (donor) Lewis-type NBOs and "empty" (acceptor) non-Lewis NBOs, evaluated by means of the 2<sup>nd</sup>-order perturbation theory (2<sup>nd</sup>-order stabilisation energies).

Calculations of the vibrational spectra were based on the optimised geometries. The harmonic vibrational wavenumbers, as well as their Raman activities and IR intensities, were obtained at the same level of theory. Frequencies above 500  $\text{cm}^{-1}$  were scaled by 0.9614<sup>43</sup> before comparing them with the experimental data, in order to correct for the anharmonicity of the normal modes of vibration. The theoretical INS transition intensities were obtained from the calculated normal mode eigenvectors of the quantum mechanical calculations, and the spectra simulated using the dedicated aCLIMAX program.<sup>29</sup> This program accommodates the impact of the external modes of extended molecular solids through the choice of a suitable value for  $\alpha_i$  (eq. (1)) – in the program this is the 'A-external' parameter and was chosen as 0.035 Å.

For the solid state, plane-wave expansions, as implemented in the PWSCF code from the Quantum Espresso package,<sup>44</sup> were used, based on DFT methods within the generalized gradient-corrected PBE functional.<sup>45</sup> The pseudopotentials employed were of the Vanderbilt ultrasoft type.<sup>46</sup> A cutoff energy of 35 Ry and a Monkhorst-Pack grid<sup>47</sup> of  $3 \times 3 \times 3$  were found sufficient to attain convergence. For ferulic and cinnamic acids, the atomic coordinates were fully optimized using the published crystal structures<sup>48,49</sup> as a starting point.



**Figure 1.** Schematic representation of the most stable calculated gas-phase conformers (B3LYP/6-31G\*\*) for the phenolic acids studied here. (The atom numbering is included).

## 3. Results and discussion

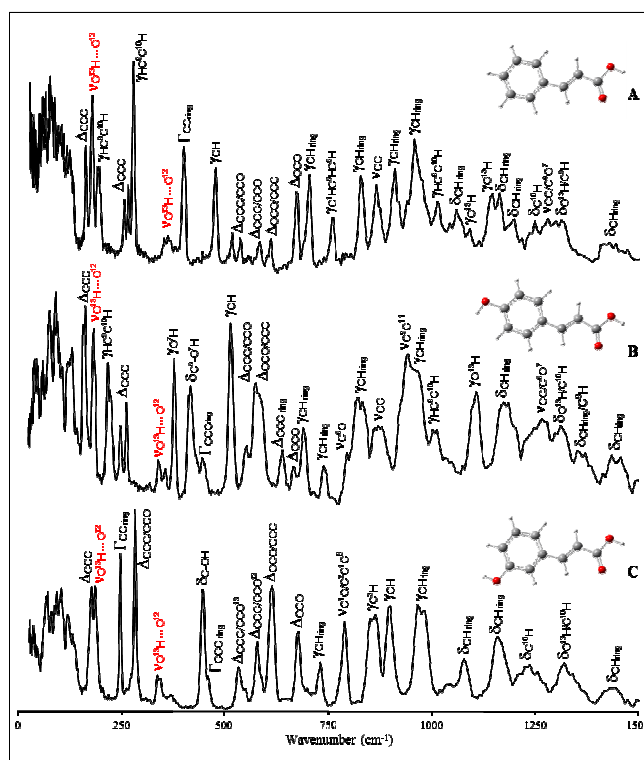
The compounds investigated here possess characteristic structural motifs that contribute to their biological activity: (i) the presence of electron-donating substituents in the aromatic ring (hydroxyl and/or methoxyl); (ii) the carboxylic moiety with an adjacent unsaturated C=C double bond, providing additional sites of attack for free radicals and also acting as an anchor to the lipid bilayer.

