

One-dimensional pnictogen allotropes inside single-wall carbon nanotubes

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ABSTRACT

The discovery of phosphorene, a single layer of black phosphorus, has kick-started tremendous research efforts investigating the pnictogen nanomaterials. Arsenene and antimonene have now also been identified and these two-dimensional nanomaterials display physical properties superior to graphene for some applications. Recently, single-wall carbon nanotubes (SWCNTs) have been filled with P_4 molecules from the melt and As_4 molecules from the vapor phase. Inside the confinement of the SWCNTs, polymerization reactions were observed yielding new one-dimensional pnictogen allotropes. Here we show using high-resolution electron microscopy that such nanostructures can also be observed upon filling SWCNTs from the vapor phase using red phosphorus as the source material. Using larger diameter SWCNTs and filling from the vapor phase favors the formation of double-stranded phosphorus zig-zag ladders observed here for the first time. SWCNTs were generally found to fill well with liquid phosphorus. However, substantial decreases in the filling yields were observed for both phosphorus and arsenic filling of narrow SWCNTs using the vapor route. The filling experiments with molten antimony gave very low filling yields. However, the antimony zig-zag ladder could be observed on two occasions suggesting that this structural motif dominates across the pnictogens. Computational predictions of the encapsulation energies of the various pnictogen nanostructures are consistent with the observed experimental trends and band gap calculations predict that the single-stranded zig-zag chains of all investigated pnictogens are fully metallic. Using SWCNTs with diameters greater than 1.5 nm displayed a plethora of complex new phosphorus nanostructures which highlights an exciting new avenue for future work in this area.

INTRODUCTION

The pnictogen group of chemical elements derives its name from the Ancient Greek word for ‘choking’ which refers to the effect that the addition of nitrogen gas has on oxidation processes.¹ The physical and chemical properties of the pnictogens are remarkably diverse starting with the colorless dinitrogen gas across phosphorus, arsenic and antimony to the silvery and shiny metal bismuth.

The recent discovery of phosphorene, a single sheet of black phosphorus with a tunable band gap,²⁻⁵ has sparked a flurry of excitement with respect to the nanomaterials of these main group elements and a wide range of future applications, for example, in electronics, optoelectronics, sensing and thermoelectrics are currently anticipated.^{2-3, 6-10} The bulk allotropes of phosphorus include white phosphorus, which consists of the highly reactive tetrahedral P₄ molecules and exhibits three different polymorphs at ambient pressure,¹¹⁻¹² amorphous red phosphorus,¹³⁻¹⁴ fibrous phosphorus,¹⁵⁻¹⁶ phosphorus nanorods,¹⁷ as well as two layered structures, Hittorf’s violet phosphorus¹⁸⁻¹⁹ and black phosphorus,^{13, 20} the latter being the most stable allotrope at ambient pressure.²¹ Recently, the ‘blue’ allotrope has been grown on gold substrates using molecular beam epitaxial growth.²² Furthermore, a ‘green’ allotrope has been predicted computationally.²³

Gray arsenic (α -As) is the most stable allotrope at ambient conditions and consists of layers of puckered hexagons in a rhombohedral crystal structure.²⁴ The black allotrope (β -As) is metastable with an orthorhombic layered crystal structure similar to that of black phosphorus.²⁵ Amorphous arsenic, with a wide range of different densities, has also been identified.^{24, 26} Analogous to white phosphorus, yellow arsenic (γ -As) consists of tetrahedral As₄ molecules.²⁷ However, due to its extreme sensitivity towards light and X-rays, which trigger the conversion to gray arsenic, no crystallographic data is available for γ -As.²⁸ In addition to the bulk allotropes,

gas-phase clusters in size up to As_5 have been identified experimentally.²⁹ Arsenene, the arsenic equivalent of phosphorene, displays a large bandgap³⁰⁻³¹ which, combined with a low thermal conductivity, makes it a highly desirable material for thermoelectric applications.³² Other electronic, optoelectronic, spintronic and sensing applications are also envisaged.¹⁰ Experimentally, arsenene can be prepared using plasma-assisted deposition,³³ aqueous shear exfoliation³⁴ and ultrasonication of gray arsenic in appropriate solvents.³⁵⁻³⁶

The stable gray allotrope of antimony is isostructural with gray arsenic and this pnictogen also displays the analogous metastable black allotrope.¹⁰ By contrast, tetrahedral Sb_4 molecules appear to only be stable in the gas phase.³⁷⁻³⁸ However, an amorphous and highly metastable ‘explosive’ antimony allotrope exists.³⁹⁻⁴⁰ Antimonene has recently been added to the family of pnictogen nanomaterials using mechanical exfoliation,⁴¹ liquid-phase exfoliation⁴² and epitaxial growth.⁴³⁻⁴⁴

