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Understanding noninvasive charge transfer doping of graphene: a comparative study

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Abstract
In this work, we systematically investigate and compare noninvasive doping of chemical vapor deposition graphene with three molecule dopants through spectroscopy and electrical conductivity techniques. Thionyl chloride shows the smallest improvement in conductivity with poor temporal and thermal stability and nitric acid induces the biggest sheet resistance reduction with modified stability. Molybdenum trioxide doping stands out, after thermal annealing, with both causing a significant sheet-resistance reduction and having superior temporal and thermal stability. These properties make it ideal for applications in advanced electronics. Theoretical studies based on the van der Waals density functional method suggest that cluster formation of molybdenum trioxide underpins the significant reduction in sheet resistance, and the stability, that arises after thermal annealing. Our comparative study clarifies charge transfer doping of graphene and brings understanding of the weak-interaction nature of such non-destructive doping of graphene. Our work also shows that we can use weak chemisorption to tailor the electronic properties of graphene, for example, to improve conductivity. This ability open up possibilities for further use of graphene in electronic interconnects, field effect transistors and other systems.

1 Introduction

Graphene is attractive due to its very high carrier mobility and current-carrying capacity. It is ideal for applications as transparent conductors, electronic interconnects, field effect transistors and more [1]. Graphene doping is being investigated as a viable technique to tailor its electronic properties and improve conductivity. Doping of graphene through functionalization, via either thermal treatment or plasma treatment create many defects in the graphene, causing damage to the electronic structure and increased scattering and very difficult to control these processes. Surface-charge transfer based doping is also being investigated in various carbon materials [2–4]. It is important that the adsorption binding is not so strong that it then disrupt the graphene properties and not so weak that there is no thermal stability. However, some of these doping mechanisms also involve adhesion of organic species that are themselves large enough that they can be held firmly in place by van der Waals (or more generally nonlocal-correlation) [5] forces. Similarly, small molecules (that may by themselves desorb) may cluster, thus gaining a large physical extension that still permits it to be held in place by weak chemisorption. With weak chemisorption one will have a finite charge transfer but one will not have any significant graphene deformation [6, 7]. In effect, one achieves the analogy to modulation doping (known from semiconductor-hetero structure designs) and can thus suppress scattering.

For example, metal chlorides [8–12], organic dopants [13–17] and volatile oxidants (HCl, H2SO4, HNO3, SOCl2) have been studied in recent years as viable dopants for surface transfer doping of graphene [18–21]. Kasry et al. [22]...
studied the effect of p-type doping on graphene using nitric acid to reduce its sheet resistance to a value closer to conventional transparent conducting oxides. They observed a reduction in sheet resistance by a factor of 3 and yielded films consisting of eight stacked layers of graphene with a sheet resistance of 90 ohms/sq. and a transmittance of 80%. They attribute the reduction in sheet resistance to the formation of charge transfer complex between graphene and nitric acid described by the following reaction which was proposed by Filiaux et al. [23]. Das et al. [24] dip casted pristine graphene in HNO3 solution of varying concentrations of HNO3. The work function was increased from 4.52 to 5.31 eV causing a shift in the Fermi level in graphene which gave it a p-type doped character with increased hole concentration. Bae et al. [25] used wet chemical doping using HNO3 on 30-inch chemical vapor deposition (CVD) graphene made using a roll to roll process and obtained films with sheet resistance of about 125 ohms/sq. and an optical transmittance of 97.4%. They further improved these values to about 30 ohms/sq. with 90% transparency by making a doped four layer film using layer by layer stacking.

Eda et al. [26] demonstrated that SOCl2 treatment reduced sheet resistance in graphene oxide films and used it as a transparent hole collecting electrode in organic photovoltaic devices. Li et al. [27] studied the effect of SOCl2 doping of CVD graphene on photovoltaic properties of the graphene/silicon Schottky junction solar cell and found that it significantly improved the conductivity of graphene and improved the conversion efficiency of the cell by threefold. Later, two articles from the same laboratory investigated several dopants for applications in graphene/silicon heterojunction solar cells [28, 29]. The first article demonstrated p-type chemical doping using chloride and nitrate ions from SOCl2 and HNO3 respectively to enhance the Schottky junction in the solar cells [28]. They found that nitrate ions were better at reducing the sheet resistance and increasing the work-function for better charge separation and transport leading to an improvement in efficiency by a factor of 1.68. The second article explored the effect of doping via four volatile oxidants namely, HCl, HNO3, H2O2 and SOCl2 on the power conversion efficiency (PCE) of graphene/silicon solar cells and this study also evaluated the thermal stability [29]. They found that all of them showed an improvement in PCE with SOCl2 showing the best out of them. PCE was found to be decreased with time and SOCl2 and HCl cases showed better stability than the rest.