The most stable geometries for the phenolic acids under study were previously determined to display an *s-cis* conformation of the carboxylic group ( $\angle(\text{H}^{24}\text{O}^{13}\text{C}^{11}\text{O}^{12}) = 0^\circ$ , Figure 1), along with a *trans* orientation of the aromatic ring relative to the terminal carboxylate ( $\angle(\text{C}^3\text{C}^9\text{C}^{10}\text{C}^{11}) = 180^\circ$ ).<sup>6,21,24,27</sup> All energy minima have a planar geometry that favours the stabilizing effect of the  $\pi$ -electron delocalization between the ring and the C<sup>9</sup>C<sup>10</sup> and C<sup>11</sup>O<sup>12</sup> double bonds. A typical structural feature of cinnamic acids and their derivatives is the partial double bond character of (C<sup>3</sup>–C<sup>9</sup>) and (C<sup>10</sup>–C<sup>11</sup>) ( $d=145\text{--}147$  pm<sup>21,24,27</sup>), reflecting this effective electron delocalization between the aromatic ring, the unsaturated pendant chain and the carboxylic moiety. In fact, the Wiberg indexes calculated for this series of compounds vary within the range 1.11–1.13 and 1.05–1.06 for the (C<sup>3</sup>–C<sup>9</sup>) and (C<sup>10</sup>–C<sup>11</sup>) bonds, respectively. Moreover, for the dihydroxylated caffeic acid, an identical orientation of the phenyl OH substituents (coplanar with the ring) was found to be strongly favoured, since it minimizes steric repulsions and enables the formation of highly stabilising medium strength intramolecular (O)H  $\cdots$  O interactions ( $d_{(\text{O} \cdots \text{H})}=211$  pm<sup>24</sup>, 2<sup>nd</sup>-order  $\text{Lp}(\text{O}^{12}) \rightarrow \sigma^* \text{O}_{10}\text{H}_{11}$  stabilising energy= $11.3$  kJ mol<sup>-1</sup>).

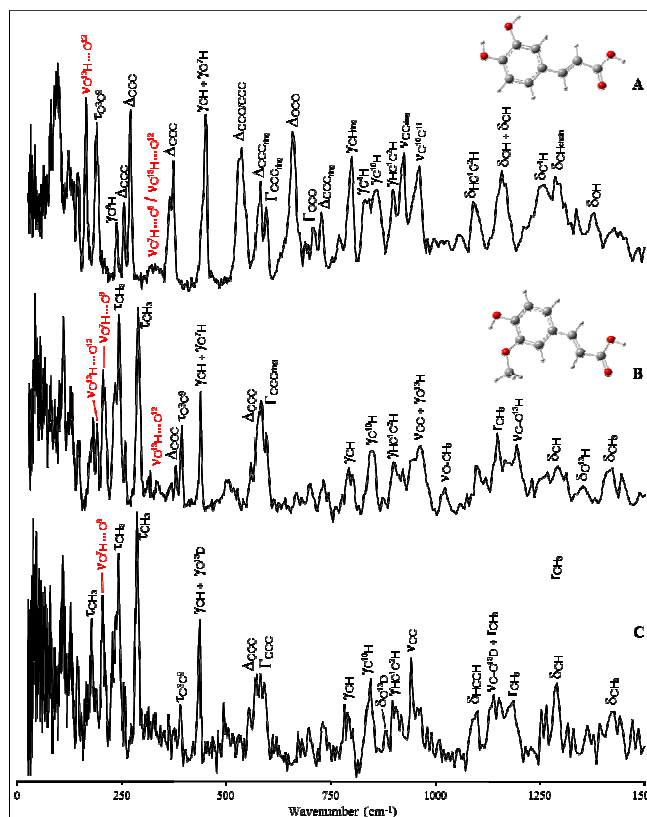


The experimental INS spectra for cinnamic and monohydroxylated cinnamic acids (*p*- and *m*-coumaric) are shown in Figure 3, while the corresponding data for the disubstituted caffeic and ferulic acids are depicted in Figure 4. Good quality spectra were obtained, allowing us to identify the main vibrational bands for these molecules, in the light of the calculated INS spectra at the B3LYP/6-31G\*\* level, for both the isolated molecule and the dimeric species (top-to-top dimers, Figure 5).

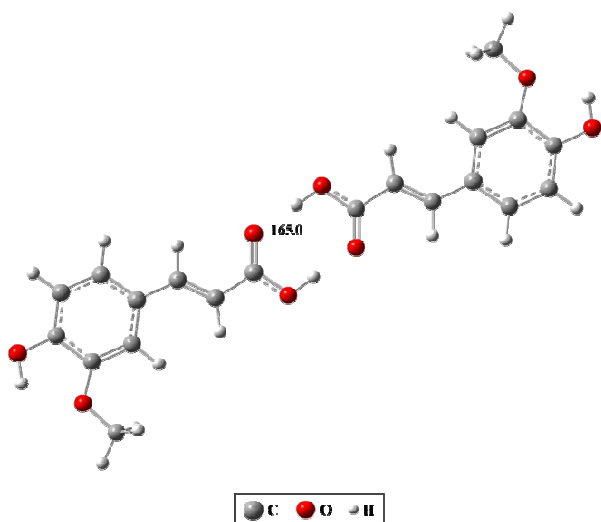
As a consequence, a different Davydov splitting pattern was observed for the monohydroxylated *p*- and *m*-coumaric acids – at 158/163, 218/224, 341/356, 414/419 and 575/588 cm<sup>-1</sup> for the former, and at 335/342, 577/588, 852/864 and 967/980 cm<sup>-1</sup> for the latter (Figure 3). This type of band splitting is due to the occurrence of distinct arrangements in the unit cell, leading to two crystallographically and energetically inequivalent species. This doubling of the unit cell was previously observed for *n*-alkanes<sup>50</sup> and linear polyamines.<sup>51</sup> Furthermore, the presence of several molecules in a crystal (or a packed solid state) generates an external force field, which is responsible for new vibrational modes – the so-called external modes (usually below *ca.* 200 cm<sup>-1</sup><sup>50,52</sup>). These are easily identified in the INS spectra presently obtained (Figures 3 and 4).