Carbon nanotubes have been used as inert molds to prepare new one-dimensional allotropes of a range of elements including carbon in its various forms,⁴⁵⁻⁴⁷ iodine,⁴⁸ sulphur,⁴⁹ selenium⁵⁰, tellurium,⁵¹ as well as the two pnictogens phosphorus⁵²⁻⁵³ and arsenic.⁵⁴ In the case of phosphorus, fibrous chains with alternating P_8 and P_2 units have been identified inside large-diameter multi-wall carbon nanotubes.⁵² Single-stranded phosphorus zig-zag chains have been prepared in narrow single-wall carbon nanotubes (SWCNTs) by polymerizing encapsulated chains of P_4 molecules.⁵³ The formation of the zig-zag chain can be regarded as the first step from white to red phosphorus, and consequently, this one-dimensional allotrope of phosphorus has been named ‘pink’ phosphorus.⁵⁵ The single-stranded zig-zag chain has also recently been observed for arsenic inside SWCNTs which also forms double-stranded zig-zag ladders.⁵⁴ In addition to stabilizing new pnictogen nanostructures inside carbon nanotubes, they also provide environments stabilizing the highly reactive P_4 and As_4 molecules against the infamous spontaneous combustion with

atmospheric oxygen to give the corresponding oxides.⁵³⁻⁵⁴ Filling carbon nanotubes with antimony has been attempted as part of the arc-discharge preparation process.⁵⁶ However, the atomic structure of the filling material could not be resolved.

Here we investigate and explore a range of questions related to the formation of one-dimensional pnictogens inside SWCNTs. These include (i) the filling with phosphorus from the vapor phase using red phosphorus as the source material, (ii) the search for the double-stranded zig-zag chain of phosphorus, which has previously been observed for arsenic,⁵⁴ (iii) the dependence of filling yields and structures of the pnictogens on the diameters of the SWCNTs, (iv) the filling of SWCNTs with antimony from the melt as well as (v) a complete set of inclusion and band-gap energies of the encapsulated pnictogens using density functional theory (DFT).

EXPERIMENTAL SECTION

Purification and opening of SWCNTs

Tuball SWCNTs were purchased from OCSiAl in raw powder form and purified with steam at 900°C for 5 hours.⁵⁷ A second heat treatment in air at 450°C for 30 minutes ensured the opening of the SWCNT tips. After this, the SWCNTs were refluxed in 6 M HCl for 2 hours and washed on a filter membrane with an excess of water. Finally, the sample was dried in an oven overnight at 60°C. HiPCO SWCNTs were purchased from Nanointegris Inc. in ethanol-wet cake form. These were dried at 100°C and 10^{-2} mbar for 4 days, and then tip-opened by heating to 400°C in air. CoMoCAT SWCNTs were purchased from Sigma Aldrich in powder form and tip-opened by heating to 400°C in air.

Filling experiments with molten white phosphorus⁵³

Caution: White phosphorus is volatile, extremely toxic and, especially after sublimation, pyrophoric. Extreme care should be taken when conducting experiments of this type. 50 mg of the SWCNTs were heated to 200°C at 10^{-6} mbar and transferred into a glove box under vacuum. Crude white phosphorus was purified by vacuum sublimation onto a liquid nitrogen-cooled probe in a Schlenk flask attached to a dual manifold Schlenk line. In a glove box, the sublimation probe was transferred immediately to a second Schlenk flask loaded with the SWCNTs and a Teflon-coated magnetic stirrer. Back on the Schlenk line, warm water was poured into the sublimation probe to melt the white phosphorus onto the SWCNTs. The black mixture was stirred at 50°C under nitrogen for one hour. After this, the sublimation probe water was poured out of the probe, and after rinsing with a little acetone, was replaced with liquid nitrogen to sublime off the excess white phosphorus from the SWCNTs. The probe was removed in the glove box and set aside for storage. The Schlenk flask was returned to the Schlenk line and the filled SWCNTs were suspended in 125 mL deionized water and sonicated for 5 minutes to help disperse the mixture. Compressed air was then gently bubbled through the stirred suspension for two days using a needle to oxidize any excess white phosphorus. The sample was filtered in air on a Hirsch funnel using glass-fiber filter paper, washed with deionized water and dried in a desiccator. Caution: if the air oxidation step is shortened or omitted, the nanotubes can be pyrophoric and ignite spontaneously on the filter.

Filling experiments with phosphorus from the vapor phase

20 mg SWCNTs and 20 mg red phosphorus were ground in an agate mortar and pestle for 5 minutes to create a homogenous mixture. The mixture was then transferred to a 200 mm long quartz ampoule which was pumped down to 10^{-6} mbar, heated to 200°C at $7^{\circ}\text{C min}^{-1}$ and held for 2 hours. The ampoule was then sealed under dynamic vacuum, heated in a tube furnace to 500°C at $10^{\circ}\text{C min}^{-1}$ and held for 12 hours with the sample end in the center of the furnace. After cooling,

the ampoule was rotated by 180° and the sample reheated as before to sublime the white phosphorus along the length of the ampoule. This step was done three times. The fourth and last heat treatment sublimed the white phosphorus onto the SWCNTs located at the cool end of the ampoule. After recovering the sample from the ampoule in a glove box, the sample was transferred to a Schlenk flask as in Section 2.2, suspended in water and the excess white phosphorus removed by bubbling air through the suspension. The nanotubes were recovered by filtration onto glass-fiber paper on a Hirsch funnel.

