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Mechanical properties of carbynes investigated by \textit{ab initio} total-energy calculations

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As \textit{sp} carbon chains (carbynes) are relatively rigid molecular objects, can we exploit them as construction elements in nanomechanics? To answer this question, we investigate their remarkable mechanical properties by \textit{ab initio} total-energy simulations. In particular, we evaluate their linear response to small longitudinal and bending deformations and their failure limits for longitudinal compression and elongation.

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\section*{I. INTRODUCTION}

The rich chemistry of carbon is due to the capability of its electronic configuration to adjust to different bonding situations. Carbon atoms realize three main hybridization schemes of the valence orbitals: \textit{sp}, \textit{sp}\textsuperscript{2}, and \textit{sp}\textsuperscript{3} hybridization. \textit{sp}\textsuperscript{3} bonding tends to form three-dimensional networks, as in diamond and the related amorphous structures. \textit{sp}\textsuperscript{2}-hybridized orbitals lead to two-dimensional networks, at the basis of graphene, graphite, fullerenes, nanotubes, ribbon structures, and other forms of wide current interest for their unique mechanical and electronic properties. \textit{sp}-hybridized carbon forms linear structures (carbynes), which, compared to their chemically more stable \textit{sp}\textsuperscript{3} (alkanes) and \textit{sp}\textsuperscript{2} (alkenes) hydrogenated counterparts, tend to be rigid, thus potentially appealing as backbones for molecular nanotechnology.

Carbynes of increasing length and varied terminations are being synthesized\textsuperscript{4–6} and characterized.\textsuperscript{5} At the same time, the fabrication of carbon atomic chains from stretched nanotubes or graphene was achieved by controlled electron irradiation in transmission electron microscopes.\textsuperscript{6–8} The remarkable robustness of these carbon chains under irradiation combined with the ease of electron-beam fabrication at the nanometer scale can provide a route to the synthesis of actual nanodevices based on carbynes. Recent investigations have elucidated the prominent role of molecular oxygen as a primary source of chemical degradation of \textit{sp} carbon.\textsuperscript{9,10} In a pure-carbon environment carbynes are far more stable.\textsuperscript{11–15}

Effects of longitudinal straining\textsuperscript{16–19} and lateral bending\textsuperscript{20} were investigated very recently by \textit{ab initio} methods. The outcome of those studies is that carbyne chains are at one time extremely stiff against longitudinal straining and very soft against a bending deformation, to the extent that even extreme bending only moderately affects their bonding properties.\textsuperscript{20} In the present paper we focus on the mechanical properties of carbynes in the perspective of exploiting them as construction materials for nanoengineering. We use \textit{ab initio} simulations\textsuperscript{13,21} to predict the stiffness of the carbynes for both longitudinal strain and for bending, and their ultimate tensile strength. The main results of these simulations is that a \textit{sp} carbon chain (and it matters little whether it is of cumulene—all double bonds, or polyynic—with alternating single-triple bonds) exhibits not only an exceptional mechanical stiffness against longitudinal deformations, but also a small but nonzero rigidity against bending. Accordingly, a \textit{C}_n chain resembles a thin beam characterized by a finite rigidity against buckling, rather than a string. We find that, e.g., a single-carbyne \textit{C}_8 “pillar” can withstand a modest but non-negligible axial compression force \(\sim 0.7\) nN before buckling.

\section*{II. THE MODEL}

In simulations, one can tune the carbyne length by fixing the position of the end carbon atoms of a selected chain section, and leave all other atoms free to relax to their equilibrium position, compatible with the imposed strain. As well known to civil engineers, as soon as the compressive stress exceeds a critical value, the straight configuration of a beam subject to a purely axial strain \(\varepsilon<0\) turns unstable and “buckles” spontaneously to a curved arc. In the simulations of highly strained configurations, it is straightforward to address either the metastable straight configuration, or the stable buckled configuration investigated in Ref.\textsuperscript{20}.

