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Substrate-assisted nucleation of ultra-thin dielectric layers on graphene by atomic layer deposition

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We report on a large improvement in the wetting of Al₂O₃ thin films grown by un-seeded atomic layer deposition on monolayer graphene, without creating point defects. This enhanced wetting is achieved by greatly increasing the nucleation density through the use of polar traps induced on the graphene surface by an underlying metallic substrate. The resulting Al₂O₃/graphene stack is then transferred to SiO₂ by standard methods. © 2012 American Institute of Physics.

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The growth of ultra-thin dielectric layers on graphene is central to unlocking several of its electronic applications. For example, RF transistors require thin high dielectric constant (high-K) gate oxide layers for good electrostatic control of the channel. Lateral spin valves require electron transparent tunnel barriers to overcome the impedance mismatch and allow spin injection while bilayer pseudospin field effect transistors require a tunnel barrier to couple the two stacked graphene layers. Interestingly, continuous 1 nm sputtered Al₂O₃ films have already been grown on few-layer graphene. Technologically, it is desirable to use atomic layer deposition (ALD) to deposit dielectrics on graphene, but so far an additional seeding or interfacial layer is needed to yield continuous ALD oxide films ≤10 nm thick, due to the poor wetting of graphene. The seed layers can be either inorganic (1–5 nm thick evaporated Al, Hf, Ti, or Ta layers oxidized in air before transfer to the ALD system) or organic (spin-coated films of low-K polyhydroxystyrene derived polymer, or 2.5 nm films of ozone activated poly-vinyl alcohol, or perylene tetracarboxylic acid). Indeed, the surface of a defect-free, clean graphene sheet is inert under the soft chemical ALD process.

Increasing the wetting of ALD dielectrics on graphene would thus allow a seed layer to be avoided. In this direction, Shin et al. proposed to increase the surface energy of graphene by an O₂ plasma, which improves wettability, but at a cost of inducing defects in the graphene lattice.

In this paper, we demonstrate an alternative method to enhance wetting of ALD dielectric films on monolayer graphene without compromising its crystalline structure or seeding its surface. We show how to strongly improve wetting of graphene by tuning its surface energy using the substrate, as the graphene/metal proximity creates a dense network of polar traps and nucleation sites on the graphene surface. We focus experimentally on 10 nm Al₂O₃ films grown by ALD on graphene materials supported on different substrates: graphite, graphene grown by chemical vapour deposition (CVD) transferred on SiO₂, CVD graphene grown on Cu, and CVD graphene grown on a Ni-Au alloy. The Al₂O₃ films are grown in a Cambridge Nanotech Savannah system using tri[methyl]aluminum as precursor and water as oxidant, both carried in a 20 sccm flow of N₂ for 100 cycles, at growth chamber temperatures Tch leading to nominal 10 nm thick Al₂O₃ films. In all presented experiments, precursors exposure doses are kept constant (1 Torr, 0.015 s pulses) and CVD-like reactions in the chamber are prevented by varying with temperature purges times between ALD half-reactions.

For reference, we first grow 10 nm Al₂O₃ films on highly oriented pyrolitic graphite (HOPG) at different Tch (80°C with 60 s purges, 150°C with 20 s purges, 200°C with 8 s purges). The resulting films are characterized using a Hitachi S5500 scanning electron microscope (SEM). We found poor wetting of the Al₂O₃ on HOPG for all Tch, as reported by others. Films grown at lower temperatures have higher coverage, but are still clearly discontinuous. SEM images of the best-wetted films on HOPG (at Tch = 80°C) in Fig. 1(a) are clearly discontinuous, indicating an island-like Volmer–Weber growth mode.

We then investigate oxide ALD on CVD graphene. We focus in particular on high quality graphene (Raman D/G ratio ~ 0.05 at 532 nm) grown on Cu at ~1000°C in controlled Ar:H₂:CH₄ atmospheres. The graphene sheet is placed on a SiO₂ substrate using a standard transfer process with a PMMA carrier film and a FeCl₃ Cu-etch and is then annealed in ultra-high vacuum (~10⁻⁶ mbar). Compared to HOPG, wetting is expected to be better on transferred CVD graphene, as the transfer step can induce some defects. However, 10 nm ALD Al₂O₃ films grown at 80–200°C on transferred CVD graphene also leads to poorly wetted films similar to those formed on HOPG (Fig. 1(b)). Direct ALD of continuous ≤10 nm Al₂O₃ films on HOPG or on transferred CVD graphene on SiO₂ is compromised.

