Exclusive Substitutional Nitrogen Doping on Graphene Decoupled from an Insulating Substrate

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ABSTRACT: The on-surface synthesis of atomically flat N-doped graphene on oxidized copper is presented. Besides circumventing the almost standard use of metallic substrates for growth, this method allows producing graphene with \(~2.0\) at \% N in a substitutional configuration directly decoupled from the substrate. Angle-resolved photoemission shows a linear energy-momentum dispersion where the Dirac point lies at the Fermi level. Additionally, the N functional centers can be selectively tailored in sp\(^2\) substitutional configuration by making use of a purpose-made molecular precursor: dicyanopyrazophenanthroline (C\(_{16}\)H\(_{6}\)N\(_{6}\)).

INTRODUCTION

The Nobel Prize recognition to the discovery of graphene in 2010 boosted the efforts to produce this two-dimensional (2D) material, which opened a new paradigm of relativistic condensed matter physics. One of the most remarkable features of graphene originates from the linear dispersion of its \(\pi\) and \(\pi^*\) bands at the Dirac point in the vicinity of the Fermi level, which accounts for the description of pristine graphene as a zero-gap semiconductor.\(^1,2\) This material with outstanding physical properties has inspired the seek for a mechanism to control its behavior in a tunable manner, and nonsurprisingly, studies on the functionalization of graphene with the presence of heteroatoms such as N, B, and Si have appeared in an exponential way.\(^3\)–\(^6\) For instance, nitrogen-doped graphene (N-graphene) was first reported in 2009 from the pyrolysis of pyridine and ammonia in the arc discharge of graphite electrodes,\(^7\) but later studies have optimized its growth with a wide scope of techniques.\(^8\)–\(^13\) Several applications of N-graphene have already been tested toward a new generation of batteries,\(^14\)–\(^16\) oxygen reduction,\(^17\) hydrogen generation,\(^18\) biosensors,\(^19,20\) supercapacitors,\(^21,22\) and quantum dots.\(^23,24\) Ideally, the method used for the synthesis of graphene should be chosen according to the intended application.

Especially in applications where the control of the electronic properties plays a critical role in the performance, the growth mechanism should be understood. Different N bonding environments can affect the local electronic structure, so for N-graphene, the goal is to gain control on the electronic properties incorporating substitutional functional centers and the least possible vacancy-related defects. Also, the work function and carrier concentration of N-graphene are changed by the dopants.\(^25,26\) A decoupling from the surface can also open interesting application pathways. With all this in mind, on-surface synthesis can be one of the most versatile strategies to build such complex low-dimensional materials and use them as-grown. To some extent, this method can be related to chemical vapor deposition (CVD), in which vapors of pyrazine,\(^7\) ammonia,\(^27\) acetonitrile,\(^14\) and triazine\(^28\) among others have been used, but making N-graphene with mainly substitutional configuration is still extremely challenging with such methods. Studies carried out with ion implantation have shown that N can be incorporated onto the surface,\(^29,30\) and further work has proven the incorporation of the heteroatoms at the local level with techniques such as scanning tunneling microscopy, which require a metallic surface to ease characterization.\(^26,28,31\) However, CVD can offer a scalability advantage, but the effect of catalytic surfaces used must be carefully taken into account. For instance, a metallic substrate can negatively interfere with the performance of graphene, and this is the reason why fabricating graphene-based electronic devices requires several transfer processes to place either directly onto insulators or onto high \(\kappa\)-dielectric materials.\(^32\)–\(^34\) These
processes undermine the quality of graphene and make it prone to wrinkling, explaining, in turn, the considerable amount of research still looking for a clear route to fabricate high-quality graphene on suitable substrates.\(^{13,36-39}\)