There are contracting observation reported for HNO3 and SOCl2 doping case. Cui et al. [29] reported that SOCl2 doped Graphene/Si Schottky heterojunction solar cells shows better power conversion stability than HNO3 doped case. However, Li et al., [28] demonstrated that nitrate ions from HNO3 were found to be more effective for reducing the sheet resistance and enlarging the work function of graphene for effective charge separation and transport compared to SOCl2 case. Meanwhile, Chen et al. [30] and Xie et al. [31] demonstrated effective and non-destructive p-type doping of epitaxial graphene by depositing thin films of MoO3 on its surface. Their results show that the large difference in work function between the epitaxial graphene (4.0–4.3 eV) and MoO3 (6.8 eV).

In case of MoO3 different doping mechanisms are reported. Some papers discuss charger-transfer induced due to annealing [32] and some discuss MoO3 cluster formation due to multistage doping, without annealing [33–37]. Formation of metal carbide (Mo–C) with annealing at very high temperature is also reported [38]. Wu et al. [39] conducted a similar electronic structure study of interface between MoO3 and CVD graphene and observed a hole density of about 5.44×10^{12} cm^{-2} and a lowering of Fermi level by 0.28 eV.

They report that the formation of surface negative dipoles due to electron transfer from the graphene to MoO3 causes an increase in the samples work function and the presence of gap states due to oxygen vacancies significantly reduces the hole injection barrier at the interface. Hellstrom et al. [32] demonstrated strong and stable doping of graphene and CNT films using MoO3 for transparent electrodes. They observe a reduction in sheet resistance after the interaction with the MoO3, and also observed that the sheet resistance further reduced upon annealing these systems in an inert gas atmosphere. In their experiments with graphene MoO3 bilayers, they observed a reduction of sheet resistance from 465 ohms/sq. to 340 per square. which was further reduced to 260 ohms/sq. 24 h after annealing with an overall improvement by 42%. Hellstrom et al. [32] have also demonstrated doping stability, noting that the sheet reduction of the conductivity was observed upon high temperature annealing. They claimed that the annealing drives the partial oxidation of the nano carbon and partial reduction of MoO3 much further than mere deposition and refer to it as activation of MoO3 towards chemical oxidation of nanocarbon which they found to be optimal around 450 °C. Quite recently, Meyer et al. [33] studied the interface structure of graphene with thermally evaporated MoO3 films and found that it gave rise to large 1.9 eV interface dipoles and a downward bending of the MoO3 conduction band towards the Fermi level of graphene causing a nearly ideal alignment of the transport levels. The Fermi level shifted downwards by 0.25 eV manifesting as a p-type doping with sheet resistance values lower than 50 ohms/sq. for few layer graphene. Santiago et al. [40] reported a two order improvement of electrical properties carbon nano tubes after evaporation with MoO3. There is no annealing effect in these carbon nanotubes-MoO3 systems; merely changing thickness gave rise to improve the electrical conductivity. It is therefore interesting to determine if different oxidation states on MoO3 are created [7] due to the formation of MoOx clusters.
The current work seeks and tests techniques that can effectively dope the graphene while causing minimal to no damage to the graphene structure. This is necessary for its application in advanced electronics but we have found no other comparative studies of non-destructive charge transfer doping of graphene using three kinds of dopants thionyl chloride (SOCl₂), nitric acid (HNO₃) and molybdenum tri-oxide (MoO₃) in the literature. Our work has the following key objectives: (i) establish protocols for non-destructively doping graphene using charge transfer doping to improve its conductivity for application in electronics; and (ii) determine the most effective doping scenario for applications in advanced electronics by comparing the merits of using SOCl₂, HNO₃ and MoO₃ as p-type dopants for graphene. Finally, we comment on the most effective doping scenario for applications in advanced electronics by comparing their merits.