To characterize the mechanical properties of a carbyne segment, and in particular to evaluate the buckling transition, one needs to identify the correct linear-response parameters in a simple model where the effects of different kinds of deformations can be decoupled and studied separately. The simplest continuum model describes a carbyne as a strained thin beam of a homogeneous material, thereby ignoring its discrete atomic structure. Three basic deformations can be applied to the beam: longitudinal traction or compression, lateral bending (as sketched in Fig. 1), and torsion around its axis. Correspondingly, the elastic deformation energy can be decomposed as

\begin{equation}
E_{el} = E_{\text{tens}} + E_{\text{bend}} + E_{\text{tors}},
\end{equation}

where \(E_{\text{tens}}\) is the tensile energy increase due to longitudinal elongation/shortening, \(E_{\text{bend}}\) is the energy due to the pure flexion of the beam, and \(E_{\text{tors}}\) is the energy increase due to the twisting of the beam. Remarkably, even this third term is important for the essentially one-dimensional cumulenes.\textsuperscript{13} In the present work we will, however, mostly ignore torsion, and concentrate on the first two terms.

In the linear-response (small-strain) regime, we write the tensile energy in terms of the strained length \(l\) relative to the fully relaxed length \(l_0\) as

\begin{equation}
E_{\text{tens}} = \frac{1}{2} \chi \frac{\lambda}{l_0} (l - l_0)^2,
\end{equation}

where \(\lambda\) is a strain.

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We identify the end-to-end distance \( l_0 \) from four lateral constraints, which induce two bending moments on the beam central section, which, as a result, bends as the arc of a circle. We compute the total adiabatic energies by means of the plane-wave DFT code QUANTUM ESPRESSO.\textsuperscript{25,39} We consider the two different limiting structures of carbynes: polyynes (or \( \alpha \) carbynes) with alternating single-triple bonds (\( \ldots \equiv C\equiv C\equiv C\equiv \ldots \), contrasted to cumulenes or \( \beta \) carbynes, with nearly-equal-length (double) bonds (\( \ldots =C=\equiv C=\ldots \). A typical polyynic chain is obtained when each end carbon atom forms a single bond with a ligand (e.g., a hydrogen atom, as in diacetylene), while a typical cumulene is obtained when each end carbon forms a double bond, for example to a \( \text{CH}_2 \) group, as in ethylene.

We focus on a \( C_8 \) chain segment, and compare its mechanical response in its \( \text{HC}_2C_8\text{C}_2H \) polyynic and \( \text{H}_2\text{C}-\text{C}_8\text{-CH}_2 \) cumulenic realizations. For the polyyne, we select hydrogenacetylide terminations rather than a single hydrogen because the \( \text{HC}_2 \) group allows us to impose a bending moment at the ends of the \( C_8 \) chain as in Fig. 1, by displacing a \( C \) atom along the chain continuation, rather than a chemically different \( \text{H} \) atom, thereby probing the intrinsic properties of carbyne, rather than those of a specifically terminated compound. We also consider \( C_n \) polyyne of different lengths, to check for size effects, and a \( 90^\circ \) end-twisted \( C_8 \) cumulene, which we must treat within the local spin-density approximation (LSDA) since twisting induces a peculiar total spin-1 electronic state, with a twofold-degenerate level occupied by two parallel-spin electrons.\textsuperscript{13} We do not attempt any serious size scaling, for two main reasons: (i) very long carbynes are mechanically unstable (thus unsuitable for nanomechanical applications) anyway, and (ii) the long-range interaction effects demonstrated recently\textsuperscript{17} would make a proper \textit{ab initio} size-scaling determination of the mechanical (in particular, bending) properties of long carbynes prohibitively expensive. In the present work we do not deal with the exotic odd-\( n \) \( C_n \) chains, which are chemically less stable and more difficult to synthesize.\textsuperscript{26,27}

To evaluate \( \chi \), starting from the fully relaxed carbyne we stretch or compress a straight \( C_8 \) chain section by fixing the positions of the end carbon atoms, and letting all other atoms free to relax to their resulting equilibrium positions under stress. We extract an estimate of \( \chi \) by fitting Eq. (2) to the resulting values of the excess energy as a function of the chain elongation, as reported in Fig. 2.