Filling experiments with arsenic from the vapor phase⁵⁴

20 mg SWCNTs and 20 mg gray arsenic were ground for 5 minutes using an agate pestle and mortar to create a homogenous mixture. The mixture was then placed in a 200 mm long quartz ampoule which was pumped down to 10^{-6} mbar, heated to 200°C at 7°C min⁻¹ and held for 2 hours. The ampoule was sealed under dynamic vacuum, heated in a tube furnace to 615°C at 10°C min⁻¹ and held for 12 hours with the sample end in the center of the furnace. The ampoule was reheated three more times, each time rotating the ampoule by 180° to sublime the arsenic along the ampoule. The fourth and last heat treatment sublimed the arsenic onto the SWCNTs located at the cool end of the ampoule. After recovering the sample from the ampoule, excess arsenic was removed by stirring in 100 mL 0.5 M nitric acid solution for 4 hours followed by filtration and washing with water.

Filling experiments with molten antimony

30 mg SWCNTs and 30 mg gray antimony were ground for 5 minutes in an agate pestle and mortar to create a homogenous mixture. The mixture was then placed in a 200 mm long quartz ampoule which was pumped down to 10^{-6} mbar, heated to 200°C at 7°C min⁻¹ and held for 2 hours. The

ampoule was sealed under dynamic vacuum, heated in a muffle furnace to 700°C at 10°C min⁻¹ and held for 66.5 hours. After recovering the sample from the ampoule, excess antimony was removed by stirring in 150 mL 1 M nitric acid solution for 4 hours followed by filtration and washing with water.

High-resolution transmission electron microscopy (HRTEM)

The filled SWCNTs were prepared for HRTEM by sonication in absolute ethanol for 1 minute and then dropping the dispersions onto lacey carbon Agar 300 mesh copper grids. HRTEM was conducted using a FEI Titan 80/300 STEM/TEM instrument fitted with a Cs (image) corrector. The microscope was operated at 80 kV to reduce electron beam damage to the samples. The image corrector was necessary to image the structure of the SWCNTs by reducing the effects of spherical aberrations. All images were taken with the image corrector corrected to at least 18 mrad. Simulated TEM images were obtained using the SimulaTEM software⁵⁸ using an accelerating voltage of 80 kV, a defocus value of -310.271850 Å and a spherical aberration constant of 0.15 mm.

X-ray photoelectron spectroscopy (XPS)

X-ray photoelectron spectroscopy of the samples on indium foil was carried out using a Thermo Scientific K-Alpha instrument with a monochromated Al K α X-ray source and flood gun. Survey scans were obtained by accumulating three spectra with a resolution of 1 eV and pass energy of 200 eV. High-resolution scans in the P2p, As3d and Sb3d regions were obtained from the average of 10 spectra recorded with a resolution of 0.1 eV and pass energy of 50 eV. A 400 μ m spot size and 50 ms dwell time was used for each scan.

Computational methods

DFT calculations were performed using the Vienna ab initio Simulation Package (VASP) where the core electrons were described with projector augmented wave (PAW) potentials.⁵⁹⁻⁶⁰ An energy cutoff of 400 eV was used to expand the wave functions. The optB86b-vdW van der Waals inclusive exchange correlation functional was used.⁶¹ An empirical cylindrical confining potential was implemented to simulate the confinement of carbon nanotube following the procedure used in refs 53, 62. The empirical potential has a Morse-type formula defined as $V(x) = D [(1 - \exp(-A(x - R)))^2 - 1]$ in eV where x is the distance of a pnictogen atom to the carbon nanotube wall in Å. The various parameters are: $D = 0.125, 0.172, 0.203, A = 1.060, 1.052, 0.974$ and $R = 3.640, 3.565, 3.733$, respectively for P, As, and Sb. The parameters used were fitted to the DFT interaction between an X_4 molecule ($X = P, As$ or Sb) and graphene.

RESULTS AND DISCUSSION

Filling experiments with elemental pnictogens

To investigate the various aspects of filling SWNCTs with elemental pnictogens mentioned earlier, SWCNTs with a range of different diameters were used including CoMoCat SWCNTs with diameters smaller than 1 nm,⁶³ HiPCO SWCNTs, which are in the 0.8 – 1.3 nm diameter range,⁶⁴ and Tuball SWCNTs with diameters exceeding 1.5 nm.⁶⁵

The filling yields of the pnictogens were determined using XPS as shown in Figure 1. In all cases, peaks associated with both elemental and oxidized pnictogens were observed. The complete set of XPS spectra are provided in Figures S1 and S2. The elemental pnictogens are presumed to be protected inside the SWNCTs, while the oxidized material is the result of some residual elemental pnictogens adhering to the outside walls or the tips of the tubes, and able to react with atmospheric oxygen or with the nitric acid used for purification.