2 Experimental section

2.1 Methods: synthesis of graphene, chemical doping and theoretical studies

The graphene was grown using a thermal CVD process with copper as the catalyst. Copper foil was used as the growth substrate which was first cleaned and then subjected to high temperatures in a thermal CVD setup in the presence of a carbon source to result in graphene growth. Three kinds of dopants SOCl₂, HNO₃ and MoO₃ were used for chemical doping experiment. All reagents were received from Sigma-Aldrich and used as-received without further purification. More details about the thermal CVD process and chemical doping are given at Supporting Information section.

Our theoretical analysis is based on first-principle density functional theory (DFT), using the van der Waals density functional (vdW-DF) method [41–43]. We use the consistent-exchange vdW-DF-cx version [44] as implemented in the Quantum Espresso (QE) code package [45]. We determine the adsorption-induced doping by computing the charge distribution on each atom using Bader analysis [46]. We also perform Car–Parrinello molecular dynamics (CPMD) simulations [47], again using QE package, to track the stability of binding and cluster formation.

The graphene is modelled by a 5 × 5 unit cell containing 50 atoms and a vacuum region of at least 12 Å. Ultrasoft pseudopotential with a cutoff energy of 40 Ry and a 4 × 4 × 4 Monkhorst-Pack k-point sampling is used in our DFT simulations. All atoms are allowed to move till the interatomic forces are smaller than 0.008 eV/Å.

For all of the CPMD simulations the Brillouin zone integration is performed using the Gamma-point only. The equation of motion is integrated using a timestep of 8 a.u. and an electron mass of 400 a.u. All adsorption systems are heated up to 350 K in canonical (NVT) ensembles with Nosé-Hoover thermostats. The stability of the adsorption systems is then studied in microcanonical (NVE) ensembles.

2.2 Characterization

Raman spectroscopy was carried out on the graphene samples before and after the doping process. A Raman microscope with a 638 nm laser along with a 1200 lines/mm grating was used for all measurements. The morphology of graphene was characterized by scanning electron microscopy (SEM). This process was carried out in a Zeiss Supra 60 VP instrument. X-ray photoelectron spectroscopy (XPS) was used to measure the composition, chemical and electronic states of the graphene samples before and after the doping process. Sheet resistance measurement was carried out in a 4-point probe setup (CMT-SR2000N). Temporal and temperature stability study of the doped graphene as a function of time was evaluated by measuring the sheet resistance of the doped samples over an extended period.

3 Results and discussion

In present study, CVD graphene has been non-destructively doped using a surface-charge transfer method to improve its conductivity for application in electronics. The efficacy of three kinds of dopants SOCl₂, HNO₃ and MoO₃ were evaluated with respect to the extent of improvement in the conductivity of graphene, effect on the graphene quality and most importantly, their stability with increasing time and temperature. The latter criteria is essential to determine the viability of their use in electronic applications. So, we comment on the most effective doping scenario for applications in advanced electronics by comparing their merits after careful evaluation.

The graphene samples were grown using thermal CVD on the copper foil, its morphology was observed using SEM. Details of the synthesis and transfer were given at the Supplementary Information section (Supplementary Figs. 1–5). Raman spectroscopy was performed on the transferred graphene samples to check the quality of the transferred graphene and indicate that the graphene obtained is predominantly single layer. Raman spectrum of one of the transferred graphene samples along with a map of ratios between the 2D and G peak intensities showed at Supplementary Fig. 6. The D band around 1340 cm⁻¹ indicates the level of disorder or defects in the graphene. A perfect crystal of graphene would not have a D band. As seen from the figure, the D peak is quite low and shows that the graphene obtained is of good quality. The two most prominent features expected in graphene, the G and the 2D bands are also visible. The G
band around 1601 cm\(^{-1}\) arises due to the in-plane stretching of carbon–carbon bonds and is a signature for all carbon allotropes with \(sp^2\) bonding. The most intense feature is the 2D band around 2645 cm\(^{-1}\) and is the second order of the D peak. The ratio between the 2D and G peak intensities (I\(2\text{D}/I\text{G}\)) are above 1 in all cases and most of them are close to or above 2. This ratio above 1, along with the narrow FWHM of the 2D band close to 30 cm\(^{-1}\) indicate that the graphene obtained is predominantly a single layer and verify the observations using SEM. It is worth mentioning that although the same process was used to prepare the graphene samples, there was some degree of variation in the quality of graphene obtained and the Raman spectra between samples. Therefore, each sample was analyzed individually before and after doping.