**Figure 3.** Experimental INS spectra (30 – 1500 cm<sup>-1</sup>) for cinnamic (A), *p*-coumaric (B) and *m*-coumaric (C) acids. (The atoms are numbered according to Figure 1).



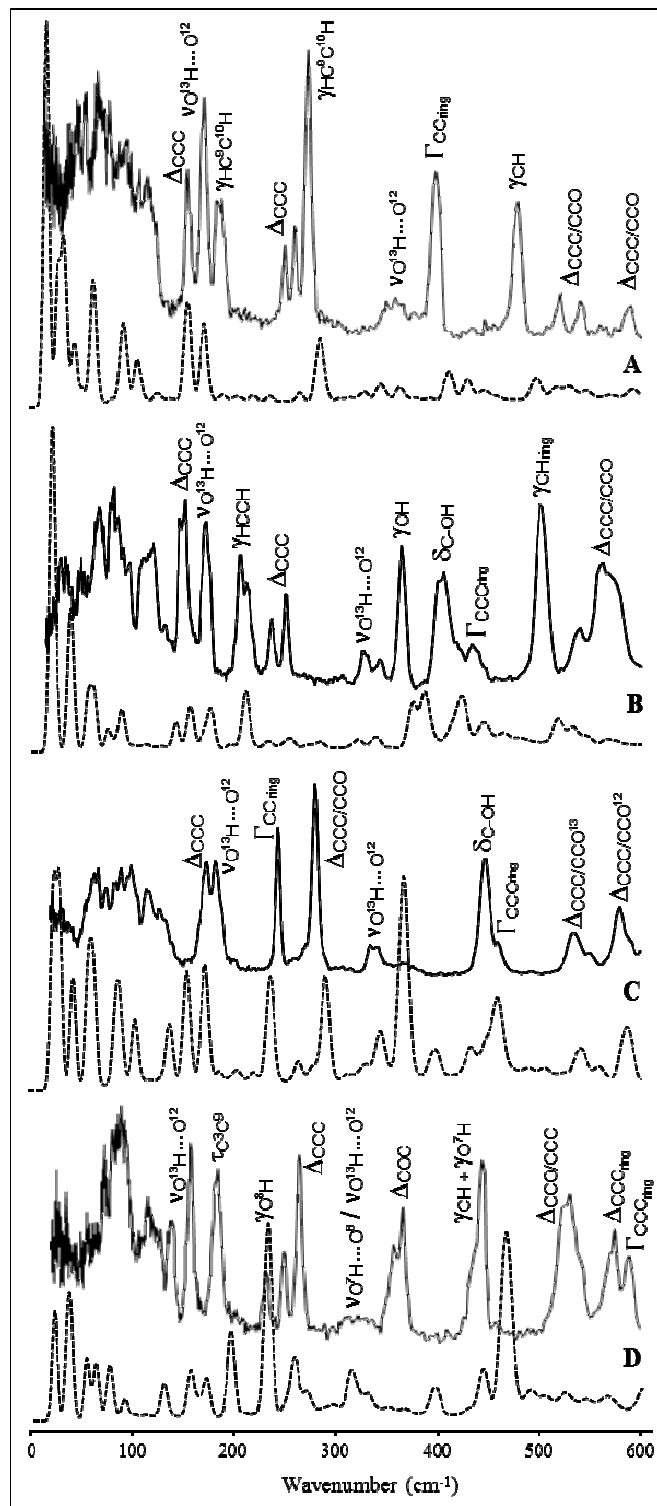
**Figure 4.** Experimental INS spectra (30 – 1500  $\text{cm}^{-1}$ ) for caffeic (A), ferulic (B) and  $\text{O}^{13}$ -deuterated ferulic (C) acids. (The atoms are numbered according to Figure 1).



**Figure 5.** Schematic representation of the most stable calculated structure (B3LYP/6-31G\*\*) for the centrosymmetric top-to-top dimer of ferulic acid. (Distances are represented in pm).