We instead tried performing the ALD step directly onto the CVD graphene on the Cu foil step (Fig. 2(a)), before the transfer. Importantly, the study is carried out on a similar piece of graphene/Cu foils as used previously for transferred graphene on SiO₂. Hence, the structural quality should be better on the un-transferred sheet and thus one might expect the wetting to be worse. The ALD process is then repeated at each temperature. Surprisingly, the resulting Al₂O₃ films show dramatically improved wetting in the SEM. At Tch = 200°C, the coverage is already much improved compared to that on graphene/SiO₂, with coverage increasing from ~60% to >90%. The improvement is even
more notable at $T_{ch} = 80^\circ C$ (Fig. 2(b)), as SEM reveals no structural defects in the $\text{Al}_2\text{O}_3$ film, in clear contrast to that found on graphite (Fig. 1(a)) or on transferred graphene/SiO$_2$ (Fig. 1(b)). Concerning the growth of thinner oxide films by ALD, Fig. 2(c) shows a 3 nm $\text{Al}_2\text{O}_3$ film after 30 ALD cycles deposited at $T_{ch} = 80^\circ C$. Clearly, wetting of a 3 nm $\text{Al}_2\text{O}_3$ film on graphene/Cu is greatly improved compared to wetting of a thicker 10 nm film on graphene/SiO$_2$.

Many applications require the graphene film to be ultimately on an insulating substrate. An $\text{Al}_2\text{O}_3$/graphene layer, equivalent to that in Fig. 2(b), was transferred on a SiO$_2$ substrate using a standard transfer process (Fig. 3(a)). Significantly, the resulting $\text{Al}_2\text{O}_3$/graphene/SiO$_2$ stack can then be obtained over large areas following this scalable transfer process. The structural quality of the graphene in this stack was checked using micro-Raman mapping (Renishaw InVia system) as shown in Fig. 3(b). The observed uniform low intensity of the Raman D peak relative to the G peak in Fig. 3(c) confirms that the sp$^2$ structure of the graphene is preserved, even after ALD and transfer steps. The simple inversion of the order of the transfer and of the ALD growth steps allows us to benefit from the increased coverage of the $\text{Al}_2\text{O}_3$ film observed in Fig. 2(b), providing a high quality encapsulated graphene layer.

To confirm the significance of our observations, complementary experiments are conducted at different $T_{ch}$ and on different graphene-like systems. In particular, monolayer graphene is also grown by CVD on a Ni-Au alloy, specifically tuned to allow growth of high quality monolayer graphene at a low ($450^\circ C$) temperature, under a flow of C$_2$H$_2$. Also, on a different set of samples, varying the exposure conditions yields the growth of multilayer graphene sheets ($\sim 5$–10 layers) on either Cu or Ni-Au. Again, 10 nm of $\text{Al}_2\text{O}_3$ are deposited by ALD on these samples at three different $T_{ch}$. All the observations made in terms of $\text{Al}_2\text{O}_3$ coverage for each $T_{ch}$ are reported in Fig. 4(a). Two distinct trends are seen. A first set of experiments (on HOPG, monolayer/SiO$_2$, multilayer/Cu, and multilayer/Ni-Au) gives results corresponding to those in Fig. 1: A decrease of $T_{ch}$ leads to improved coverage but with a poor wetting in all cases, the $\text{Al}_2\text{O}_3$ coverage remaining below 90%. A second set (on monolayer/Cu, monolayer/Ni-Au) gives results corresponding to those in Fig. 2. The $\text{Al}_2\text{O}_3$ coverage is much improved compared to set 1 (above 90% in all cases) and films appear continuous for $T_{ch} = 80^\circ C$. We have observed similar trends for other ALD grown oxides (in particular, HfO$_2$ grown from water and tetrakis[dimethylamido]hafnium precursors).
by only an atomically thin graphene layer, our coverage results suggest a nucleation density of $\sim 10^7 \times 10^6$ per $\mu m^2$, and that other processes help increase wetting. The graphene/metal proximity has modified graphene’s overall wettability.