After the graphene samples were synthesized, they were doped with three kinds of dopants \(\text{SOCl}_2\), \(\text{HNO}_3\) and \(\text{MoO}_3\), and analyzed to observe the effect of doping through various characterization techniques. The details of the doping were given in method section and supplementary information section. The morphology of the graphene films after the different doping treatments were observed using SEM. Figure 1a, b and c, d show SEM images of graphene samples after the \(\text{SOCl}_2\) and \(\text{HNO}_3\) treatment respectively. The graphene flakes and grain boundaries are visible. An organic substance was observed after the graphene was exposed to vapors. The wrinkles that were observed in the untreated graphene seem to have reduced after the vapor treatment process in both the cases. There were no major rips or tears observed in the films for either of the cases showing that the process of vapor doping is not strong enough to cause major tears to the graphene as in other chemically converted graphene. The morphology of graphene after \(\text{MoO}_3\) deposition is showed at Fig. 1e, f. The \(\text{MoO}_3\) seems to have covered all of the graphene while the grain boundaries and wrinkles of the graphene underneath are still visible. It also appears that the \(\text{MoO}_3\) deposition is slightly more aggregated near the wrinkles and is more visible. SEM images of the sample after annealing have been shown in Fig. 1g–i. At a lower magnification, the graphene does not look much different compared to the undoped graphene. However, on closer examination the \(\text{MoO}_3\) has formed smaller islands which uniformly cover the underlying graphene. This was not observed in the samples after deposition and indicates that the \(\text{MoO}_3\) undergoes dewetting from the graphene film during the annealing.

Sheet resistance for the samples was measured for three samples before and after doping for each dopant species. The sheet resistance values before and after \(\text{SOCl}_2\) treatment in three different samples that have been numbered for convenience (Fig. 2a). It can be seen that a reduction in sheet resistance is consistently observed in all three samples. Sample 1.1 shows the biggest reduction of 41% while the smallest reduction is 37% in sample 1.3. The average reduction in sheet resistance from these three samples is about 38%. Figure 2b highlights the effects found in the case of \(\text{HNO}_3\) treatment. Here there is a significant reduction in sheet resistance consistently observed with the biggest reduction of 73% and an average reduction of about 70% which is significantly more than the \(\text{SOCl}_2\) case. Figure 2c shows the comparison between sheet resistance values for three samples before \(\text{MoO}_3\) deposition, after deposition and after annealing. It can be seen that for all three samples, the sheet resistance values initially increase after deposition. This is probably because the graphene materials are not coated with \(\text{MoO}_3\) immediately after they are transferred so they could be slightly degraded or contaminated when exposed in air. This verified by experiment results that shows the graphene resistance change with different storage duration, as shown in Fig. S11 in the Supporting Information. However, the sheet resistance values reduce significantly after annealing at 450 °C in argon atmosphere.

The previous reports on graphene doping with \(\text{MoO}_3\) all report a reduction in sheet resistance soon after the deposition, while Hellstrom et al. [32] reported a more significant reduction after annealing. A reduction in sheet resistance was not observed immediately in any of the samples and was only observed after the annealing process. The annealing process possibly helped improve the interaction between the graphene and the \(\text{MoO}_3\) causing the charge transfer and thus the reduction in sheet resistance. The biggest reduction in sheet resistance was 59% while the remaining two samples showed a reduction in sheet resistance by about 49% with an average reduction of about 52%. This variation in the extent of reduction in sheet resistance between the samples despite the same doping process could be due to the variation in the initial quality of the graphene and possibly some minor changes during the processing.