Herein, we report an on-surface synthesis method to grow N-graphene using a nitrogen-rich polycyclic aromatic hydrocarbon as feedstock, namely, dicyanopyrazophenanthroline (see Figure 1).\(^{40}\) This derivative possesses a high C/N ratio (16:6); all the nitrogen atoms are incorporated into the aromatic framework in sp\(^2\) or other sp-N configuration, and it can be deposited by sublimation. The molecular deposition studies on different surfaces include single-crystalline Cu(111), polycrystalline copper foil, and the corresponding oxidized surfaces. Remarkably, the use of dicyanopyrazophenanthroline leads to the exclusive incorporation of N in sp\(^2\) configuration within the graphene hexagonal lattice from the first step with a doping level of \(\sim 2.0\) at \%, which represents at least double the amount of substitutional atoms reported for N-graphene grown on an insulating material (Table 1). It is important to remark that this value corresponds to substitutional doping if it is compared to the literature, where the doping values reported combine the portion of different bonding environments. Furthermore, N-graphene grown on oxidized Cu(111) shows an energy-momentum dispersion with the Dirac point lying at the Fermi level, demonstrating that the oxide layer decouples graphene from the substrate.

![Figure 1. Dicyanopyrazophenanthroline (C\(_{16}\)H\(_{6}\)N\(_{6}\)) molecule was used as a precursor for the synthesis of N-graphene. Its structure has favored the formation of N substitutional configuration. Left: C\(_{16}\)H\(_{6}\)N\(_{6}\) molecule. Right: N-graphene.](https://example.com/figure1.png)

#### METHODS

A dicyanopyrazophenanthroline C\(_{16}\)H\(_{6}\)N\(_{6}\) molecule was used as C/N feedstock, and its synthesis and capability to induce the incorporation on the graphene network have been reported elsewhere.\(^{40}\) We have grown N-doped graphene, and its spectroscopic characterization has been carried out in situ in ultrahigh vacuum (UHV) via photoemission spectroscopy (PES) in different energy ranges as explained later, as well as with low-energy electron diffraction (LEED). PES was performed using a dedicated spectrometer equipped with a monochromated Al K\(\alpha\) source (1486.6 eV) and a Scienta RS4000 analyzer. It is also equipped with a He II lamp operating at 40.8 eV for ultraviolet PES. Polycrystalline copper foil (0.025 mm) from Alfa Aesar and a Cu(111) single crystal (\(\phi 5\) mm \(\times\) 2 mm) from MaTeck GmbH were used as starting materials. For the sample preparation, the Cu(111) single crystal and the Cu foil were cleaned by Ar\(^{+}\) sputtering for several cycles for 60 min and then annealed at 800 K for another 60 min. C\(_{16}\)H\(_{6}\)N\(_{6}\) was subsequently deposited on the clean Cu(111) and the clean Cu polycrystalline copper foil at room temperature in UHV using a Knudsen cell working at \(\sim 400\) K. The deposition rates were monitored using a quartz micro balance. To obtain oxidized surfaces, the samples were exposed to air during 5 min, while additional metallic Cu(111) and Cu foil samples were kept under UHV conditions. The following annealing treatment at 1250 K was performed on all samples. Scanning transmission electron microscopy (STEM) measurements were performed with a Nion UltraSTEM 100 electron microscope operated at 60 keV.\(^{41}\) The TEM grid preparation was carried out using the polymethylmethacrylate (PMMA)-based technique, which included the coating of graphene with PMMA, etching of the copper foil with FeCl\(_3\) to place the suspended graphene on TEM grids, and, finally, the PMMA removal with acetone.\(^{42}\) Scanning electron microscopy was carried out using a Zeiss Supra 55 VP microscope.

#### RESULTS AND DISCUSSION

The use of C\(_{16}\)H\(_{6}\)N\(_{6}\) results in the effective growth of graphene on the substrates in question. Nevertheless, significant differences appear looking in their spectroscopic responses. First, our samples show graphene portions ranging from 0.5 to 2 \(\mu\)m independently from the surface where they were grown, which hints that the grain size distribution is linked to the feedstock rather than to the nature of the substrate (see a representative scanning electron micrograph of this type of material in Figure 2). Further microscopy measurements were carried out by STEM. This was carried out for analytical and structural inspection on suspended graphene transferred to TEM grids in the feasible cases as later described together with the surface structural analysis via LEED and ARPES. With these pieces of information, we are able to gather valuable information on the electronic dispersion (momentum vs energy) of N-graphene on Cu(111) compared to the oxidized counterpart. All these results are explained below in detail.