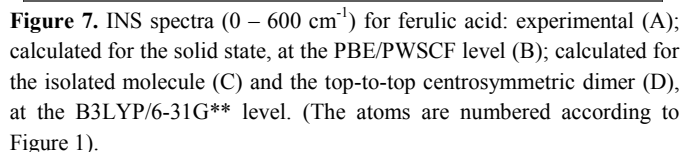
The main peaks reflecting H-bond interactions, which play an essential role in the conformational equilibrium of the phenolic derivatives under study, were unequivocally identified in their INS spectra, namely the features due to the ((H)O<sup>-</sup>H) stretching modes associated to the top-to-top (O<sup>13</sup>)H<sup>24...</sup>O<sup>12</sup> close contacts within the dimeric species present in the solid.<sup>22,27</sup> at *ca.* 180 and 350 cm<sup>-1</sup> (Figures 3, 4 and 6, Tables 1 and 2). Additionally, the occurrence of these centrosymmetric dimers formed through two intermolecular H-bonds between carboxylic groups (*d*<sub>(H...O)</sub> equal to 165 pm in the dimer (Figure 5) and 153 pm in the solid (Figure 2)), leads to an expected shift of the calculated carbonyl stretching vibration to lower wavenumbers as compared to the monomer (isolated molecule): *e.g.* 1701 *vs* 1742 cm<sup>-1</sup>, for ferulic acid, the former being in good accordance with the observed value at 1698 cm<sup>-1</sup> (Table 1). Moreover, the presence of intramolecular H-bonds between the hydroxyl hydrogen in the *para* position and the *meta* oxygen atom (from the hydroxyl in caffeic and the methoxyl in ferulic acids) were also detected.

Since the scattering by deuterium is roughly two orders of magnitude weaker than that from hydrogen, specific deuteration of the compounds is a very useful approach for an accurate assignment of the vibrational modes due to H-bonding interactions. For ferulic acid, spectra were acquired for both the hydrogenated solid and the O-deuterated molecule at the carboxylic moiety ( $\text{O}^{13}\text{D}^{24}$  for  $\text{O}^{13}\text{H}^{24}$  substitution, Figure 1), leading to the unequivocal assignment of the vibrations associated with the ( $\text{O}^{13}\text{H}\cdots\text{O}^{12}$ ) intermolecular close contacts (in the low wavenumber region) (Table 1, Figure 4(B) and (C)). As expected, deuteration alters the vibrational pattern according to two main factors: (i) increase in the mass of the oscillators involving the deuterated carboxylic group; (ii) disruption of the ( $\text{O}^{13}\text{H}^{24}\cdots\text{O}^{12}=\text{C}$ ) interactions (responsible for the formation of centrosymmetric dimeric species in the condensed phase, Figures



**Figure 6.** INS spectra (0 – 600 cm<sup>-1</sup>) for cinnamic (A), *p*-coumaric (B), *m*-coumaric (C) and caffeic (D) acids: experimental – solid line; calculated at the B3LYP/6-31G\*\* level, for the top-to-top centrosymmetric dimer – dotted line. (The atoms are numbered according to Figure 1).

the B3LYP/6-31G\*\* level (Table 1). The lower  $\text{O}^{13}\text{-H}$  stretching frequencies at the PW level are essentially due to the different density functionals employed, which yield a shorter hydrogen bond for the PBE functional and PW (153 pm) than for B3LYP/6-31G\*\* (165 pm). For further comparison, we have calculated the dimer frequencies at the PBE/6-31G\*\* level, *i.e.* with the same density functional used in the PW calculations. The results thus obtained evidence an H-bond distance of 155 pm and  $\nu(\text{O}^{13}\text{H})$  frequencies equal to 2485 and 2662  $\text{cm}^{-1}$ , with corresponding  $\nu(\text{O}^{13}\text{D})$  frequencies of 1868 and 1937  $\text{cm}^{-1}$ . In both cases, these values compare well with those obtained at the PW level, which are much lower than the B3LYP/6-31G\*\* wavenumbers: the shorter hydrogen bonds cause the  $\text{O}^{13}\text{H}$  distance to elongate, thus decreasing its stretching frequency. On the other hand, the much larger  $\gamma(\text{O}^7\text{H})$  frequency at the PW level is due to the solid-state structure, namely, to the fact that the hydrogen of the hydroxyl  $\text{O}^7\text{-H}$  bond of one molecule is involved in an H-bond with the oxygen of the  $\text{C}^{11}\text{-O}^{12}$  carbonyl of another molecule (Figure 2). This close contact is absent in the isolated monomer and dimer representations.