Raman spectroscopy was carried out on the graphene samples before and after the doping process to see the effect of doping on the graphene through changes in the spectra. Raman data obtained from the samples with the highest improvements (for each of the three doping candidates) are shown in the figures in this section. Figure 3a, b shows the Raman spectra of the samples before and after \(\text{SOCl}_2\) where the Fig. 3b shows the G band of the same spectra. One of the prominent features of p-type doping in graphene is an upshift of the G band. This can be clearly observed in this sample where the peak position shifts from 1599 cm\(^{-1}\) before doping to 1606 cm\(^{-1}\) showing an upshift by 7 cm\(^{-1}\). This shift in the G band is an indication of p-type doping and is known to depend on the amount of charge removed per carbon atom. We note that greater shift implies more charge is withdrawn, although this dependence is not linear. This prominent shift in G peak was observed in all the samples. A reduction in I\(2\text{D}/I\text{G}\) ratio was observed along with
Fig. 1 SEM images of graphene treated with three different dopants with different magnification; SOCl₂ vapors (a 10 kx and b 50 kx); treated with HNO₃ vapors (c 10 kx and d 50 kx); after MoO₃ deposition (e 10 kx and f 50 kx); after MoO₃ deposition and annealing (g 10 kx h 50 kx i 150 kx)
Fig. 2 Figure showing the change in sheet resistance of graphene sample after doping.  

a SOCl₂, b HNO₃, and c MoO₃. In case of SOCl₂ and HNO₃ sheet resistance values reduce after exposure, however in case of MoO₃ the sheet resistance values are increased after deposition and reduced significantly after annealing at 450 °C in argon atmosphere. (Color figure online)
a narrowing of the G band as can be seen from the Raman spectra which is expected from doping graphene due to suppression of the 2D band. Interestingly, it was observed that the ID/IG ratio reduced from 0.32 to 0.14 after the SOCl₂ treatment which corresponds to fewer defects in the graphene. This reduction was also observed to varying degrees in the other samples treated with SOCl₂ which suggests that the treatment reduces defects in the graphene and confirms its non-destructive nature.

Figure 4 compares the Raman spectra of the sample before and after HNO₃ treatment. In this case, the upshift in the G band position is even more pronounced where it shifts from 1601 to 1610 cm⁻¹ with an increase of about 9 cm⁻¹. A similar upshift was observed in the other samples treated with HNO₃ and was always larger than that observed in the SOCl₂ treated samples. Therefore more significant charge transfer effect takes place in this case and agrees with the extent of reduction in sheet resistance when comparing both the vapor doping scenarios. There was no major changes in the ID/IG ratio which suggests that the HNO₃ treatment does not introduce significant defects in the graphene. Figure 5 shows the Raman spectra of a graphene sample before MoO₃ deposition, after deposition and after annealing. It can be seen that although there is a minor upshift of the G band after the deposition from 1604 to 1605 cm⁻¹, the major shift by 15–1619 cm⁻¹ was observed after annealing indicating that the charge transfer takes place during annealing and agrees with the results of the sheet resistance measurements. Another interesting feature is the broadening of the G band and a second component seems to exist in the G band after annealing. The origin of this second component is unclear and is associated with either deposition or adsorption of some material on the graphene, strain in graphene or presence of local charge inhomogeneity. A reduction of the I₂D/IG ratio was also observed which is expected. There seems to be an increase in the ID/IG which suggests a slight increase in the defects in the graphene. However,
this was not observed in all the samples and would require further investigation.

X-ray photoelectron spectroscopy also used to confirm the presence of the dopant species and to observe their interaction. The XPS also allowed us to track shifts in the Fermi level of graphene. We studied XPS obtained from four different samples; undoped graphene, SC_{l_{2}} treated graphene, HNO_{3} treated graphene, MoO_{3} coated and annealed graphene. Precise quantification of the XPS spectra allows us to obtain composition but this approach also requires very exact fitting of the peaks with proper constraints. Survey spectra obtained from each of our four samples are given at Fig. 6. The predominant peaks have been labeled and the respective elements that would be expected were observed in each of the cases. Carbon, oxygen, and silicon peaks were observed in all the samples from the graphene and the underlying SiO_{2} substrate. Moreover, The Cl_{1s} and Cl_{2p} peaks were visible in the sample treated with SC_{l_{2}} vapors while N_{1s} peak was detected in the sample treated with HNO_{3} vapors. The sample doped with MoO_{3} shows the Mo_{3d}, Mo_{3p} and Mo_{3d} peaks. Apart from the expected peaks, there were also strong unexpected peaks such as the fluo-
rine and aluminum peaks which appear on all the samples the origin of which is unclear. A high resolution C_{1s} spectra were acquired for each of the sample.