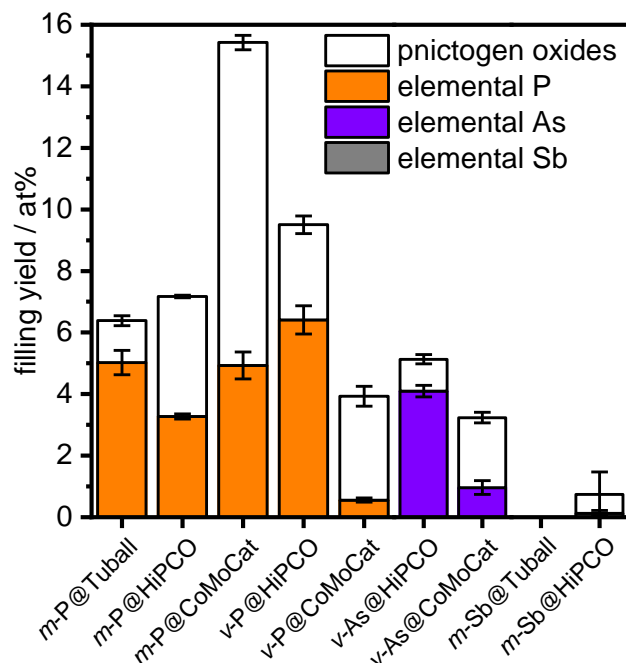


Figure 1. Filling yields of elemental and oxidized pnictogens in various types of SWCNTs. The prefix ‘m’ indicates filling from the melt whereas ‘v’ denotes vapor phase filling. The filling yields were determined by fitting the survey and high-resolution pnictogen spectra shown in Figure S1 and S2. The error bars were calculated by measuring at least three XPS spectra for a given sample. The filling yields of m-P@CoMoCat and v-As@HiPCO were reported in our earlier studies.⁵³⁻⁵⁴

Some of the highest filling yields were observed for elemental phosphorus using the melt filling technique and they were found to be relatively independent of the diameter of the SWCNTs. Filling with phosphorus from the vapor phase was successful as well and gave the actual highest filling yield out of all experiments in the case of HiPCO tubes. The filling yield using the smaller diameter CoMoCat tubes was considerably lower. These findings suggest that if P₄ molecules fit inside the SWCNTs, then filling from the melt is generally viable. However, the vapor filling does seem to

work better for larger-diameter SWCNTs suggesting some limitations for the vapor condensation of P₄ molecules into small-diameter SWCNTs.

As previously reported, HiPCO SWCNTs also fill well with elemental arsenic⁵⁴ from the vapor phase delivering comparable filling yields as obtained for phosphorus. Similar to the v-P@CoMoCat experiment, CoMoCAT SWCNTs also gave a quite low filling yield for elemental arsenic. Because As₄ is notoriously thermally unstable,⁶⁶ the filling of SWCNTs with elemental arsenic from the liquid phase appears to be impossible.

Filling experiments with molten antimony were not very successful giving a filling yield of elemental antimony of only 0.13 at% for m-Sb@HiPCO. Unlike molten white phosphorus, liquid antimony does not seem to contain any molecular species.⁶⁷ So the poor filling yield is likely to be due to the high surface tension of molten antimony of 368 mN m⁻¹⁶⁸ which exceeds the often quoted limit for filling SWCNTs of 200 mN m⁻¹.⁶⁹ Even if Sb₄ molecules formed upon entering the SWCNTs, they would be expected to be highly unstable. For example, attempts at isolating yellow antimony on a graphite surface resulted in the grow of a complex selection of extended nanostructures around defect sites.⁷⁰

Microstructures of the one-dimensional pnictogen allotropes

Three different types of phosphorus and arsenic nanostructures have been identified inside the SWCNTs with HRTEM as shown in Figure 2. These include lines of tetrahedral P₄ or As₄ molecules analogous to peas in a pod, the double-stranded zig-zag ladder and the single-stranded zig-zag chain. Notably, the phosphorus double-stranded zig-zag ladder is reported here for the first time whereas the other structures have been observed in our previous studies.⁵³⁻⁵⁴ It is also worth mentioning that the lines of P₄ molecules and the phosphorus zig-zag chain were previously

observed inside CoMoCat SWCNTs filled from the melt⁵³ whereas they are imaged here for the first time inside the larger diameter HiPCO SWCNTs after filling from the vapor phase.