## Conclusions

The results shown here evidence that the structural stability and conformational behaviour of these systems relies on the balance between intra- and intermolecular hydrogen-type interactions, which are very accurately probed by inelastic neutron scattering techniques coupled with *ab initio* calculations. Access to the low frequency region of the spectrum unveiled the dependence of the vibrational modes occurring in this energy range relative to the molecular packing in the condensed phase, namely through the detection of characteristic Davydov splittings and of vibrational modes assigned to intermolecular H-bonds.

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Table 1. Main experimental and calculated INS wavenumbers (cm<sup>-1</sup>) for caffeic and ferulic acids.

CA			FA				<sup>a</sup> FA <sub>deut</sub>			<sup>b</sup> Approximate description
<sup>c</sup> Exp	<sup>d</sup> Calc/mon	<sup>e</sup> Calc/dim	<sup>c</sup> Exp	<sup>d</sup> Calc/mon	<sup>e</sup> Calc/dim	Calc/PW	Exp	<sup>d</sup> Calc/mon	Calc/PW	
					3025	2602-2641		2208	1892–1918	$\nu(\text{O}^{13}\text{H}); \nu(\text{O}^{13}\text{D})$
					2926	2488-2489		2151	1867–1869	$\nu(\text{O}^{13}\text{H}); \nu(\text{O}^{13}\text{D})$
(1640)	1741	1702	1698	1742	1701	1614-1623	~1694	1734		$\nu(\text{C}=\text{O})$
(1614)	1624	1626	(1628)	1630	1629	1612-1614			1609–1616	$\nu(\text{C}^9=\text{C}^{10}) + (\text{C}=\text{C})_{\text{ring}}$
			(1601)	1595	1596	1582-1587			1581–1590	$\nu(\text{C}=\text{C})_{\text{ring}} + \nu(\text{C}^9=\text{C}^{10})$
1575 (1570)	1598	1587								$\nu(\text{C}=\text{C})_{\text{ring}}$
			1448	1446	1446	1451-1456	1447	1446	1447–1453	$\delta_{\text{a}}(\text{CH}_3)$
1433	1426	1428								$\delta(\text{O}^{13}\text{H})$
			1417 (1421)	1421	1420	1413-1416	1430	1421	1415–1417	$\delta_{\text{s}}(\text{CH}_3)$
1382 (1388)	1383	1375	1356 (1371)	1381	1376	1377-1380				$\delta(\text{O}^7\text{H}) + \delta(\text{O}^{13}\text{H}) + \nu(\text{CC})_{\text{ring}}$
1344 (1353)	1358	1374								$\delta(\text{O}^7\text{H}) + \delta(\text{O}^{13}\text{H}) + \nu(\text{CC})_{\text{ring}}$
1296 (1298)	1298	1293	1296	1304	1300	1295-1302	1295	1303	1296–1298	$\delta(\text{CH})_{\text{chain}}$
1262 (1265)	1248	1244	1263	1266	1264	1256-1262	1260	1264	1258–1268	$\delta(\text{C}^4\text{H})$
1217 (1220)	1225	1243								$\delta(\text{CH})$
			1197 (1194)	1191	1199	1184-1187				$\nu(\text{C}-\text{O}^{13}\text{H}) + \delta(\text{O}^7\text{H}) + \delta(\text{CH})_{\text{chain}}$
							1191	1192	1178–1188	$\text{r}(\text{CH}_3)$
1174 (1177)	1172	1174								$\delta(\text{O}^7\text{H}) + \delta(\text{CH})$
1162	1149	1148	1151	1147	1146	1147-1157	1160	1169	1157–1171	$\delta(\text{O}^7\text{H}) + \delta(\text{O}^8\text{H}) + \delta(\text{CH}); \text{r}(\text{CH}_3)$
							1142	1147	1154–1155	$\nu(\text{C}-\text{O}^{13}\text{D}) + \text{r}(\text{CH}_3)$
								1135	1135-1146	$\text{r}(\text{CH}_3)$
1098	1096	1094	1104 (1118)	1105	1107	1101-1109	1101	1108	1093–1110	$\delta(\text{HC}^1\text{C}^2\text{H})$
			1025	1029	1029	1019-1020				$\nu(\text{O}-(\text{CH}_3)) + \nu(\text{CC})_{\text{ring}}$
964	989	966	966		966	960-966	966			$\nu(\text{C}^{10}\text{C}^{11})$
927	954	945	~948	952	946	950-953	947	940	931–933	$\nu(\text{CC})_{\text{ring}} + \gamma(\text{O}^{13}\text{H})$
904	907	908	903	908	907	904-909	908	908	904–909	$\gamma(\text{HC}^1\text{C}^2\text{H})_{\text{op}}$
							887	883	762–765	$\delta(\text{O}^{13}\text{D}) + \nu(\text{C}^{10}\text{C}^{11})$
861	866	849	850 (867)	844	848	853-854	848	844	837–838	$\gamma(\text{C}^{10}\text{H})$
840	848	814		828	827	836-837				$\gamma(\text{C}^4\text{H})$
802 (802)	795	798	805		801	789-794	808	800	789–797	$\gamma(\text{CH})_{\text{ring}}$
775		788	794	798	799	807-808	798	797	806–807	$\Delta(\text{CCC})_{\text{ring}}$
731		751	735	732	736	746-754	737	728	730–738	$\Delta(\text{CCC})_{\text{ring}}$
712		712	702	723	713	700-702	708	715	695–702	$\Gamma(\text{O}^{12}\text{C}^{11}\text{O}^{13})$