The doped samples show varying degrees of shift when compared to the pristine sample (Fig. 7). The C_{1s} binding energy for the undoped sample is about 284.69 eV, while that of the SC_{l_{2}} has downshifted by 0.40–284.29 eV. In the case of the sample doped with MoO_{3}, the binding energy of C_{1s} is centered around 284.40 eV showing a shift of 0.29 eV. However, a very small shift of 0.02 eV for HNO_{3} doped sample was observed which is much smaller than expected. Therefore, the XPS measurements would need to be repeated and fitted carefully to extract precise changes in Fermi level of the graphene after doping. A high resolution spectra of Cl_{2p}, N_{1s} and Mo_{3d} levels acquired from the samples doped with SC_{l_{2}}, HNO_{3} and MoO_{3} respectively (Fig. 8). The Cl_{2p} peak around 199 eV seems to consist two peaks, one near 198.5 eV which corresponds to the presence of Cl^{-1} ions while the other around 200.3 eV which corresponds to Cl–C covalent bonds. The N_{1s} spectra from the HNO_{3} doped sample shows two main peaks, a large peak at 407.2 eV corresponding to C–N covalent bonding and another one around 400 eV which corresponds to the NO_{3}^{-1} ionic bonds. These results suggest that the C–Cl and C–N interactions take place due to the doping process and contribute to the enhancement in the electronic transport [24, 28, 48]. The Mo–3d peak seems to consist of three main peaks. The peaks around 232.6 and 235.8 eV correspond mainly to Mo_{6+} oxidation states and possibly contain smaller components from Mo_{5+}. The peak around 229.8 eV corresponds to the Mo_{4+} oxidation states. Previous works on doping of graphene with MoO_{3} where the samples were not subjected to annealing did not show any peak corresponding to Mo_{4+} oxidation states [30, 32, 33]. These Mo_{4+} oxidation states seem to become visible only after the annealing which strongly suggests reduction of MoO_{3}, indicating that it receives electrons from the graphene, thus causing more efficient charge transfer. These results agree with the observation by Hellstrom et al. [32]. Moreover, no major evidence of Mo–C bonding was observed from the Mo_{3d} spectra.

The temporal and temperature stability of the doped graphene as a function of time was evaluated by measuring the sheet resistance of the doped samples over an extended period. The results for the doping stability over time are reported at Fig. 9. We find that samples treated with SC_{l_{2}} show the least stability with time. In fact it quickly reaches 99.6% of the initial sheet resistance value before doping. HNO_{3} also shows a poor short term stability: while the initial reduction in sheet resistance is large, it goes up to 56.8% of the undoped value in just 2 days. However, thereafter it remains fairly stable beyond that reaching 66.3% of the undoped value at the end of 4 weeks. The increase in
sheet resistance with time is possibly due to reaction with the moisture in the air and also desorption of the dopants from the graphene to some extent. This could be reduced by adding a protective layer on top of the graphene after doping, however it would lead to other complications during patterning and depositing contacts for electronic applications.

The sample doped with MoO$_3$ shows fairly good stability with time where the sheet resistance increases gradually to 63% of the undoped value at the end of 4 weeks and 66% at the end of 6 weeks. In this case, the increase in sheet resistance is predominantly due to the effect of moisture in the atmosphere. It is worth mentioning that some of the increase in sheet resistance may be due to the damage caused by the probes during each measurement which is why the measurement was not carried out very often. This possible damage due to the probes could be avoided by depositing metal contacts on the graphene instead of using the probes directly on the graphene.

