

Thermodynamics of Linear Carbon Chains

Nathalia L. Costa^{1,*}, Keshav Sharma^{2,*}, Yoong Ahm Kim³, Go Bong Choi³, Morinobu Endo⁴,
Newton M. Barbosa Neto^{5,†}, Alexandre R. Paschoal^{1,‡} and Paulo T. Araujo^{2,§}

¹*Department of Physics, Federal University of Ceara, 60455-760 Fortaleza, Ceara, Brazil*

²*Department of Physics and Astronomy, University of Alabama, Tuscaloosa, Alabama 35487, USA*

³*Department of Polymer Engineering, School of Polymer Science and Engineering, and Alan G. MacDiarmid Energy Research Institute, Chonnam National University, 77 Yongbong-ro, Buk-gu, Gwangju 61186, Republic of Korea*

⁴*Faculty of Engineering, Shinshu University, 4-17-1 Wakasato, Nagano-shi 380-8553, Japan*

⁵*Institute of Natural Sciences, Graduate Program in Physics, Federal University of Para, 66075-110 Belem, PA, Brazil*



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Linear carbon chains (LCCs) are one-dimensional materials with unique properties, including high Debye temperatures and restricted selection rules for phonon interactions. Consequently, their Raman C-band frequency's temperature dependence is a probe to their thermal properties, which are well described within the Debye formalism even at room temperatures. Therefore, with the basis on a semiempirical approach we show how to use the C band to evaluate the LCCs' internal energy, heat capacity, coefficient of thermal expansion, thermal strain, and Grüneisen parameter, providing universal relations for these quantities in terms of the number of carbons atoms and the temperature.

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One-dimensional (1D) linear carbon chains (LCCs) are one of the simplest materials presenting properties associated with both molecules and solids [1–8], making them ideal for many electronic [9–13], mechanical [14,15], and thermal applications [2,16]. They exhibit *sp* hybridization and can be either semiconducting, known as polyynes with alternate single and triple bonds ($\cdots - \text{C} \equiv \text{C} - \text{C} \equiv \text{C} - \cdots$), or metallic, also known as cumulenes ($\cdots \text{C} = \text{C} = \text{C} = \text{C} = \cdots$) [8,12]. Historically, host-free LCCs have posed interesting questions regarding their stability [6,17,18] but recently, single-wall (SW), double-wall (DW), and multiwall (MW) carbon nanotubes (CNT) are considered ideal environments for fabricating stable LCCs with up to 6000 C atoms [1,5,9,11,15]. The polyyne structure is often the most stable configuration with band gaps typically around 2.13 eV [19,20].

The LCCs possess simple phonon structures presenting longitudinal and transversal modes when they are host-free [8]. When encapsulated, however, LCCs share their main axis with their host CNTs and this configuration inhibits transversal modes [20–23]. These phonons possess long mean free paths (~ 0.5 – $2.5 \mu\text{m}$) and lifetimes (~ 30 – 110 ps), making phonon-phonon interactions (ph-ph) inefficient in LCCs [8] when compared with other carbon materials [24–26]. The optical longitudinal mode (in-axis out-of-phase C-atom vibrations), so-called C band, with frequencies ω_{LCC} around 1850 cm^{-1} , is an easily trackable Raman spectroscopic signature and due to the weak nature of ph-ph it is an excellent probe to study mechanical and thermal properties [27–32]. The C band has been characterized in the literature via room-temperature [19,20] and pressure-dependent Raman spectroscopy [21–23]

but its use to obtain important mechanical [22] and thermal observables (this work) is a novel concept. Additionally, it is well known that ω_{LCC} is proportional to N^{-1} , where N is the number of carbon atoms [17,22].

Other complex chainlike structures have also been reported [33–39]. Chorro *et al.* [33], Rols *et al.* [34], and Cambedouzou *et al.* [35] showed that C_{60} carbon peapods inserted in SWNTs submitted to various pressures (P) and temperatures (T) could undergo, for example, polymerization (for pressures around 4 GPa), high orientational mobility (even for $T < 100$ K), and fast diffusional reorientations at $T = 200$ K. In another study, Bousige *et al.* [36] showed evidences of melting and liquid phases for C_{60} chains inside CNTs, revealing that such a quasi-1D system behaves as a harmonic crystal for $T \leq 550$ K (suggesting inefficient ph-ph), transitions to a liquid phase for $T \approx 650$ K and evolves to a complete disappearance of any structural correlations for $T \geq 850$ K. All these studies suggest that the interactions between C_{60} molecules and the CNTs' walls are mostly a second order effect. More recently, sulfur chains inside SWNTs have also demonstrated enhanced field-emission properties and outstanding gas-sensing properties [37–39]. In this context, the study of LCCs' thermodynamic properties could greatly benefit other encapsulated 1D systems.

