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Mechanisms of Postsynthesis Doping of Boron Nitride Nanostructures with Carbon from First-Principles Simulations

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Electron-beam-mediated postsynthesis doping of boron-nitride nanostructures with carbon atoms [Nature (London) 464, 571 (2010); J. Am. Chem. Soc. 132, 13692 (2010)] was recently demonstrated, thus opening a new way to control the electronic properties of these systems. Using density-functional theory static and dynamic calculations, we show that the substitution process is governed not only by the response of such systems to irradiation, but also by the energetics of the atomic configurations, especially when the system is electrically charged. We suggest using spatially localized electron irradiation for making carbon islands and ribbons embedded into BN sheets. We further study the magnetic and electronic properties of such hybrid nanostructures and show that triangular carbon islands embedded into BN sheets possess magnetic moments, which can be switched on and off by electrically charging the structure.

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Graphene [1] is a two-dimensional (2D) material with a honeycomblike arrangement of carbon atoms. Its unique mechanical and electronic properties [2] have stimulated vigorous research on possible applications of graphene in nanoelectronics. Concurrently with the studies of graphene, individual flakes of hexagonal boron-nitride (h-BN) were isolated and characterized [3,4]. This material has the same hexagonal atomic network as graphene, but alternating boron and nitrogen atoms substitute for carbon. This system is a semiconductor with a gap of more than 5 eV [5], while graphene is a semimetal.

Taking into account the similarities in the atomic structure of these two materials and differences in the electronic characteristics, it is tempting to explore the possibilities for creating hybrid BCN nanosystems, with the electronic properties defined by the relative concentration of B, C, and N atoms. Numerous attempts to manufacture graphene with nitrogen or boron dopants have indeed been made [6–8], including chemical vapor deposition growth of BCN sheets [9]. However, it was shown [10] that such hybrid materials are thermodynamically unstable and that h-BN and graphene tend to segregate, as also confirmed by theoretical studies [11]. An alternative approach to engineering the composite BCN 2D materials was recently demonstrated [12,13]. Post-synthesis substitutional carbon doping of h-BN sheets [13] and BN nanotubes [12] was implemented via in situ electron-beam irradiation inside a transmission electron microscope (TEM), with the system being transformed from electrical insulator to conductor. It was found that carbon atoms coming from the decomposition of hydrocarbon molecules take mostly the positions of boron atoms, and it was deemed that the substitution process is dominated by the knock-on electron damage followed by mending of the boron-vacancies with carbon atoms. Such an interpretation, however, is inconsistent with nearly the same probabilities for knocking out B and N atoms [14,15] at high (300 keV) electron energies used in the experiments. Moreover, using essentially the same technique, C impurities were introduced into BN sheets at electron energies of 60 keV [16], which is below the threshold for any atom displacement [14,15].

In this Letter, by employing ab initio simulations, we show that although radiation damage should definitely play a role in the substitution process, the main driving force for the substitution is the energetics of the system, as it costs less energy to substitute a B-atom than N, especially when the system is positively charged. Taking into account recent experimental results reporting the development of triangular multivacancies under the beam, we further study the electronic structure of triangular carbon islands in BN sheets, and show that, in perfect agreement with Lieb’s theorem [17], such quantum dots possess magnetic moments which can be switched on and off by charging the system.

In our simulations we used the spin-polarized density-functional theory (DFT) as implemented in the plane-wave-basis-set VASP [18] code. We used projector-augmented wave potentials [19] to describe the core electrons and the generalized gradient approximation of Perdew, Burke, and Ernzerhof (PBE) [20] for exchange and correlation. As the PBE approach normally underestimates the band gap width, more accurate calculations were also carried out using the Heyd-Scuseria-Ernzerhof (HSE) functional [21]. A kinetic energy cutoff of 400 eV was used in all simulations. We used supercells of different sizes (32–200 atoms) with different numbers of B and N atoms substituted with C atoms. The k points for the integration over the Brillouin zone were generated by the Monkhorst-Pack scheme with the account for the supercell size. Such a simulation setup has been demonstrated to be adequate for...
the modeling of defects and impurities in 2D systems with
covalent bonding [15,22,23]. The finite-supercell-size cor-
corrections [24–26] for charged configurations were analyzed
by extrapolating the total-energy results to the dilute limit,
converging the formation energies within 0.2 eV. The DFT
molecular dynamics (MD) simulations of atom dynamics
after electron impacts were carried out as in Ref. [15].