To evaluate the bending stiffness \( g \), we force the chain to bend by imposing a small lateral displacement of the two atoms adjacent to the \( C_8 \) section of the chain, while keeping...
FIG. 2. (Color online) The DFT-LDA tensile excess energy computed for the strained straight HC\textsubscript{2}-C\textsubscript{8}-C\textsubscript{2}H polyyne (bullets), as a function of $\varepsilon = (l - l_0)/l_0$. Solid line: a quadratic fit of Eq. (2), restricted to the seven data points within $\pm 0.8\%$ strain, where deviations from the linear-response regime are negligible. Crosses: the total excess energy without the constraint that the chain should remain straight; buckling starts to build up for strain exceeding $\varepsilon_\text{crit}$. The very similar data and fit for the cumulene H2C-C8-CH2 is not reported for clarity’s sake.

The first and last atom in the chain section bounded to remain laterally undisplaced, following the scheme of Figs. 1 and 3 (inset). We then let all other degrees of freedom (including the axial position of the laterally bounded atoms) of the C\textsubscript{8} section relax to form a circular arc, the axial position of the laterally bounded atoms) of the (inset). We then let all other degrees of freedom (including the axial position of the laterally bounded atoms) of the C\textsubscript{8} section relax to form a circular arc, the axial position of the laterally relaxed in-plane bent configuration (cumulene) (in-plane bent cumulenic chain: squares and dotted line; the out-of-plane bent cumulenic chain results in very similar energies and is not shown for clarity. Inset: an example of fully relaxed in-plane bent configuration of the cumulene H2C-C4-CH2. The resulting elasticity parameters $\chi$ and $g$ are collected in Table I. Despite the different bonding configuration of polyyne and cumulene, the overall longitudinal elasticity $\chi$ values are remarkably similar. The value of $\chi$ is compatible with a sound velocity $v_s = (\chi/\mu)^{1/2} \simeq 31.5$ km/s for long-wavelength longitudinal vibrations ($\mu$ is the linear mass density). In view of the long-range interactions demonstrated recently, the weak dependence of $\chi$ on the chain length is remarkable. The bending stiffness $g$ is comparatively more sensitive to the chain length (see also Fig. 4). If this velocity is used to evaluate the highest-frequency Brillouin-zone-boundary phonon in a simplest mass-spring chain, one

![Diagram](image)

FIG. 3. (Color online) The bending energy $E_\text{bend} = E_\text{mech} - E_\text{ext}$ [see Eq. (7)] as a function of the squared bending angle of -C\textsubscript{8}- chains. Here $E_\text{mech}$ is the excess energy in the bent configuration induced by the four-point fixed lateral displacement (at most 30 pm) sketched in Fig. 1, and $E_\text{ext}$ is the value obtained by Eq. (2), using $\chi$ as obtained in the straight-chain calculation of Fig. 2. Symbols are the DFT values, lines are linear fits, whose slopes, according to Eq. (4), represent the values of $g$. Polyynic chain: circles and dashed line. In-plane bent cumulenic chain: squares and dotted line; the out-of-plane bent cumulenic chain results in very similar energies and is not shown for clarity. Inset: an example of fully relaxed in-plane bent configuration of the cumulene H2C-C4-CH2.

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![Table](image)

**Table I.** The computed mechanical properties of selected carbynes. The elastic tension $\chi$ and bending stiffness $g$ define the elastic response, Eq. (7). The ultimate strain $\varepsilon_\text{ult}$ and tension $F_\text{ult}$ define the maximum stretching that the beam can stand before it breaks. For a C\textsubscript{8} chain section with both ends pinned (hinged, free to rotate), $F_\text{ult}$ measures the minimum compressive force with a corresponding strain $\varepsilon_\text{ult} = (l_\text{ult} - l_0)/l_0$ relative to the reported equilibrium length $l_0$ that must be applied to induce buckling.