The responses of materials to changing temperatures are closely related to the lattice thermal expansion (LTE) and anharmonic effects [27–32]. The former is associated with electron-phonon couplings (*e-ph*) and the latter is associated with ph-ph. Usually, ph-ph renormalize phonon energies, playing marginal roles in the understanding of

electronic and thermal properties of materials. When it comes to thermal properties, LCCs become particularly interesting and seem to be ahead of other materials such as fullerenes and transition metal dichalcogenides (TMDs) [40–43]. However, detailed studies of *e*-ph, ph-ph, and thermal effects on LCCs inside CNTs are still missing. Recently, theoretical results by Wong *et al.* reported the LCC's coefficient of thermal expansion (CTE) to be around $7.0 \times 10^{-5} \text{ K}^{-1}$ [44], which is much higher than that of CNTs [45], graphene [46], TMDs [47,48] and hexagonal boron nitride [49]. In fact, very little has been explored about LCCs' temperature-dependent phenomena, and important works available in the literature are theoretical simulations [4,8,41]. In 2016, Shi *et al.* studied LCCs inside DWCNT and showed that the C-band linewidth (lifetime) of LCCs increases (decreases) with increasing temperature [1]. They explained their findings in terms of ph-ph and interactions between LCCs and the DWCNTs' inner walls, despite their weak van der Waals interactions [22,33–35,50–52]. To the best of our knowledge, there is no quantitative temperature-dependent study on ω_{LCC} and its connection to thermodynamic observables.

In this Letter, the ω_{LCC} temperature dependence of distinct LCCs is used as a direct probe to their thermodynamic observables in a wide range of temperatures, showing that fundamental thermodynamic relations follow universal behaviors that only depend on T and N . Our findings show that these true one-dimensional systems are well described by the Debye model even for T at and above 300 K, indicating that ph-ph are negligible indeed.

The LCCs encapsulated by multiwalled CNTs (LCC@MWCNTs) were synthesized using arc discharge [19], while the LCCs encapsulated by double-walled CNTs (LCC@DWCNTs) were synthesized using catalyzed chemical vapor deposition [53,54]. The MWCNTs' purity with regard to nanoparticles is $\approx 80\%$ with average diameters (length) of 10.4 nm (2.3 mm). The DWCNTs' synthesis yielded over 95% of highly pure and crystalline DWCNTs with an outer diameter of ≈ 1.6 nm. The encapsulated LCCs were then obtained by submitting the DWCNTs to thermal treatment for 30 min at 1500°C in a graphite furnace containing high purity argon gas. The LCC@CNTs filling ratio is $\approx 80\%$ [19,53,54]. The samples were dispersed in acetone and sonicated for 2 h and then drop casted onto a Si wafer of $\approx 1 \text{ cm}^2$ area. Raman spectra were acquired with a $50\times$ objective in a backscattering geometry using an Andor iDus 401 series CCD coupled to a Shamrock spectrometer (1200 lines/mm grating; spectral resolution of $\approx \pm 0.5 \text{ cm}^{-1}$). Samples were resonantly excited with 532 (2.33) and 632 nm (1.96 eV) lasers under constant power density of $0.25 \text{ mW}/\mu\text{m}^2$.

The correlation between LCC's C-band frequency and N indicates LCCs with 36 to 86 carbon atoms in LCC@MWCNTs and LCCs with 35 to 39 carbon atoms in LCC@DWCNTs [1,7,11,17,21,23,55]. Representative