Figure 10 shows a quantification of doping stability with temperature. We track the percentage increase in sheet resistance for different dopants with increasing temperatures. The figure reports the percentage increase in sheet resistance which is plotted as a function of temperature. The percentage increase in sheet resistance is
much more pronounced for samples doped with HNO₃ and SOCl₂ than with MoO₃. The SOCl₂ doped sample reached 43.2% of its initial doped value while HNO₃ doped sample reached around 40% at 200 °C. This increase in sheet resistance is attributed to desorption of the dopants from the graphene at elevated temperatures. It should be noted that although the percentage increase in both these cases is quite similar, the initial reduction in the case of HNO₃ is significantly larger and therefore the increase in the value of sheet resistance when compared to the undoped value is less significant than for the SOCl₂ doped sample. The effect of temperature on the sheet resistance is not very significant for the samples doped with MoO₃ where it reaches only 12.4% of the initial doped value. This shows that the samples doped with MoO₃ has much better stability when compared to the other two doping scenarios.

3.1 Theoretical investigation

We also theoretically investigated the mechanisms of graphene doping by HNO₃, SOCl₂ and MoO₃ molecules. Our focus was on understanding the charge transfer effects that follow the weak chemisorption of these molecules on graphene. First, we simulated the adsorption structures in DFT and tracked the stability using CPMD performed at 350 K.

Figure 11 reports the results of our CPMD simulations for HNO₃ on graphene. We find that there will be a dissociation of hydrogen or of hydrogen and oxygen atoms leaving NO₃ or NO₂ molecules to diffuse on the graphene. Below, we also document that the NOₓ adsorption causes charge transfer, i.e., that the residual molecules are technically chemisorbed. However, the diffusion documented in the CPMD studies reflect a weak chemisorption nature for the NOₓ–graphene system and we do, in fact, find limited adsorption energies (per molecule or molecular unit) in all of the cases. The adsorption energy is calculated from $E_{\text{ads}} = -(E_{\text{gr}} + E_{\text{mol}} - E_{\text{gr} + \text{mol}})$, where $E_{\text{gr}} + E_{\text{mol}}$ is the adsorption system energy, $E_{\text{gr}}$ and $E_{\text{mol}}$ are the energy of isolated graphene and molecule. Regular chemical-bond formation does not account for behavior in the set of investigated adsorption cases and use of the vdW-DF method (instead of traditional semilocal DFT) is important for our theoretical characterizations.

Figure 12 summarizes our study of weak chemisorption of MoO₃ on graphene. In this study, we simulated the adsorption of a MoO₃ single molecule, a MoO₃ dimer as well as of three- and four-molecule clusters. Our CPMD simulations indicates that all (MoO₃)n adsorption structures are stable at 350 K. Our CPMD results from the four MoO₃ weak chemisorption problems further suggest that are significant clustering energies to drive formation of large adsorbate clusters and (at sufficient deposition) even assembly of MoO₃ coatings. Having identified optimal...
adsorption configurations, we extract total energies in the four systems and, in turn, define energy differences that characterize clustering among weakly chemisorbed MoO$_3$ on graphene. This clustering cohesive energy is calculated from $E_{cc} = - \left( E_{ads}^{MoO_3} - nE_{ads}^{MoO_3} - E_{Gr}^n + nE_{Gr}^{MoO_3} \right) / n$ and grows with the cluster size: it is 1.35 eV per molecule for a MoO$_3$ dimer, while it is 1.60 and 1.65 eV per molecule for (MoO$_3$)$_3$ and (MoO$_3$)$_4$ clusters.

The tendency for nucleation and assembly also explains why the thermal stability of the MoO$_3$ increases with thermal annealing: the MoO$_3$ clustering stabilizes the adhesion. Comparing the total energy of the adsorbed systems, of the clean graphene surfaces and of the molecule in the gas phase, we define adsorption energies $E_{ads}$. For larger adsorbates, weak chemisorption can easily provide binding in excess at the eV scale and beyond [5–7, 49]. Table 1 includes a comparison of the adhesion energy per molecules for the set of investigated dopants and dopant-cluster configurations. The table confirms that the MoO$_3$ cluster adhesion energy exceeds 1 eV already when reaching a three-molecular size.

Finally, our calculations permit us to track the doping and graphene modification that occurs at optimal adsorption configurations. Table 1 reports our vdW-DF-cx results for the adsorption distances, for the associated mean absolute displacement of the carbon atoms sitting immediately below the adsorbates, and for the resulting charge transfer (as asserted in a Bader analysis of the electron distribution in the optimal configurations).