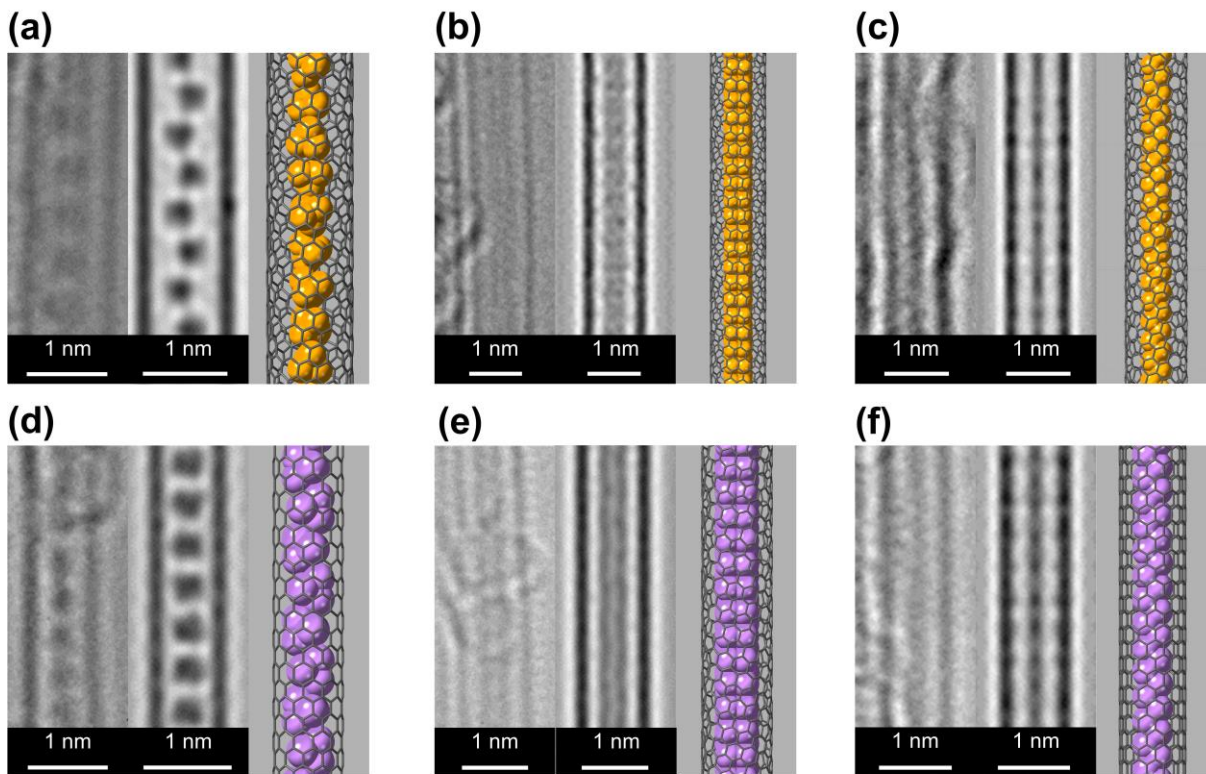


Figure 2. High-resolution transmission electron microscopy (HRTEM) images of the three different nanostructures adopted by elemental phosphorus and arsenic inside SWCNTs. Each panel shows three images including the experimental (left) and simulated micrographs (middle) as well as the corresponding DFT geometry-optimized atomistic models (right). The data for $v\text{-P@HiPCO}$ and $v\text{-As@HiPCO}$ are shown in (a-c) and (d-f), respectively. The imaged nanostructures include (a,d) the chain of P_4/As_4 beads, (b,e) the double-stranded zig-zag ladder and (c,f) the single-stranded zig-zag chain. The simulated micrographs were obtained by rotationally averaging the filling materials (about the center of mass for the molecular species and about the tube axis for the chain structures).

As can be seen in Figure 2, the experimental HRTEM images agree well with calculated images based on structures that were geometry-optimized with DFT. The best agreement between the experimental HRTEM images and the simulated pictures was obtained by rotationally averaging the P_4 and As_4 molecules about their centers of mass, and the linear chains by rotation about the axis of the SWCNT. Similar dynamic rotations of the filling materials have been previously observed in the case of encapsulated graphene nanoribbons⁴⁷ and may most likely be due to the exposure to the electron beam inside the HRTEM.

The two polymerized pnictogen chain structures can be clearly distinguished on the basis of their widths. The zig-zag ladder manifests as a broad feature in HRTEM with a gap between the two parallel strands whereas the single zig-zag chain is seen in the form of a single, narrow line. The distance between the two strands of the zig-zag chains is on average 2.38 Å for arsenic and 2.43 Å for phosphorus. As shown in Figure S3, the distance between the two strands increases slightly as the diameter of the confining SWCNT increases. This effect is seen both in the experimental data as well as for the DFT-optimized structures.