<table>
<thead>
<tr>
<th></th>
<th>HC\textsubscript{2}-C\textsubscript{8}-C\textsubscript{2}H \hspace{5pt} (polyyne)</th>
<th>HC\textsubscript{2}-C\textsubscript{8}-C\textsubscript{2}H \hspace{5pt} (polyyne)</th>
<th>H\textsubscript{2}C-C\textsubscript{8}-CH\textsubscript{2} \hspace{5pt} (cumulene) \hspace{5pt} (planar)</th>
<th>H\textsubscript{2}C-C\textsubscript{8}-CH\textsubscript{2} \hspace{5pt} (cumulene) \hspace{5pt} (90° twisted)</th>
<th>H\textsubscript{2}C-C\textsubscript{12}-C\textsubscript{2}H \hspace{5pt} (polyyne)</th>
<th>H\textsubscript{2}C-C\textsubscript{16}-C\textsubscript{2}H \hspace{5pt} (polyyne)</th>
</tr>
</thead>
<tbody>
<tr>
<td>$l_0$ (C\textsubscript{8}) (pm)</td>
<td>377</td>
<td>886</td>
<td>888</td>
<td>889</td>
<td>1395</td>
<td>1903</td>
</tr>
<tr>
<td>$\varepsilon_\text{ult}$ (%)</td>
<td>16.2%</td>
<td>16.3%</td>
<td>17.9%</td>
<td>16.8%</td>
<td>16.4%</td>
<td>16.8%</td>
</tr>
<tr>
<td>$F_\text{ult}$ (nN)</td>
<td>12.7</td>
<td>12.3</td>
<td>14.7</td>
<td>12.7</td>
<td>12.1</td>
<td>12.1</td>
</tr>
<tr>
<td>$\chi$ (nN)</td>
<td>150</td>
<td>156</td>
<td>151</td>
<td>148</td>
<td>149</td>
<td>148</td>
</tr>
<tr>
<td>$g$ [nN (nm)$^{-2}$]</td>
<td>$7.91 \times 10^{-2}$</td>
<td>$5.69 \times 10^{-2}$</td>
<td>$5.68 \times 10^{-2}$</td>
<td>$5.95 \times 10^{-2}$</td>
<td>$5.62 \times 10^{-2}$</td>
<td>$5.11 \times 10^{-2}$</td>
</tr>
<tr>
<td>$\varepsilon_\text{crit}$ (%)</td>
<td>$-3.7%$</td>
<td>$-0.458%$</td>
<td>$-0.470%$</td>
<td>$-0.493%$</td>
<td>$-0.473%$</td>
<td>$-0.174%$</td>
</tr>
<tr>
<td>$F_\text{crit}$ (nN)</td>
<td>5.49</td>
<td>0.715</td>
<td>0.710</td>
<td>0.744</td>
<td>0.702</td>
<td>0.259</td>
</tr>
</tbody>
</table>
obtains a frequency $\nu_{\text{RHz}} = \nu_0/(\pi d_{\text{CC}}) \simeq 79$ THz, i.e., a wave number $\sim 2600$ cm$^{-1}$, expectedly slightly above the region of the observed high-frequency Raman and infrared modes emerging as the characteristic spectroscopic signatures of the carbynes.\textsuperscript{12,13,27–29}

The cumulene is anisotropic: one can make it bend either within the molecular plane, as sketched in the inset of Fig. 3, or perpendicularly to it, resulting in differently oriented double bonds\textsuperscript{28} being perturbed. We find that bending is significantly softer for the out-of-plane distortion than for the in-plane one, with an $\sim 5\%$ anisotropy.

The proposed cosine expression for the strain energy given in Eq. (9) of Ref. 20 is compatible with our Eq. (4) for small curvature (although a small amount of longitudinal strain energy is neglected there). The relation $g \simeq d_{\text{CC}} F_d$ obtained based on the average bond length $d_{\text{CC}} = 129$ pm and orientation-dependent bonding strength $F_d = 2.54$ eV as obtained in Ref. 20 yields $g = 5.26 \times 10^{-2}$ nN/(nm)$^2$, in good agreement with our result, especially with the $C_{12}$ chain value. This agreement is even more surprising in view of the extreme deformations introduced in Ref. 20, and indicates that the linear-response coefficient of Eq. (4) extends well outside its validity region, provided that the cosine expression Eq. (9) of that work is adopted.

According to elementary continuum mechanics,\textsuperscript{31} the critical axial load that a beam can sustain in its straight configuration before the onset of the buckling instability is given by Euler’s formula

$$F_{\text{crit}} = \pi^2 \frac{g}{I^2},$$

(8)

corresponding to a critical strain

$$\varepsilon_{\text{crit}} = \frac{F_{\text{crit}}}{F_0} \simeq -\frac{F_{\text{crit}}}{\chi} \simeq -\pi^2 \frac{g}{\chi l_0^2},$$

(9)

for a critical sinusoidal lateral deformation profile. By substituting the computed values for $\chi$ and $g$, we find that the $C_8$ carbyne sustains a critical strain of nearly $-0.5\%$, under the action of a critical force of approximately 0.7 nN, before buckling, as reported in Table I. The corresponding critical strain and force for cumulene (in the softer out-of-plane direction) are essentially equal. The computed values are relevant when both ends of $C_8$ are pinned (hinged, free to rotate), while if both ends were frozen, the critical force and strain would be four times larger. Likewise, following Eqs. (8) and (9), $F_{\text{crit}}$ decays rapidly with size. For example, for $C_{16}$, whose length is approximately 2.15 times that of $C_8$, $F_{\text{crit}} = 0.14$ nN, i.e., approximately 20% of the $C_8$ value only.