ARTICLE							Journal Name			
662	649	657	669 (672)	625	673	661-696				$\Delta(\text{O}^{12}\text{C}^{11}\text{O}^{13})$
598		583	596	604	585	593-595	594	593	593-595	$\Gamma(\text{CCC})_{\text{ring}}; \delta(\text{C}^{11}\text{O}^{13}\text{D})$
			584	563	584	575-586	584	587	575-585	$\Gamma(\text{CCC})_{\text{ring}}$
			578 (576)	558	564	566-574	575	558	565-570	$\Delta(\text{CCC})_{\text{chain}}$
581	578	581	562 (565)	543	554	525-538	560	533	523-537	$\Delta(\text{CCC})_{\text{ring}}$
539	560	545								$\Delta(\text{C}^9\text{C}^{10}\text{C}^{11}) + \Delta(\text{O}^{12}\text{C}^{11}\text{O}^{13})$
			528	543	527	512-517			512-517	$\Delta(\text{C}^5\text{O}^8\text{C}^{17}) + \Delta(\text{C}^{10}\text{C}^{11}\text{O}^{13})$
532	521	527	508 (503)	515	510	494-503			491-494	$\Delta(\text{C}^{10}\text{C}^{11}\text{O}^{12}) + \Delta(\text{C}^3\text{C}^9\text{C}^{10})$
466	471	470	501	489	488	721-730	497	489	721-735	$\gamma(\text{O}^7\text{H})$
453										
444	449	445	440 (446)	450	450	442-443	438	446	442-443	$\gamma(\text{C}^1\text{H}) + \gamma(\text{C}^4\text{H}) + \gamma(\text{O}^7\text{H}); \gamma(\text{O}^{13}\text{D})$
374	395	397	393	398	399	395-397	393	395	395-397	$\tau(\text{C}^3-\text{C}^9)$
366			379	363	380	374-392				$\Delta(\text{C}^5\text{O}^8\text{C}^{17})$
~330		320	335		311	258-277				$\nu(\text{O}^{13}-\text{H}^{24} \cdots \text{O}^{12})$
~330		314	315			327-351				$\nu(\text{O}^7-\text{H}^{18} \cdots \text{O}^8)$
			290	281	282	284-294	293	281	284-291	$\tau(\text{CH}_3)$
272	278	272								$\Delta(\text{C}^4\text{C}^3\text{C}^9) + \Delta(\text{C}^9\text{C}^{10}\text{C}^{11})$
258	257	251	259		256	248-277	260			$\Delta(\text{C}^9\text{C}^{10}\text{C}^{11})$
240	225	233	244		226	246-247	245			$\gamma(\text{O}^8\text{H}); \tau(\text{CH}_3)$
			235	225	225	233-240	235	225	239-247	$\tau(\text{CH}_3)$
			207		222	195-197	206		194-197	$\nu(\text{O}^7-\text{H}^{18} \cdots \text{O}^8) + \Delta(\text{C}^5\text{O}^8\text{C}^{17})$
193	195	196								$\tau(\text{C}^3-\text{C}^9)$
166		165	191		217	156-181				$\nu(\text{O}^{13}-\text{H}^{24} \cdots \text{O}^{12})$
			181	187	188	183-186	181	187	183-186	$\tau(\text{CH}_3)$
			173		171	176				external libration
146		156	146		156	146	146			external libration
123		131	129				128			external libration
			110		113	115	110			external libration
~97		92	90 (92)		92	95	89			external libration
80		78	80	80	81	76	82	80	82	external libration; $\tau(\text{C}^5\text{O}^8)$
64		63	65	35	61	63	65	35	59	$\tau(\text{C}^3-\text{C}^9)$
			53							external libration
43		54	42		50	46				external libration