One origin is the wetting “transparency” of graphene monolayers, noted by Rafiee et al. $^{26}$ These authors compared water contact angles on Si, Au, and Cu substrates with and without graphene to show that water adhered better on graphene when it is in intimate contact with a metallic substrate. This wetting transparency was also noted by Roos et al. $^{27}$ for bis(terpyridine) molecules adsorbed on graphene/Ru(0001). Moreover, Roos et al. $^{27}$ emphasised finite molecular adsorption on graphene/metal systems due to corrugations and direct interactions with graphene. In this direction, Pollard et al. $^{28}$ and Zhang et al. $^{29}$ showed by ab initio calculations and scanning tunnelling spectra that interaction between a monolayer graphene and an underlying metallic Rh(111) or Ru(0001) substrate creates a network of polar sites on the top surface, with lower adsorption energies for molecules like phtalocyanine, pentacene, or perylene tetracarboxylic diimide. This effect was previously seen for naphtalhlocyanine and cuprate-phtalocyanine on a h-BN monolayer on Rh(111).$^{30-32}$

Polar sites are a peculiarity of monolayer two-dimensional crystal/metal interactions. $^{27-32}$ They result from combined charge transfer (constrained by the difference of graphene and metal work functions) and wrinkles of the graphene sheet (constrained by the mismatch of the graphene crystal with the underlying metal). These studies suggest that the surface density of such polar traps ($N_0$) is $\sim 10^7 \times 10^6$ per $\mu m^2$. $^{25}$ As $N_0 \ll N_n$, this indicates that defects may contribute to wetting $^{18,22}$ but are not dominant. By contrast, when the surface is separated from a metallic substrate

We now consider the origin of the difference between the two experiments in Fig. 4(a). We first estimate from Fig. 1 the nucleation density ($N_n$) on HOPG or graphene/SiO$_2$ to be 2000–3000 per $\mu m^2$, while the Raman D/G ratio in Figs. 3(b) and 3(c) corresponds to a defect density ($N_d$) of $\sim 100–200$ per $\mu m^2$. $^{25}$ As $N_d \ll N_n$, this indicates that defects may contribute to wetting $^{18,22}$ but are not dominant. By contrast, the graphene transparency and the polar site network artificially modify the surface energy of the graphene sheet. This in turn alters the energy balance during the first ALD cycles from a water cluster formation/high desorption case $^{22}$ to a more homogeneous adsorption of water (see Fig. 4(b)). This explains the transition from an island-like Volmer–Weber ALD growth (Fig. 1) to a more two dimensional Frank–van der Merwe ALD growth mode (Fig. 2) as water molecules adsorb more homogeneously over the graphene surface. Critically, the metal/graphene combination provides enhanced wetting effect without creating structural defects in the graphene$^{18,22}$ (Figs. 3(b) and 3(c)).

While our experiments show that the effect of the substrate can result in a drastically higher Al$_2$O$_3$ coverage (from Fig. 1 to Fig. 2), we believe that a further enhancement is possible, and could allow better wetting at $T_{ch} > 80^\circ C$. Indeed, it has been shown that the energetic landscape of the graphene sheet can be tuned by the selection of the metallic substrate: For instance, the trapping energy of the surface polar sites and their distribution on the surface as a regular superstructure template depend on the graphene/metal couple $^{27-33}$ which may mean further optimisations of the water/surface interaction is possible. One could also use substrate assistance to provide a more controlled molecular coating of the graphene surface (Ref. 29 reports anisotropic trap energies forcing specific orientations of polar molecules), e.g., when using self-assembled monolayers as seeds for the ALD growth. Also investigation of graphene interactions with certain insulators may prove to lead to similar effects.
In summary, we have shown a dramatic increase in the wetting ability of ALD Al$_2$O$_3$ films by the assistance of a metallic substrate (Cu, Ni-Au) beneath the graphene layer, compared to thicker graphite layers or inert SiO$_2$. This assistance is conveniently fulfilled by the metallic catalysts commonly used during the CVD growth of graphene. This allows the nucleation of ultrathin Al$_2$O$_3$ films on clean graphene sheets, without introducing additional defects or seeds. The resulting Al$_2$O$_3$/graphene stack can then be routinely transferred onto a substrate of choice (here SiO$_2$). We highlight the role of the graphene/substrate interaction, which modifies the energetic landscape of the graphene surface and allows a more efficient trapping of water molecules during our Al$_2$O$_3$ growth by ALD. We envision this phenomenon to be also of importance for other controlled growths and coatings on graphene.

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