Raman spectra at 300 K and C-band spectra fitted with Lorentzian curves are shown in Figs. S1(a)–S1(c), Sec. S1 in the Supplemental Material [56]. The MWCNTs host six distinct LCCs and the DWCNTs host four distinct LCCs. For LCC@MWCNTs, each LCC is labeled LCC_i^{MW} ($i = 1, 2, 3, 4, 5$, and 6), where $i = 1$ ($i = 6$) represents the longest (shortest) LCC. Their respective room-temperature frequencies are $\omega_{\text{LCC}_1}^{\text{MW}} = 1796.0$, $\omega_{\text{LCC}_2}^{\text{MW}} = 1804.0$, $\omega_{\text{LCC}_3}^{\text{MW}} = 1839.0$, $\omega_{\text{LCC}_4}^{\text{MW}} = 1843.0$, $\omega_{\text{LCC}_5}^{\text{MW}} = 1854.0$, and $\omega_{\text{LCC}_6}^{\text{MW}} = 1863.0 \text{ cm}^{-1}$. Similarly, for the LCC@DWCNTs, each LCC is labeled LCC_i^{DW} (longest LCC) and LCC_4^{DW} (shortest LCC), whose frequencies are $\omega_{\text{LCC}_1}^{\text{DW}} = 1852.0$, $\omega_{\text{LCC}_2}^{\text{DW}} = 1856.0$, $\omega_{\text{LCC}_3}^{\text{DW}} = 1860.0$, and $\omega_{\text{LCC}_4}^{\text{DW}} = 1863.0 \text{ cm}^{-1}$. The T dependence of $\omega_{\text{LCC}_i}^{\text{MW}}$ and $\omega_{\text{LCC}_i}^{\text{DW}}$ shows that $\omega_{\text{LCC}}(T) = \omega_{\text{LCC}}^0 - (d^2\omega_{\text{LCC}}/dT^2)T^2$, where ω_{LCC}^0 is the frequency at 0 K [Figs. S2(a)–2(c), Sec. S1 in the Supplemental Material [56]]. In the case of inefficient ph-ph, $\Delta\omega_{\text{LCC}}(T) = \omega_{\text{LCC}}(T) - \omega_{\text{LCC}}^0$ [see Fig. 1(a)] will depend mostly on LTE, being directly linked to observables such as specific heat [$c_v(T)$], coefficient of thermal expansion—CTE [$\alpha(T)$], thermal strain (ϵ_T), and the Grüneisen parameter (γ_P) ($P = \text{constant pressure}$).

The literature [20–23,33–36,52] has shown that (i) 1D chains of atoms or molecules are independent of force fields in the CNTs' radial direction [33–36]; (ii) spectral features from the hosting CNTs and LCCs are independent, even under high pressures [20–23]. The CNTs provide conditions that are sufficient to stabilize the LCCs and inhibit transversal vibrations but not sufficient to entangle CNTs and LCCs properties; and (iii) interactions between distinct LCCs are not strong enough to affect their structures even at high pressures [20–23]. Therefore, transverse vibrations and mutual interactions between distinct LCCs, and LCCs and CNTs are second order effects. These, together with inefficient ph-ph, lead us to hypothesize that (i) LCCs are truly 1D Debye materials even at $T = 300 \text{ K}$ [note that most materials are well described within the Debye formalism only at very low temperatures ($T \ll 300 \text{ K}$) [57,58]]; (ii) any phonon frequency variation with T is connected solely with lattice expansion or contraction; and (iii) since ph-ph are inefficient, γ_P is T independent.

The Debye model does not take into account ph-ph and it is known to work very well for temperatures such that $T_D/T \rightarrow \infty$, where T_D is the Debye temperature. The Debye temperature is proportional to the maximum excitable phonon frequency obtained from phonon density of states or dispersion [57,58]. For LCCs, the maximum excitable phonon frequency is set by ω_{LCC} , resulting in $T_D = T_{\text{LCC}} = \hbar\omega_{\text{LCC}}/k_b$ between 2584 (longest LCC) and 2680 K (shortest LCC), where k_b is the Boltzmann constant and $\hbar = h/2\pi$ (h is Planck's constant). Under the Debye model formalism (DMF), the canonical partition function for LCCs is given by (Sec. S2 in the Supplemental Material [56] for additional details)