<sup>a</sup>Deuterated at O<sup>13</sup>. <sup>b</sup>Atoms are numbered according to Figure 1;  $\nu$  – stretching,  $\delta$  – in-plane deformation,  $\tau$  – rocking,  $\gamma$  – out-of-plane deformation,  $\Delta$  – in-plane deformation of skeleton atoms,  $\Gamma$  – out-of-plane deformation of skeleton atoms,  $\tau$  – torsion; *s* and *as* stand for symmetric and antisymmetric, while *ip* and *op* represent in-phase and out-of-phase modes. <sup>c</sup>Raman experimental data in parenthesis<sup>26</sup>. <sup>d</sup>For the isolated molecule (monomer), at the B3LYP/6-31G\*\* level, wavenumbers above 550 cm<sup>-1</sup> are scaled according to Ref. 43. <sup>e</sup>For the top-to-top centrosymmetric dimer, at the B3LYP/6-31G\*\* level, wavenumbers above 500 cm<sup>-1</sup> are scaled according to Ref. 43. For the solid, at the PBE/PWSCF level.

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**Table 2.** Main experimental and calculated INS wavenumbers (cm<sup>-1</sup>) for cinnamic, and *p*- and *m*-coumaric acids.

Cin			<i>p</i> -C		<i>m</i> -C		<sup>a</sup> Approximate description
Exp	<sup>b</sup> Calc/mon	<sup>b</sup> Calc/PW	Exp	<sup>b</sup> Calc/dim	Exp	<sup>b</sup> Calc/dim	
1650	1643	1620-1638		1626			$\nu(\text{C}^9=\text{C}^{10})$
1560	1567	1586-1592		1562			$\nu(\text{CC})\text{ring} + \nu(\text{C}=\text{O})$
~1430	1450	1476-1480	1437/1457	1431	1435	1429	$\delta(\text{CH})\text{ring}$
			1364	1351			$\delta(\text{CH})\text{ring} + \delta(\text{C}^9\text{H})$
1319	1319	1295-1309		1332			$\delta(\text{O}^{13}\text{H}) + \delta(\text{C}^9\text{H})$
1303	1312	1281-1290	1315	1318	1320	1318	$\delta(\text{O}^{13}\text{H}) + \delta(\text{C}^{10}\text{H})$
1283	1271	1260-1266	1268	1269			$\nu(\text{CC})\text{ring}; \nu(\text{C}^6\text{O}^7)$
1252	1226	1202-1204			1237	1226	$\delta(\text{C}^{10}\text{H})$
1203	1205	1162-1164					$\delta(\text{C}^1\text{HC}^2\text{HC}^4\text{HC}^5\text{H})$
1168	1186	1145-1149	1176/1188	1187	1160	1178	$\delta(\text{CH})\text{ring}$
1148	1125	1034-1038		1125			$\gamma_s(\text{O}^{13}\text{H})$
1095	1089	1006-1022	1106	1089			$\gamma_a(\text{O}^{13}\text{H})$
1064	1077	988-989			1076	1075	$\delta(\text{CH})\text{ring}$
1018	1014	971-972	1009	1010			$\gamma(\text{HC}^9\text{C}^{10}\text{H})$
					967/980	985	$\gamma(\text{C}^4\text{HC}^5\text{HC}^6\text{H})$
962	971	959-961	969	978			$\gamma(\text{C}^1\text{HC}^2\text{HC}^4\text{HC}^5\text{H})$
			943	949			$\nu(\text{C}^{10}\text{C}^{11})$
913	922	915-916			897	902	$\gamma(\text{C}^2\text{HC}^6\text{HC}^4\text{HC}^{10}\text{H})$
			881	894			$\gamma(\text{C}^{10}\text{H})$
871	837	845-848	864	841			$\nu(\text{C}^3\text{C}^9) + \nu_s(\text{C}^2\text{C}^3\text{C}^4)$
					852/864	869	$\gamma(\text{C}^3\text{H})$
833	834	827-834	834	843			$\gamma(\text{CH})\text{ring}$
			823	815			$\gamma(\text{CH})\text{ring}$
			798	787			$\nu(\text{C}^6\text{O}) + \nu_s(\text{C}^1\text{C}^6\text{C}^5)$
					789	777	$\nu(\text{C}^1\text{O}) + \nu_s(\text{C}^2\text{C}^1\text{C}^6)$
762	771	760-763					$\gamma(\text{C}^1\text{HC}^6\text{HC}^5\text{H})$
			741	755	729	721	$\gamma(\text{CH})\text{ring}$
706	703	700-704	692	695			$\gamma(\text{CH})\text{ring}$
676	675	668-698	668	659	675	678	$\Delta(\text{OCO})$
			640	653			$\Delta(\text{CCC})\text{ring}$
613	600	611-612			613	604	$\Delta(\text{O}^{12}\text{C}^{11}\text{O}^{13}) + \Delta(\text{CCC})\text{ring}$
586	571	582-586	575/588	582	577/588	582	$\Delta(\text{C}^3\text{C}^9\text{C}^{10}) + \Delta(\text{C}^{10}\text{C}^{11}\text{O}^{12})$
					547	547	$\Delta(\text{C}^2\text{C}^1\text{C}^6) + \Delta(\text{C}^3\text{C}^4\text{C}^5)$
539	538	519-547	553	540	532	540	$\Delta(\text{C}^2\text{C}^3\text{C}^9) + \Delta(\text{C}^3\text{C}^9\text{C}^{10}) + \Delta(\text{C}^{10}\text{C}^{11}\text{O})$
521			516	539			$\gamma(\text{CH})$
479	508	476-484					$\gamma(\text{CH})$
402	422	397-400	448	444	459	472	$\Gamma(\text{C}^1\text{C}^2\text{C}^4\text{C}^5)$
			434	438	446	458	$\delta(\text{C}-\text{OH}) + \Delta(\text{CC}^3-\text{C}^9)$
			414/419	410			$\delta(\text{C}^6-\text{O}^7\text{H})$
395sh	413	374-375					$\Delta(\text{C}^2\text{C}^3\text{C}^9)$
			378	392	370	379	$\gamma(\text{O}^7\text{H})$
363	369	362-363	341/356	352	335/342	356	$\nu(\text{O}^{13}-\text{H}^{24} \cdots \text{O}^{12})$