The examples of the studied structures are shown in
Fig. 1. Throughout this paper, we use the following nota-
tions: CB means that a single B atom was substituted with
C atom, while 4C3B1N stands for a triangular island formed
by substituting 3 B and 1 N atoms with 4C atoms, etc. As
expected, the PBE calculations for a pristine h-BN sheet
underestimate the gap (4.56 eV), while the HSE functional
provides a semiconducting gap of 5.56 eV, the value which
is in a good agreement with the results of quasiparticle GW
calculations [27].

We base our analysis of the electron-beam-mediated sub-
stitution process on calculations of the formation en-
ergy $E_f$ of substitutional defects composed from $K_C$
carbon atoms embedded into the h-BN matrix as follows:

$$E_f(K_C^q; e) = E_{tot}(K_C^q) - E_{tot}(BN) + K_B\mu_B + K_N\mu_N$$

$$- K_C\mu_C + q(\mu_e - E_V),$$

where $E_{tot}(K_C^q)$ and $E_{tot}(BN)$ are the total energies of an
h-BN system with defect and defect free structure, respec-
"ively; $K_B$ and $K_N$ are the numbers of B and N atoms
replaced by $K_C$ C atoms ($K_B + K_N = K_C$). $\mu_B$, $\mu_N$, and
$\mu_C$ are the chemical potentials of the elements present in
the system. In an h-BN sheet in equilibrium

$$\mu_{BN} = \mu_B + \mu_N,$$

where $\mu_{BN}$ is the total energy per BN pair in the h-BN
sheet. We chose $\mu_C$ the same as in graphene. The choice
of $\mu_B$ and $\mu_N$ is normally defined by the growth condi-
tions of a material. However, we deal with post-growth
conditions and assume that the original system is
stoichiometric. As $N_2$ molecules should be present in the

![FIG. 1 (color online). Examples of the structures studied in our calculations, including single carbon atom impurities in the h-BN sheet, four carbon atom islands with boron (4C3B1N structure) and nitrogen (4C3B1N) termination, as well as 16-atom (16C6B10N structure) and 25-atom (25C15B10N structure) islands.](035501-2)

![FIG. 2 (color online). Formation energies $E_f$ of substitutional defects in different charge states as functions of electron chemical potential $\mu_e$ (a)–(d). Single carbon atom substitutional impurity in B-atom (a) and N-atom (b) positions. Four C atoms substituting 3 B and 1 N-atom (c) and 1 B and 3 N atoms (d). All the results are for the 200-atom supercell with different defects (PBE calculation). The numbers stand for the charge states. $E_f$ as a function of $K_C$ for various charge states and for preferential substitution of N and B atoms (e). $E_f$ vs $K_C$ for positive charge states and for preferential B substitution giving the lowest formation energies for each $K_C$ (f). Note that the values of $E_f$ are negative.](035501-2)
conduction band $E_C$ [e.g., $\mu_e = 3$ eV for preferential N substitution, as in Fig. 2(e)], then substitution of N atoms is also possible indicating that by charging the structure during irradiation, an additional control over substitution process can be gained.

We received qualitatively the same results for all other structures we studied (up to 81 C atoms in the 200-atom BN supercell): it is energetically favorable to have an N-terminated triangular defect with a larger number of missing B atoms, especially for positive charge states of the system, as evident from Figs. 2(e), 2(e), and 2(f). We further studied effects of charging on the energetics of the substitution process by comparing the total energies of neutral and charged reference systems including a C adatom on pristine BN sheet as well as those with C atoms in substitutional positions and the corresponding B/N adatoms on BN sheet. In agreement with the results of $E_f$ calculations presented in Fig. 2, we found that positive charge makes the energy required to “swap” a B atom with a C lower than to swap an N with a C atom, while negative charge gives the opposite result.

Keeping in mind the well-known deficiencies of the DFT-PBE approach, we checked the formation energies of the substitutional defects presented in Fig. 2 by repeating the calculations in a 32-atom supercell using the HSE functional. Figure 3 shows the lowest (among all charge states) formation energies of the substitutional defects presented in Fig. 2 as functions of $\mu_e$ calculated using the PBE and HSE functionals. Here the range of $\mu_e$ is normalized to the corresponding value of the gap. The PBE and HSE results agree well, indicating that the PBE approach can be used for the calculations of the formation energy of substitutional defects in BN systems if $\mu_e$ is rescaled using the ratio of the PBE and HSE gaps.

Although the N-rich environment matches better the experimental situation, we also studied the dependence of the results on the choice of $\mu_B$ up to the B-rich environment by the example of substitutional impurities formed by single C atoms. We found that even if $\mu_B = -7.62$ eV corresponding to the lowest energy $\alpha$-rhombohedral phase of boron [29] is taken as the reference, the substitution of B atoms is still preferable for positively charged systems.