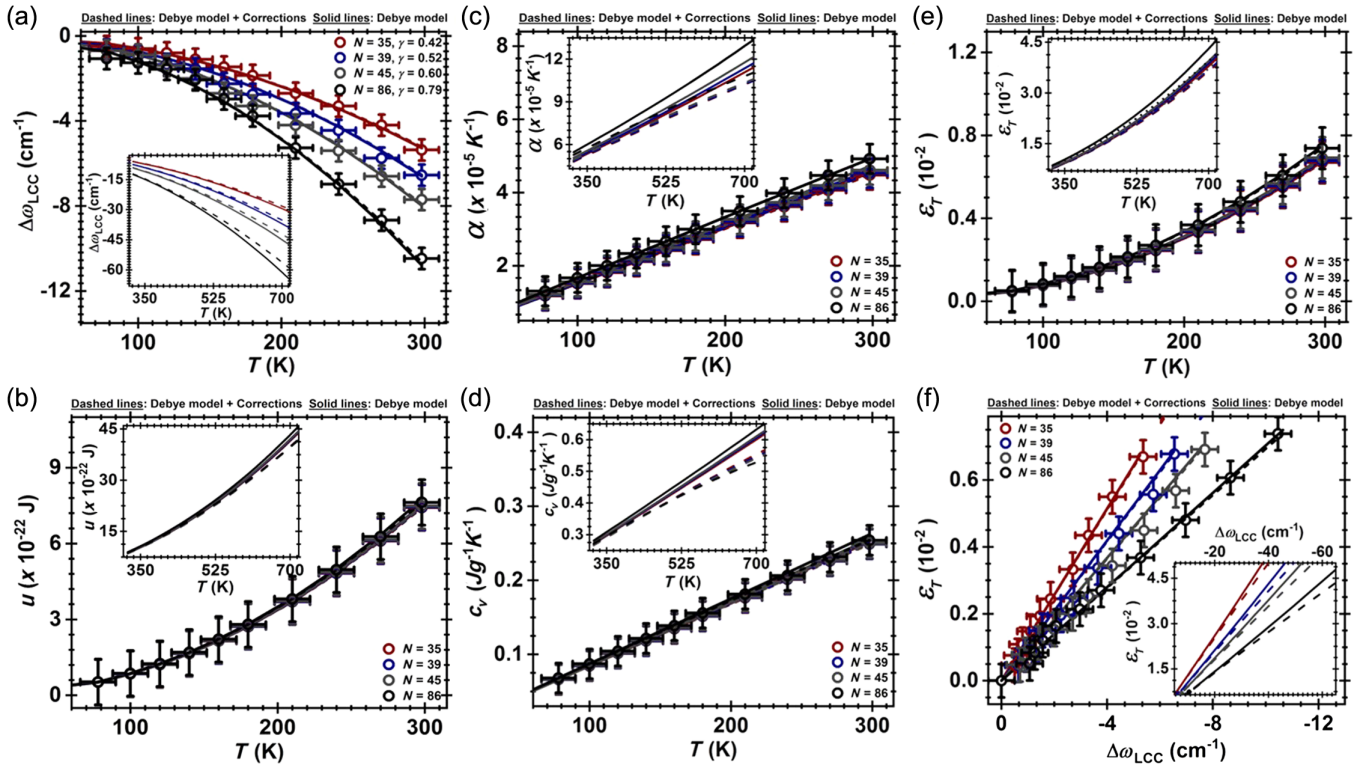


FIG. 1. Representative experimental results for LCCs ($N = 35, 39, 45$, and 86): (a) experimental $\Delta\omega_{\text{LCC}}(T)$ evolution with T . (b) The energy per N , $u(T)$, presents a quadratic, universal, and unified behavior with T ; (c) $\alpha(T)$ shows a linear universal behavior with T , where LCCs with N from 35 to 45 atoms present very similar $\alpha(T)$. For $N = 86$, however, $\alpha(T)$ increases at a slightly higher rate with increasing T . (d) The heat capacity per N , $c_v(T)$, presents a linear, universal, and unified behavior with T . (e) A T^2 universal dependence is observed for the thermal strain $\varepsilon_T(T)$ with N from 35 to 45 atoms presenting very similar $\varepsilon_T(T)$, while for $N = 86$ $\varepsilon_T(T)$ increases at a slightly higher rate with increasing T ; (f) Every LCC presents a distinct linear dependence of ε_T with $\Delta\omega_{\text{LCC}}(T)$. The curves show the predictions of the pure Debye model (solid lines) compared with the model proposed (dashed lines) and the insets in (a)–(f) suggest that for temperatures $300 \text{ K} \leq T \leq 700 \text{ K}$ the observables evolve more slowly with increasing T . Figures S3 and S4 in the Supplemental Material [56] show the results including all the 10 LCCs measured in this work.

$$\text{Ln } Z = -N \frac{T}{T_{\text{LCC}}} \int_0^{T_{\text{LCC}}/T} \text{Ln}[1 - \exp(-x)] dx, \quad (1)$$

where $x = \hbar\omega_{\text{ph}}/k_b T = T_{\text{ph}}/T$, ω_{ph} is the phonon frequency, and T_{ph} is the phonon equivalent temperature. Upon integration by parts, Eq. (1) becomes

$$\text{Ln } Z \approx N \frac{T}{T_{\text{LCC}}} \int_0^{T_{\text{LCC}}/T} \frac{x}{e^x - 1} dx. \quad (2)$$