280	302	276-280					$\gamma(\text{H}^{19}\text{C}^9\text{C}^{10}\text{H}^{20})$
267	321	268-269	263	301	282	285	$\Delta(\text{CC}^3\text{-C}^9) + \Delta(\text{C}^{10}\text{C}^{11}\text{O}^{13})$
258	287	260-261	249	270			$\Delta(\text{C}^9\text{C}^{10}\text{C}^{11})$
					246	241	$\Gamma(\text{CCC})\text{ring}$
196	205	189-199	218/224	232			$\gamma(\text{H}^{19}\text{C}^9\text{C}^{10}\text{H}^{20})$
181	187	169-195	184	182	185	178	$\nu(\text{O}^{13}\text{-H}^{24}\cdots\text{O}^{12})$
166	175	168-169	158/163	158	177	160	$\Delta(\text{C}^4\text{C}^3\text{C}^9\text{C}^{10})$
127	121	126	132		131	119	external libration
120	118	122	122		119	116	external libration
107	98	111	108	106	103	98	external libration
					93		external libration
81	77	82	91	88	79	76	external libration + $\tau(\text{C}^3\text{-C}^9)\text{op}$
			78	69			external libration
68	67	71			68	57	external libration
60	49	62	60	60			external libration
	39	46	46	47	36	35	external libration + $\tau(\text{C}^3\text{-C}^9)\text{ip}$
			40	41			external libration

<sup>a</sup>Atoms are numbered according to Figure 1;  $\nu$  – stretching,  $\delta$  – in-plane deformation,  $r$  – rocking,  $\gamma$  – out-of-plane deformation,  $\Delta$  – in-plane deformation of skeleton atoms,  $\Gamma$  – out-of-plane deformation of skeleton atoms;  $s$  and  $a$  stand for symmetric and antisymmetric. <sup>b</sup>For the isolated molecule (monomer), at the B3LYP/6-31G\*\* level, wavenumbers above 500 cm<sup>-1</sup> are scaled according to Ref. 43. <sup>c</sup>For the top-to-top centrosymmetric dimer, at the B3LYP/6-31G\*\* level, wavenumbers above 500 cm<sup>-1</sup> are scaled according to Ref. 43. For the solid, at the PBE/PWSCF level.

Notes and references

<sup>a</sup> Research Unit “Molecular Physical Chemistry”, Faculty of Science and Technology, University of Coimbra, 3004-535 Coimbra – Portugal.  
<sup>b</sup> Department of Life Sciences, Faculty of Science and Technology, University of Coimbra, 3001-401, Coimbra – Portugal. E-mail: pmc@ci.uc.pt.  
<sup>c</sup> ISIS Facility, SFTC Rutherford Appleton Laboratory, Chilton, Didcot, OX110QX – United Kingdom.

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## Table of Contents

**An Inelastic Neutron Scattering Study  
of Dietary Phenolic Acids**

M.P.M. Marques, L.A.E. Batista de Carvalho  
R. Valero, N.F.L. Machado and S.F. Parker

The conformational preferences of dietary phenolic compounds were determined by INS spectroscopy and DFT calculations, with the aim of understanding their beneficial activity and to establish reliable SARs. Access to the low energy spectral region enabled the identification of vibrational modes associated with H-type interactions, that are the determinant of the antioxidant capacity.