The lower formation energies of defects with N termination are consistent with the preferential substitution of B atoms reported in the experiments [12,13]. Moreover, the energetics may actually explain why mostly B atoms are substituted, as the asymmetry cannot be understood simply as a dynamical effect: at electron energies of 300 keV used in the experiments the cross section for knock out of B and N atom is roughly the same [14,15], so that if C atoms simply take the place of missing atoms at vacancies, the ratio of B/N atoms should be the same as before irradiation. Our results indicate that substitution of B atoms may be thermodynamically preferable, so that irradiation may simply facilitate the substitution process by driving the system away from equilibrium (electrons with impact parameters exceeding the knock-out value transmit kinetic energy to the recoil atoms, which may be high enough for local chemical reactions, e.g., to swap host and C atoms), and may explain the apparent contradiction. The fact that C impurities were introduced into BN sheets at electron energies below the knock-out threshold [16] also indicates that irradiation-mediated reactions can occur, opening new avenues for irradiation-assisted engineering of nanostructures [30].

Nevertheless, to assess the role of kinetics, we calculated the displacement threshold $T_d$ for C impurities and the neighboring B and N atoms. $T_d$ is the minimum kinetic energy transferred to an atom in the system required to sputter the atom. $T_d$ governs the displacement rate, and the corresponding electron threshold energy can be estimated [14,15]. We found that $T_d$ for the C atom is 1.5 eV less than for B atom in the pristine system (independent of the type of the neighboring atoms), giving rise to a 40% increase in the displacement rate (for 300 keV used in the experiments) as compared to that for a B atom in the pristine material. This means that substitution of individual atoms can hardly be explained by dynamical effects only. $T_d$ for the neighboring B atoms ($C_N$ defect) drops by 1.2 eV, and by 3.6 eV for N atoms ($C_B$ defect) as compared to the corresponding values in the pristine system. The displacement rates for these atoms will increase by 20%–30%, which may facilitate the growth of triangular defects. We stress, however, that the actual substitution process may be more complicated, and involve beam-stimulated chemical reactions. Overall, it is much easier to explain the experimental results by the thermodynamics than kinetics of the systems.

Finally, we studied the electronic structure and magnetic properties of substitutional defects representing individual C atoms and triangular islands embedded into the $\text{h-BN}$ matrix. The latter may be manufactured if a triangular vacancy-type defect is produced first in a BN sheet by irradiation, [31,32] then carbon deposition, or by focusing the beam in a certain area when hydrocarbon molecules are deliberately introduced into the TEM chamber [12,13]. It was predicted that freestanding triangular graphene flakes of finite sizes should exhibit intriguing magnetic properties [33].
FIG. 4 (color online). Magnetic moments of triangular carbon islands embedded into BN matrix as a function of the difference of the number of substituted nitrogen ($K_N$) and boron ($K_B$) atoms. The insets show the magnetic moments of individual C atoms and four-atom defects as a function of $\mu_e$. Green and red lines correspond to preferential N/B substitution. Depending on the defect type, the magnetic moment can be switched on and off.

We found that such structures, if produced, should be stable and that although $h$-BN is a nonmagnetic material, BN sheets with triangular carbon islands possess magnetic moments. The value of the moment depends on the number of substituted atoms and naturally on the charge state of the defect. Figure 4 shows the magnetic moments of the defects as a function of the difference of the number of substituted nitrogen ($K_N$) and boron ($K_B$) atoms. The removed B and N atoms can be associated with C atoms in different sublattices in graphene, so that, in perfect agreement with Lieb’s theorem [17], the total magnetic moment of a carbon flake (neutral charge state) is simply the difference in the number of atoms on the two sublattices. The analysis of the electronic structure of the substitutional defects shows that the magnetism comes from spin-polarized defect-induced states in the gap, which are spatially localized at the defects. Our results also indicate that the magnetic moment can be varied by electrically charging the system, i.e., by changing the number of electrons in the quantum dots associated with the defects, as demonstrated in the insets presented in Fig. 4.

To conclude, using DFT simulations we showed that the electron-irradiation-mediated substitution of B/N atoms in $h$-BN nanostructures with C atoms is governed not only by the response of the system to irradiation, but also the energetics of different defect configurations, which explains the puzzling experimental observations [12,13,16] of C doping of $h$-BN systems. As the substitution process can be controlled by electrically charging the system during irradiation, our results indicate that by using a focused electron beam and simultaneously charging the sample one can manufacture spatially localized hybrid structures which should exhibit intriguing electronic properties [34,35], and controllable magnetism. This technique, which is a combination of electron irradiation and beam-assisted deposition of the material deliberately introduced into the TEM chamber, can be applied to any 2D material (or 1D systems like nanotubes), including recently reported inorganic 2D structures [36].

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