For $T \ll T_{\text{LCC}}$, $T_{\text{LCC}}/T \rightarrow \infty$ and Eq. (2) is simplified to

$$\text{Ln } Z \approx N \frac{T}{T_{\text{LCC}}} \int_0^\infty \frac{x}{e^x - 1} dx = N \frac{T}{T_{\text{LCC}}} \left(\frac{\pi^2}{6} \right) = \frac{N k_b \pi^2}{6 \hbar} \frac{T}{\omega_{\text{LCC}}(T)}, \quad (3)$$

where $\int_0^\infty [x/(e^x - 1)] dx = \pi^2/6$. In our experiments, the temperature variation is not sufficiently large to take LCCs out of the harmonic oscillation regime but it is sufficient to renormalize ω_{LCC} leading to important T -dependent corrections with increasing T . Therefore, the LCCs' internal energy per N , $u(T)$, $c_v(T)$, and $\alpha(T)$ must contain one term representing the Debye approximation plus corrections involving derivatives of $\omega_{\text{LCC}}(T)$. This way $u(T)$ is

$$u(T) = \frac{k_b T^2}{N} \frac{d(\text{Ln } Z)}{dT} = \frac{k_b^2 \pi^2}{6 \hbar} \frac{T^2}{\omega_{\text{LCC}}(T)} \left[1 - \frac{T}{\omega_{\text{LCC}}(T)} \frac{d\omega_{\text{LCC}}}{dT} \right]. \quad (4)$$

Figure 1(b) shows that $u(T)$ is size independent. Equation (4) leads to the heat capacity per N - $c_v(T)$:

$$c_v(T) = \frac{du}{dT} = \frac{(k_b\pi)^2}{3\hbar} \frac{T}{\omega_{\text{LCC}}(T)} \left\{ 1 - 2 \frac{T}{\omega_{\text{LCC}}(T)} \frac{d\omega_{\text{LCC}}}{dT} + \frac{T^2}{\omega_{\text{LCC}}^2(T)} \left[\left(\frac{d\omega_{\text{LCC}}}{dT} \right)^2 - \frac{\omega_{\text{LCC}}}{2} \frac{d^2\omega_{\text{LCC}}}{dT^2} \right] \right\}, \quad (5)$$

with $\alpha(T)$ given by [56–59]

$$\alpha(T) = -\frac{\delta}{2\Theta^2} \frac{c_v(T)}{a_{c-c}} = \frac{42(k_b\pi)^2}{3\hbar m a_{c-c}^2 \omega_{\text{LCC}}^3(T)} \left\{ 1 - 2 \frac{T}{\omega_{\text{LCC}}(T)} \frac{d\omega_{\text{LCC}}}{dT} + \frac{T^2}{\omega_{\text{LCC}}^2(T)} \left[\left(\frac{d\omega_{\text{LCC}}}{dT} \right)^2 - \frac{\omega_{\text{LCC}}}{2} \frac{d^2\omega_{\text{LCC}}}{dT^2} \right] \right\}, \quad (6)$$

where $a_{c-c} = 1.37 \text{ \AA}$ is the average C-C distance. The parameters Θ and δ are obtained from the second and third derivatives of the series expansion:

$$U(R) \approx U(a_{c-c}) + \left(\frac{dU}{dR} \right)_{R=a_{c-c}} (R - a_{c-c}) + \frac{1}{2} \left(\frac{d^2U}{dR^2} \right)_{R=a_{c-c}} (R - a_{c-c})^2 + \frac{1}{6} \left(\frac{d^3U}{dR^3} \right)_{R=a_{c-c}} (R - a_{c-c})^3 + \dots,$$