Our calculations show that all three molecules can (p-) dope via weak chemisorption but also confirms that it is imperative to check stability. The HNO$_3$ adsorption causes dissociation and we have instead analyzed this strategy as a case of NO$_2$ and NO$_3$ weak chemisorption. Also, given the tendency for cluster nucleation, it is more important to characterize the doping impact of MoO$_3$ clusters instead of individual MoO$_3$ molecules.

Table 1 shows that p-doping efficiencies (as obtained in DFT) are in the order, SOCl$_2$ < (MoO$_3$)$_n$ < NO$_2$/NO$_3$. The adsorption of SOCl$_2$ molecules does not seem to lead to any significant charge transfer. Given the anchoring of dissociated NO$_3$ and NO$_2$, one can expect that the addition of HNO$_3$ molecules produce the highest doping efficiencies. In addition, one should note that adding HNO$_3$ molecules also causes a strong chemisorption of excess hydrogen and oxygen atoms, shown in Fig. 11c, d, that further affects the scattering (and doping).

Interestingly, our calculations show that the assembly of MoO$_3$ clusters causes the doping nature to change: we predict a n-type doping for the (unstable) case of single-molecule adsorption while clustering converts the MoO$_3$ adsorption into p-doping agents. Given our documentation of a clustering driving force, our theoretical studies corroborate the finding that annealing plays a critical role in the MoO$_3$-induced (p-)doping.

In fact, our set of theory results seem fully consistent with the experimentally observed trends in conductivities, but we can only provide a qualitative discussion subject to an assumption of the original nature of conduction in the CVD-grown samples. In general, chemical doping increases the density of free (electron or hole) charge carriers by charge transfer mechanisms that modify the Fermi
level of the doped. However, we do not know if the CVD-grown graphene samples were p or n doped from the outset and so we cannot trace the net impact that follows with predictions of the size of p-type doping. However, we can say this: if the samples was originally p-doped, the set of theory characterizations are consistent with the measured conductivity changes.

Subject to the stated assumption, the theoretical study does confirm the experimental observation that HNO₃ treatment gave rise to the highest resistance reduction but suffered from less stability. Overall, the theoretical characterization suggests that, with annealing and cluster assembly, MoO₃ holds advantages in terms of both thermal stability and by having a limited deformation at a fair charge transfer. However, the weak chemisorption of adsorption of MoO₃ clusters does cause some deformation of the underlying graphene sheet, and it will induce electron scattering.

### 4 Conclusions

In summary, we demonstrate the efficacy of efficient, non-destructive doping of CVD graphene using thionyl chloride (SOCl₂), nitric acid (HNO₃) and molybdenum trioxide (MoO₃). The samples were characterized using microscopy, spectroscopy and electrical measurement techniques. The
samples doped with HNO₃ showed the largest (immediate) reduction in sheet resistance with an average reduction by about 70% and SOCl₂ treated samples on the other hand showed the smallest reduction in sheet resistance with an average reduction by 38%. Whereas MoO₃ deposition on graphene increased the sheet resistance initially, this was significantly reduced after the annealing process yielding an average reduction by about 52% compared to the undoped samples. Analysis of morphology using SEM showed no severe damage to the graphene from any of the doping processes. Raman spectra on the doped samples showed an upshift in the G band for all the cases confirming the transfer of electrons from the graphene to the dopants, increasing the density of holes at the interface, thus effectively giving the graphene a p-type doped character.

The presence of the dopant species on the graphene was confirmed using X-ray photoelectron spectroscopy and their interaction with the graphene was analyzed from their core level spectra. Samples doped with SOCl₂ showed the poorest temporal and thermal stability with both increasing time and temperature while samples doped with HNO₃ performed better in both the scenarios. MoO₃ doped samples showed the best stability with increasing time and temperature and seems to be the best candidate for applications in advanced electronics which require thermal stability. Our theory modeling results indicate the adsorption configurations of the three molecules are intact (SOCl₂), dissociated (NO₂) and clustered (MoO₃)ₙ. The p-type doping efficiencies are positively related to the resistance reduction in experiments and we have provided an interpretation of why annealing is essential when we seek to use MoO₃ as a (stable) p-type dopant for graphene. The results could be very useful for graphene based interconnects and transparent electrodes where the resistance is a key factor.

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