Focusing first on the molecular precursor that is a derivative of the phenanthroline molecule, featuring an extended \(\pi\)-conjugated system, it provides the initial advantage that it bears...

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**Table 1. Reported Studies on N-Graphene Directly Grown on Insulators or High \(k\)-Dielectric Materials**

| feedstock                              | N (at %) | N-configuration                  | reference |
|----------------------------------------|----------|----------------------------------|-----------|
| molecular nitrogen (N\(_2\))           | N.A.     | pyridinic/pyrrolic               | 36        |
| ammonia (NH\(_3\))                    | N.A.     | pyridinic                        | 37        |
| ammonia (NH\(_3\))                    | 0.1      | pyrrolic/sp\(^2\)                | 38        |
| hexaazotriphenylene (C\(_{18}\)N\(_{12}\)) | 7.8      | pyridinic/pyrrolic/sp\(^2\)      | 39        |
| pyridine (C\(_{4}\)H\(_{6}\)N)         | 0.6      | sp\(^2\) (2.5 eV)\(^{a}\)        | 13        |
| dicyanopyrazophenanthroline (C\(_{16}\)H\(_{6}\)N\(_{6}\)) | 2.0      | sp\(^2\) (1 eV)\(^{a}\)         | here      |

\(^{a}\)Observable fwhm at the N 1s in XPS.
multiple nitrogen sites susceptible to coordinate with the metal, including a strong chelating phenanthroline ring, a bridging pyrazine ring, and cyano groups (see Figure 1). These coordination sites are crucial to prevent the desorption of the molecular precursor during the high-temperature on-surface synthesis process we applied. In particular, at these temperatures, the molecules decompose and rearrange on the surface to provide the most stable lattice, given the local chemical composition under the given conditions. Here, the chemical nature of the feedstock will dominate the decomposition rates that in turn will affect key factors such as the local composition. However, a mechanistic overview of the reaction or any intermediate cannot be matched with well-established polymerization mechanisms.