where $U(R) = \varepsilon[(a_{c-c}/R)^{12} - 2(a_{c-c}/R)^6]$ is the Lennard-Jones potential (LJ) (R corresponds to the C-atom coordinate and ε is the potential depth). The LJ is chosen based on recent calculations by Wang and Lin [8], whose work describes LCCs' phonon dispersions quite accurately by modeling single and triple carbon bonds using such potential. Therefore, $(dU/dR)_{R=a_{c-c}} = 0$, $\frac{1}{2}(d^2U/dR^2)_{R=a_{c-c}} = \frac{1}{2}(72\varepsilon/a_{c-c}^2)$ and $\frac{1}{6}(d^3U/dR^3)_{R=a_{c-c}} = \frac{1}{6}(-1512\varepsilon/a_{c-c}^3)$. The second derivative is associated with the spring-mass harmonic potential. This means that $\Theta = \frac{1}{2}(d^2U/dR^2) = m[\omega_{\text{LCC}}(T)]^2/2$, $\delta = -21m[\omega_{\text{LCC}}(T)]^2/a_{c-c}$ and $\varepsilon = mR_m^2(\omega_{\text{LCC}}^0)^2/72$. With $\omega_{\text{LCC}}(T)$ and $d\omega_{\text{LCC}}/dT = -2(d^2\omega_{\text{LCC}}/dT^2)T$ we calculate $u(T)$, $c_v(T)$, $\alpha(T)$, and $\varepsilon(T)$ for each LCC.

The experimental $\alpha(T)$ [Fig. 1(c)] and $c_v(T)$ [Fig. 1(d)] endorse that LCCs are well described under DMF. Multiwalled CNTs [60], graphene [46], bilayer graphene [46], and graphite [61] (quasi-one-, two-, and three-dimensional sp^2 materials, respectively) present negative $\alpha(T)$ and ph-ph are fundamental to understand their $\Delta\omega(T)$ even for the temperatures in the range $0 \text{ K} \leq T \leq 300 \text{ K}$. Here, we see that for LCCs (sp materials) these dependencies are modified: $\alpha(T)$ is always positive and linearly dependent on T with a value of $5.0 \times 10^{-5} \text{ K}^{-1}$ ($4.5 \times 10^{-5} \text{ K}^{-1}$) at 300 K for the longest (shortest) chains, which is close to those reported for SWCNTs [$\alpha(T) = 2 \times 10^{-5} \text{ K}^{-1}$ at 300 K] [62] and $10\times$ higher in magnitude than the value reported for graphene [$\alpha(T) = -8.1 \times 10^{-6} \text{ K}^{-1}$ at 300 K] [46]. These values are in remarkably good agreement with the theoretical prediction of $7 \times 10^{-5} \text{ K}^{-1}$ by Wong *et al.* [44]. To the best of our knowledge, we present the first experimental evaluation of the LCCs' $c_v(T)$, which shows a unified linear dependence with T . The value 0.25 J/gK found at 300 K is close to those for graphite (0.7) and graphene (0.5 J/gK) [63,64]. We note that 0.25 J/gK is 4.6

times smaller than the value 1.16 J/gK predicted by Zhang *et al.* [4], who performed first-principles calculations for very short LCCs having a (5, 5) carbon nanotube as reference (non-negligible boundary conditions), what could explain such discrepancy.

The heat extracted from the LCCs leads to their shrinkage, which changes the LCCs' T -dependent internal pressure. This internal pressure is associated with both the system's entropy and thermal strain (ε_T). The thermal strain possesses a T^2 dependence [Fig. 1(e)] in excellent analogy with the P^2 dependence observed by Sharma *et al.* [22]. Therefore, in terms of behaviors, strains caused by either T or P differ only in magnitude. In Fig. 1(f), ε_T is plotted against $\Delta\omega_{\text{LCC}}(T)$. The graphic shows the linear behavior expected for small magnitude strains [22,57,58], which allow us to conclude that, for a given ε_T , larger LCCs experience larger $\Delta\omega_{\text{LCC}}(T)$, in further agreement with Sharma *et al.* [22].

For inefficient ph-ph, $\Delta\omega(T)$ should be fully described in terms of the LTE as

$$\Delta\omega_{\text{LTE}}(T) = \omega_{\text{LCC}}^0 (e^{-\gamma_P \int d\varepsilon} - 1), \quad (7)$$

where $d\varepsilon = \alpha(T)dT$ is the thermal strain between T and $T + dT$. Therefore, if our hypotheses are correct, by using the observables described in Eqs. (4)–(6), Eq. (7) should be equivalent to the obtained empirical relation $\Delta\omega_{\text{LCC}}(T) = -(d^2\omega_{\text{LCC}}/dT^2)T^2$ using γ_P as a fitting parameter. In the quasiharmonic regime there is no substantial distinction between the values of γ_P and γ_T (Grüneisen parameter at constant T), i.e., $\Delta\gamma \approx 0$ [65]. As shown in Fig. 01(a), Equation 07 fits the experimental $\Delta\omega_{\text{LCC}}(T)$ with γ_P ranging from 0.42 (shortest LCC) to 0.79 (longest LCC). These values agree with γ_T found in our previous work [22], which endorses the ph-ph inefficiency. Note that Sharma *et al.* [22] also showed that smaller LCCs present smaller γ_T in the quasiharmonic regime.