Direct calculation done for the $C_{12}$ polyyne in $HC_2C_{12}C_2H$ agrees with this scaling, with a small deviation due mostly to a smaller value of $g$, relative to the $C_8$ value (while $\chi$ is practically coinciding; see Table I).

IV. DISCUSSION AND CONCLUSION

We find it rather surprising that, despite the presence of in principle softer single bonds, the polyynic chain is essentially as hard to compress as double-bond-based cumulene. The same essentially equivalent stiffness of polyyne and cumulene is found against bending deformations, but here the result is less unexpected. Even a maximally twisted cumulene in a high-spin electronic state exhibits very similar mechanical properties (see Table I).

We check the obtained buckling critical point for possible effects of chain discreteness or anharmonicity, by running actual simulations. We compare pairs of DFT simulations at fixed strain: one bound to the linear configuration as in Fig. 2, and one starting off in a slightly curved geometry. In the curved simulations, below the buckling instability, the relaxing chain goes back to straight, and recovers the same energy and outward force that the chain produces on the pinned end atoms as in the straight calculation. In contrast, above the buckling instability, a curved shape stabilizes, with a net decrease in total energy and in the force acting on the pinned end atoms. Crosses in Fig. 2 report the fixed-end excess energy of the polyynic chain when allowed to relax in a buckled geometry. We find that the actual buckling instability occurs very close to the linear-response continuum-model value $\varepsilon_{\text{crit}}$ of Eq. (9). Relaxations near instability are rather delicate, since equilibrium is almost indifferent, providing a manifold of almost-equivalent geometrical configurations which give a hard time to the optimization algorithm.

The present work demonstrates that it is possible to go a long way in describing an immaterially thin object such as a monoatomic carbon chain with the mechanics of bulk construction elements, not unlike it was done for two-dimensional graphene in Refs. 32 and 33. Eventually, carbyne chains turn out rigid enough that the 886-pm-long freestanding $C_8$ polyyne can sustain a compressive strain of nearly 0.5% before buckling. On the tensile side, we extend our calculations well outside the linear-response region, to estimate the ultimate tensile strength of carbynes. The ultimate tension $F_{\text{ult}}$ is computed as the maximum force that the carbyne section produces in sustaining an externally imposed longitudinal strain $\varepsilon_{\text{ult}}>0$. The result in the 10 nN region agrees with previous determinations\textsuperscript{17,34} (but disagrees significantly with the Tersoff-Brenner-model determination of Ref. 35), and indicates that a single molecular chain could be used as a rope to lift a mass as heavy as one microgram without breaking.
If one could pack carbynes with a lateral density of one per \( A = 0.2 \times 0.2 \text{ nm}^2 \) cross-section, one would obtain a material characterized by a remarkable ultimate tensile strength of the order 300 GPa, comparable to that of carbon nanotubes.\(^{36}\)

Despite the remarkable mechanical properties of carbynes, their chemical reactivity makes them unsuitable to many applications in real world mechanical situations, where carbon fiber or carbon nanotubes provide superior stability with comparable mechanical properties. However in clean well-isolated nanoengineered devices one could envisage that the usage of C\(_n\) chains as structural or elastic elements may lead to consistent advantages over traditional solutions. For example, the soft bending degree of freedom could be exploited for the construction of sensitive accelerometers. Indeed, according to our evaluation of \( g \), a 0.1 \( \mu \text{g} \) mass hanging at the end of a 37-nm-long C\(_{30}\) chain (with the other end bonded to a fixed substrate) would deflect laterally by as much as 1 nm under an acceleration of 0.01 m s\(^{-2}\), i.e., one thousandth of the Earth’s gravitational field.

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