The effects of increasing the SWCNT diameter and of changing from melt to vapor-phase filling on the relative occurrences of the various nanostructures was investigated next with HRTEM. In this context, it is worth mentioning that the CoMoCat and HiPCO SWCNTs used in this study displayed the expected diameter distributions in HRTEM with CoMoCat SWCNTs mainly in the 0.6 – 0.9 nm and the HiPCO SWCNTs in the 0.9 – 1.1 nm diameter range.⁶³⁻⁶⁴ As shown in Figure 3(a), the narrow CoMoCat SWCNTs filled with phosphorus from the melt showed lines of P_4 molecules and zig-zag chains predominately at lower diameters and, as expected due to their larger size, zig-zag ladders at larger diameters. Upon increasing the diameters to HiPCO SWCNTs, while still filling from the melt, the relative occurrence of the various phosphorus

nanostructures changes quite dramatically (Figure 3(b)). Due to the increased availability of larger diameter SWCNTs, the zig-zag ladder is now clearly the dominant species. Interestingly, this effect becomes even more pronounced for the vapor-filled P@HiPCO sample where the zig-zag ladder is found almost exclusively (Figure 3(c)). Compared to the filling from the melt, vapor-filling using red phosphorus as the source material is carried out at much higher temperatures. The dominance of the zig-zag ladder in the v-P@HiPCO sample therefore suggests that increasing the temperature favors the formation of the zig-zag ladder inside the SWCNTs. As shown in Figure 3(d) and previously reported,⁵⁴ the zig-zag ladder also dominates in the case of v-As@HiPCO. However, the line of beads and the zig-zag chain are observed somewhat more frequently than for v-P@HiPCO.

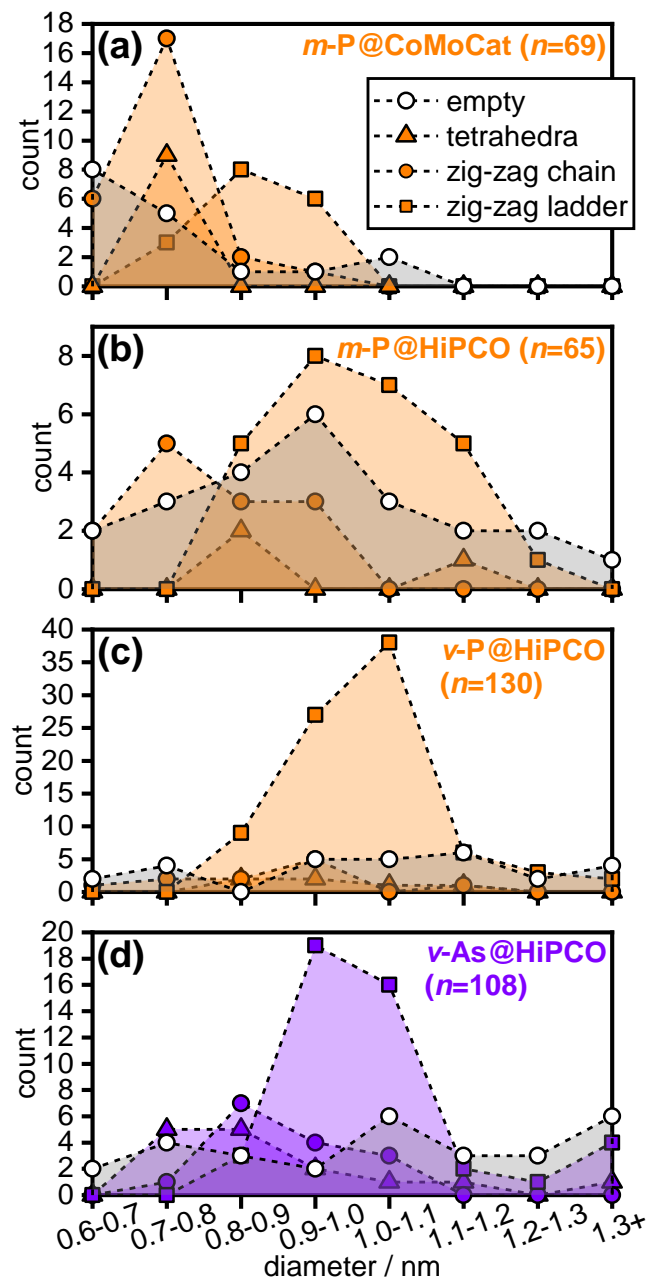


Figure 3. Occurrences of the different pnictogen nanostructures as a function of the SWCNT diameter for (a) *m-P@CoMoCat*, (b) *m-P@HiPCO*, (c) *v-P@HiPCO* and (d) *v-As@HiPCO*. Panel (d) is reproduced from ref. 54. The total number of HRTEM observations is noted for each panel.

In addition to the CoMoCat and HiPCO SWCNTs, large-diameter phosphorus-filled Tuball SWCNTs were also investigated with HRTEM (Figure 4). Interestingly, increasing the diameter of the SWCNTs above 1.5 nm leads to a complete loss of the previously observed nanostructures. Instead, several irregularly-shaped filling materials were observed ranging from what could be onion-shaped nanostructures in Figure 4(a) to potentially two zig-zag chains or ladders arranged in a double-stranded fashion in Figure 4(c). However, we emphasize that we currently do not feel confident in making definitive structural assignments. Clearly, much more work is needed to resolve these structures including further computational studies.