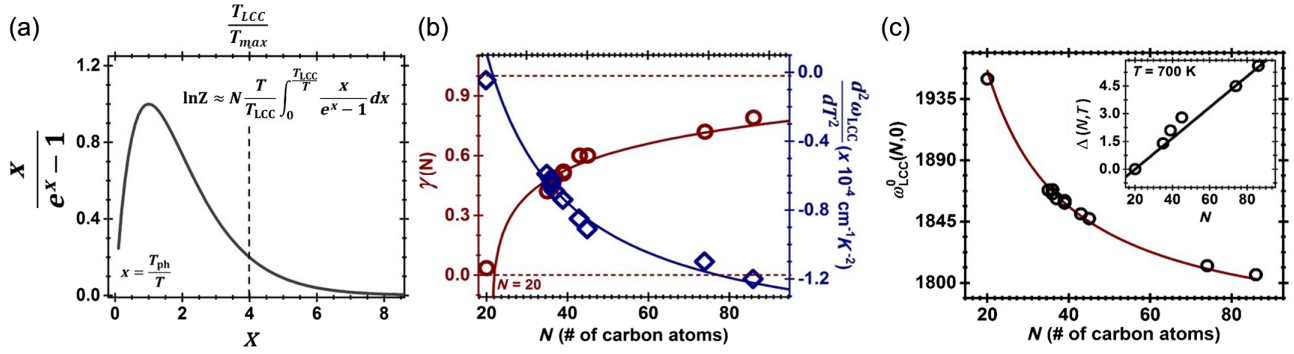


FIG. 2. (a) The distribution function $x/(e^x - 1)$ with $x = T_{ph}/T$ leads us to identify the high-temperature limit ($T_{\max}$) of the model. The distribution is most relevant for $0 \leq x \leq 4$, which implies that $T_{\max} = T_{LCC}/4 \approx 700$ K (see inset). (b) Universal dependences with N for both $\gamma_P(N) = \ln(N - 20)^{0.18}$ (red solid lines) and $d^2\omega_{LCC}/dT^2 = A + B/N$ (blue solid line with $A = -1.63 \times 10^{-4} \text{ cm}^{-1} \text{ K}^{-2}$ and $B = 35 \times 10^{-4} \text{ cm}^{-1} \text{ K}^{-2}$). The symbols (red circles and blue diamonds for γ_P and $d^2\omega_{LCC}/dT^2$, respectively) represent the experimental data. (c) $\omega_{LCC}^0(N, 0) = (1757 + 3980/N) \text{ cm}^{-1}$, where $\omega_{LCC}^0(N, 0)$ is the C-band frequency at $T = 0$ K. The open circles represent the experimental data. The inset shows a representative case at $T = 700$ K for $\Delta(N, T) = \Delta\omega_{LCC}^{\text{Corr}} - \Delta\omega_{LCC}^{\text{Debye}}$, which shows how the difference between $\Delta\omega_{LCC}^{\text{Corr}}$ proposed here [dashed lines in Fig. 1(a)] and $\Delta\omega_{LCC}^{\text{Debye}}$ [solid lines in Fig. 1(a)] progresses with N .

The ph-ph inefficiency, the successful description of LCCs' properties within DMF for $T \leq 300$ K, and the T -independent values of γ_P indicate that the corrections proposed are valid as long as the Debye temperature T_{LCC} is much larger than an upper limit ($T_{\max}$) after which the approximation $T_{LCC}/T \rightarrow \infty$ fails. Inspection of Eq. (2) shows that the LnZ behavior is determined by the integrand $x/(e^x - 1)$, which is most relevant for x between 0 and 4, as seen in Fig. 2(a). This way $T_{\max}$ is determined such that $T_{LCC}/T_{\max} = 4$, which means that $T_{\max} = T_{LCC}/4 \approx 700$ K. For $T \leq 300$ K the correction terms are negligible for every observable but for $300 \text{ K} < T < 700$ K (see insets in Fig. 1), the corrections become increasingly important. Our model predicts that $\Delta\omega_{LCC}(T)$ decreases at a slower rate with changing T [inset in Fig. 1(a)], while $\alpha(T)$ and $c_v(T)$ increase at a slower rate in comparison with the pure Debye model [insets in Figs. 1(c) and 1(d), respectively]. The magnitude of the corrections depends on $d^2\omega_{LCC}/dT^2$ and, consequently, on N [Fig. 2(b)]: the bigger the N the larger the correction. This is expected since the LCCs' thermal properties are connected to the intrinsic character of both single and triple bonds, which assume effective values reflecting the size of each LCC (see Sec. S3 in the Supplemental Material for additional details [56]).