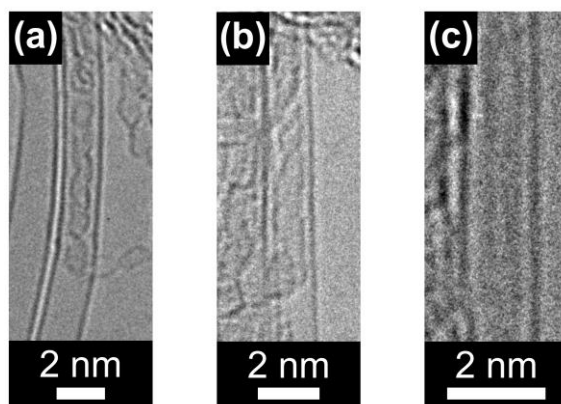


Figure 4. HRTEM images of *m-P@Tuball* showing encapsulated phosphorus nanostructures with irregular shapes.

Despite the low filling yield for *m-Sb@HiPCO* indicated by the XPS analysis shown in Figure 1, HRTEM images were recorded for this sample. On two occasions, double-stranded antimony nanostructures were observed inside the SWCNTs as shown in Figure 5. In addition to the frequent observation of phosphorus and arsenic zig-zag ladders, this structural motif therefore also seems to be realized for antimony. Extensive searching for single Sb_4 molecules or the zig-zag chain was unsuccessful.

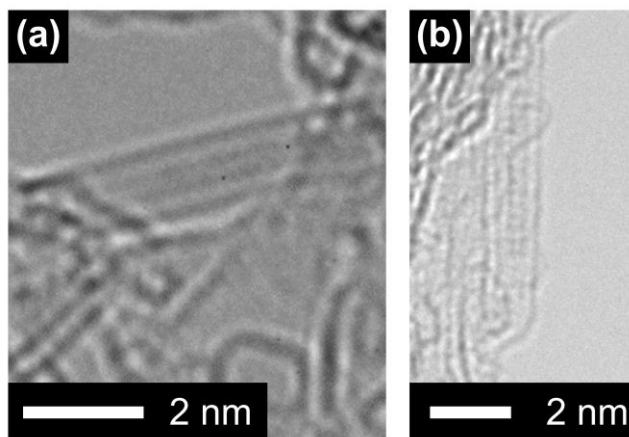


Figure 5. HRTEM images of *m-Sb@HiPCO* showing double-stranded structures in (a) 1.086 nm and (b) 0.963 nm diameter SWCNTs.

Computational energies and bandgaps of the one-dimensional pnictogen allotropes

Figure 6 shows the DFT energies of the three experimentally observed one-dimensional pnictogens. In line with the experimental filling yields shown in Figure 1, the phosphorus structures are energetically most stable whereas antimony displays the highest energies. The lowest-energy antimony structures are zig-zag ladders at large diameters which agrees well with the observations in Figure 5. In the cases of phosphorus and arsenic, the molecular forms and the single-stranded zig-zag chain are most stable at lower diameters whereas the double-stranded zig-zag ladder becomes increasingly more stable as the confinement diameter increases. All these trends are therefore in very good agreement with the experimental observations reported in Figure 3.

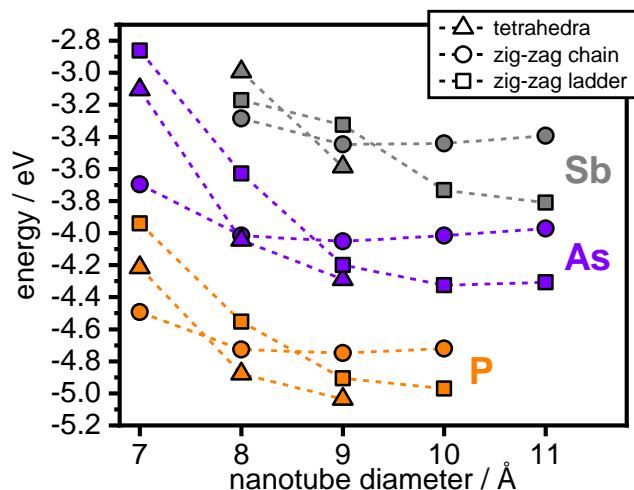


Figure 6. DFT energies of the three one-dimensional pnictogen nanostructures as a function of the nanotube diameter and defined as the per-atom binding energy relative to the isolated pnictogen atom plus the confining energy. Missing data points for a given diameter are due to the instabilities of the structures.

Figure 7 shows the DFT-calculated electronic densities of state of the three experimentally observed pnictogen nanostructures. The lines of P_4 molecules display the largest band gaps; these decrease down the pnictogen group for the various molecular species. Further closing of the band gaps to quite similar values for the various pnictogens is observed for the zig-zag ladders. Finally, the single-stranded zig-zag chains of all pnictogens are fully metallic with density of states spanning across the Fermi level. It is interesting to note that the electronic densities of state of the various nanostructures display similar features for the various pnictogens. However, depending on the pnictogen in question, the features are shifted or stretched; typically away from the Fermi level for phosphorus and towards the Fermi level in the case of antimony illustrating its general tendency towards metallic structures.

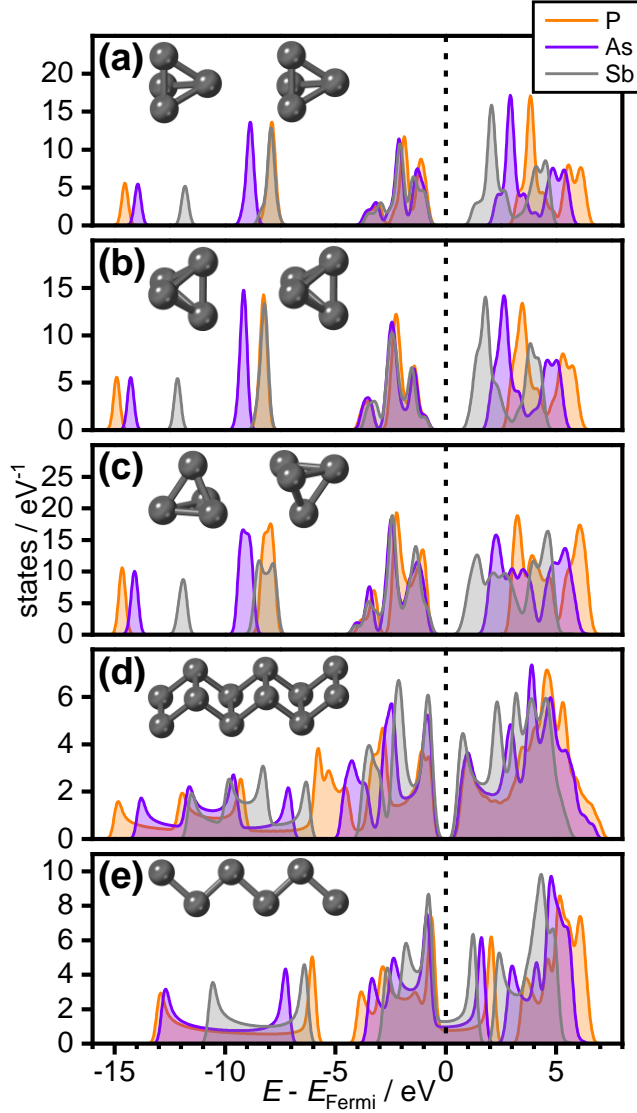


Figure 7. Electronic densities of state of confined pnictogen nanostructures including (a) corner-to-face, (b) edge-to-edge and (c) face-to-face tetrahedra as well as (d) zig-zag ladders and (e) zig-zag chains. The Fermi level is indicated by dashed vertical lines. The data for arsenic was taken from ref. 54.

CONCLUSIONS

In summary, we have gained new insights into the filling of SWCNTs with elemental phosphorus, arsenic and antimony as well as the resulting one-dimensional pnictogen allotropes. The double-

stranded phosphorus zig-zag ladder was observed for the first time and its relative occurrence upon filling from the melt was found to depend on the diameter of the SWCNTs. Filling SWCNTs with elemental phosphorus from the vapor phase using red phosphorus as the source was shown to be possible and yielded predominately the new zig-zag ladders. While the filling yield from molten phosphorus did not show a dependence on the diameter of the SWCNTs, for both phosphorus and arsenic filling from the vapor phase it was found that the larger diameter HiPCO SWCNTs fill much better than the narrower CoMoCat SWCNTs. The antimony filling yields were generally very low. However, two double-stranded structures could be imaged in HRTEM suggesting that this structural motif is a dominating one across all three pnictogens. Filling SWCNTs with diameters greater than 1.5 nm with elemental phosphorus showed a range of complex nanostructures inside the SWCNTs.

In general, the DFT-calculated energies of the various one-dimensional pnictogens were in very good agreement with the experimental findings and helped rationalizing the decreasing filling yields down the pnictogen group as well as the relative occurrences of the various nanostructures as a function of the SWCNT diameter. Furthermore, DFT calculations showed that the band gap of the confined pnictogens closes as the degree of polymerization increases and the zig-zag chains of all three pnictogens are predicted to be fully metallic. The changes in the band gaps are most pronounced for phosphorus and much weaker for antimony reflecting the general tendency of this pnictogen towards metallic behavior.

Overall, the changeovers in electronic behavior is a very interesting aspect of the one-dimensional pnictogen allotropes. To track these experimentally, future efforts will aim at preparing the new pnictogen allotropes inside insulating confinements such as boron nitride nanotubes. Furthermore, it has been pointed out that the confined phosphorus and arsenic materials

could be used for making P₄ and As₄ available for chemical reactions.^{53-54, 66, 71-73} Whether longer chain segments can be recovered intact from inside the SWCNTs using appropriate solvents is another open question. Finally, as we have shown here, moving towards larger diameter SWCNTs opens a door to ever more complex pnictogen materials. Exploring their structures and physical properties in detail will be an exciting future challenge.

ASSOCIATED CONTENT

Supporting Information. The supporting information contains the XPS spectra used for the analysis for preparing Figure 1 in the main article as well as an analysis of the distance between the two strands of the zig-zag ladder in both the experimental HRTEM images as well as the DFT-optimized structures.

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