As shown in Fig. 2(b), both γ_P and $d^2\omega_{LCC}/dT^2$ display universal dependences on N given by $\gamma_P(N) = \ln(N - 20)^{0.18}$ and $d^2\omega_{LCC}/dT^2 = A + B/N$ ($A = -1.63 \times 10^{-4} \text{ cm}^{-1} \text{ K}^{-2}$ and $B = 35 \times 10^{-4} \text{ cm}^{-1} \text{ K}^{-2}$), respectively. The $d^2\omega_{LCC}/dT^2$ relation with N lead us to the $\omega_{LCC}(N, T)$ universal relation

$$\omega_{LCC}(N, T) = \omega_{LCC}^0(N, 0) - \left(A + \frac{B}{N}\right) T^2, \quad (8)$$

where $\omega_{LCC}^0(N, 0) = (1757 + 3980/N) \text{ cm}^{-1}$ [17], see Fig. 2(c). The results for $\Delta\omega_{LCC}(T)$ also show that the LCCs' hosts do not influence the LCC's responses to T (Fig. S5 in the Supplemental Material [56]). The differences in the values observed for $\alpha(T)$, $c_v(T)$, and $\Delta\omega_{LCC}(T)$ comparing the solid (pure Debye model) and the dashed curves (pure Debye model + corrections) in Figs. 1(a)–1(d) allowed us to rewrite Eqs. (5)–(7) in terms of N and T . Such description is advantageous since it is now possible to predict the thermal properties of any chain at any $0 \text{ K} \leq T \leq 700$ K:

$$\alpha(N, T) = 985 \frac{T}{\omega_{LCC}^3(N, T)} - (3.22 \times 10^{-10}) \times \sqrt{(T - 300)^3 (N - 20)}, \quad (9)$$

$$c_v(N, T) = (1.6 \times 10^{-3}) \omega_{LCC}^2(N, T) \alpha(N, T), \quad (10)$$

where $\omega_{LCC}(N, T)$ is given by Eq. (8). Equations (9) and (10) allow for the determination of $\varepsilon_T(N, T) = \int \alpha(N, T) dT$ and $u(N, T) = \int c_v(N, T) dT$. Finally, $\Delta\omega_{LCC}(N, T) = -(d^2\omega_{LCC}/dT^2) T^2 + \Delta(N, T)$, which gives

$$\Delta\omega_{LCC}(N, T) = -\left(A + \frac{B}{N}\right) T^2 = (5.24 \times 10^{-7}) \times (T - 300)^2 (N - 20), \quad (11)$$

where $\Delta(N, T) = \Delta\omega_{LCC}^{\text{Corr}} - \Delta\omega_{LCC}^{\text{Debye}}$ is the difference between $\Delta\omega_{LCC}^{\text{Corr}}$ [dashed lines in Fig. 1(a)] and $\Delta\omega_{LCC}^{\text{Debye}}$ [solid lines in Fig. 1(a)]. The inset in Fig. 2(c) shows the representative case for $\Delta(N, 700)$.

In summary, the Letter focuses on the thermodynamic observables associated with LCCs and shows that due to

inefficient ph-ph, ω_{LCC} acts as a direct probe to the LCCs' thermal properties. Most materials are well described within DMF only at very low temperatures, but our results show that LCCs are well explained by DMF even at room temperature and likely for temperatures as high as 700 K. By observing the temperature dependence of $\Delta\omega_{\text{LCC}}(T)$, the thermodynamics associated with LCCs is successfully discussed, showing that fundamental relations such as internal energy, specific heat, coefficient of thermal expansion, and thermal strain present (N, T) -dependent universal behaviors. We present a simple semiempirical approach that is in excellent agreement with theoretical results in the literature and useful to other 1D systems.

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*These authors contributed equally to this work.

†Corresponding author.
barbosaneto@ufpa.br

‡Corresponding author.
paschoal@fisica.ufc.br

§Corresponding author.
ptaraujo@ua.edu

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