The process was followed in-situ monitoring the cleanliness, purity, and, afterward, the molecular transformation with the use of XPS with an Al Kα (1486.6 eV) monochromated source. Survey scans and a closer inspection to the C, N, and Cu regions were performed for all samples. Also, the O₁s core level signal was taken into account for the appropriate cases. First, analyzing the processes using the Cu(111) surface, Figure 3, shows the C 1s and N 1s core level spectra recorded directly after the deposition of the C₁₆H₆N₆ molecule (top) and after the transformation into N-graphene (bottom). The Shirley background was subtracted in these spectra before deconvolution. The C 1s signal corresponding to the substrate with C₁₆H₆N₆ as-deposited reveals four components at 283.7, 284.3, 284.9, and 285.6 eV, which are ascribed correspondingly to the C−N, C=C, C≡N, and C≡N bonding environments, respectively. The ratio between the integrated area of these peaks (2:3:2:1) is in good agreement with the proportions in the chemical structure of the C₁₆H₆N₆ molecule (Figure S1 in the Supporting Information). It is important to remark that the absence of a notorious shift of the C 1s peak (and implicitly of its components) confirms that our material corresponds basically to monolayer coverage (Figure S2 in the Supporting Information). On the other hand, the two components observed in the N 1s spectrum (see Figure 3b) can be attributed to N=C (397.1 eV) and N≡C (398.5 eV). The surface described so far corresponds to the raw deposition of the molecules at room temperature, but various post-deposition annealing treatments were tested. For annealing temperatures lower than 1100 K, only minor changes were observed in XPS, which lead us to use higher temperatures for the molecular transformation into N-graphene (see spectra recorded at different temperatures in Figure S3 in the Supporting Information). Once the annealing temperature reached 1250 K, a major change was evident. In particular, the C 1s peak in Figure 3a narrows down significantly in Figure 3c,
suggesting the fading of at least two components, leading to a single chemical environment. Note that previous work on sp² C systems has allowed revealing the characteristic spectra for different structures including graphene.⁵,⁶,⁰,⁵¹ Here, the spectra have been fitted with a Voigtian profile by taking into account a Gaussian contribution associated with the resolution of our spectrometer with the monochromated X-ray source. In this specific case, this C 1s spectrum can be fitted almost exclusively by a single peak at 284.4 eV with a full width at half-maximum (fwhm) of 0.8 eV, which is in very good agreement with the values corresponding to graphene in synchrotron and lab-based photoemission experiments.³,³⁵,³³,³⁴ Furthermore, a shift of the C 1s can be expected because of the N incorporation on the lattice and the interaction with the specific metallic substrate, but the largest contribution to this core response arises from the sp²-hybridized C atoms. Regarding the N incorporation into the lattice of N-graphene grown on the Cu(111) substrate, Figure 3b,d shows the core level N 1s spectra where the components below 399 eV recorded straight after the molecular deposition correspond to the nitrilic and pyridinic bonding environments in agreement with the structure of the molecule.⁴⁰ On the other hand, after annealing, the N-graphene spectrum is notoriously different. A considerable number of studies reported in the literature have focused on N-graphene, but in seldom cases, both local and bulk sensitive methods have successfully proven to be a N incorporation profile. Table 1 shows a summary of the reports on N-graphene directly grown on oxides or high-κ-dielectric materials. In contrast with those and other reports in the literature on N-graphene, where the assignment of exclusive bonding environments is clearly limited by resolution, here, the N 1s spectrum shows a single peak at 400.6 eV with an fwhm of 1 eV. Only in this way, the spectra can be safely ascribed almost exclusively to substitutional nitrogen because it is obtained from a Voigtian profile that considers the intrinsic Gaussian experimental resolution and the lifetime Lorentzian broadening.⁵⁶ Furthermore, by analyzing the relative areas of the N 1s and C 1s peaks (corrected by the atomic cross sections),⁶,³³ a N content of 2.0 at % has been found in average in these N-graphene samples. The XPS results obtained for the polycrystalline foil are very similar to those obtained from the Cu(111) single crystal from the content and bonding configuration points of view. Also, the N-graphene samples grown on the oxidized surfaces were examined in XPS. The C 1s and N 1s responses have consistently shown a shift toward lower binding energies (~0.15 eV) and ~2.0 at % N in substitutional configuration (Figure S4 in the Supporting Information). An advantage of using the foil is that preparing microscopy grids for further observation is achieved with lesser effort. With this material, we performed annular dark-field (ADF) imaging in STEM pursuing an atom-by-atom identification.⁵⁶ The collected signal from the atoms is originated from Rutherford scattering, and it increases with the atomic number, so we are able to identify the dopants clearly. Figure 4 shows an atomically resolved ADF-STEM image of the N-graphene grown from a polycrystalline foil. The images show a regular pattern of atoms forming a hexagonal lattice with a nearest-neighbor distance of 0.14 nm characteristic between the C atoms.⁵⁶,⁵⁷ A careful observation of Figure 4 unveils a higher contrast in two atoms of the image. The line profiles of these atoms are depicted in the inset of Figure 4a. As expected, N atoms (Z = 7) show a higher contrast than C atoms (Z = 6) in the ADF-STEM images. Moreover, the relative ADF-STEM intensity between the N and the C atoms (4:3) is in very good agreement with previously reported work.⁵⁶ The N atoms observed in the micrographs are clearly in substitutional configuration (see Figure S5 in the Supporting Information).

We keep in mind that the main focus has been to work toward a one-step method that enables us to obtain a high-quality N-graphene that can be used directly after the molecular transformation. Previous studies have shown that the presence of oxygen on copper can significantly influence the final quality of graphene and this is often related to the oxygen concentration during synthesis.⁵⁸–⁶² Our on-surface method offers a very controlled alternative to explore the advantage of using an oxide instead of a metallic surface to produce N-graphene. More specifically to the synthesis, after C₁₆H₆N₆ deposition, the samples were exposed to air for 5 min and then placed again under UHV conditions. This hints that the substrate plays a key role for a controllable adsorption of the molecular species and also during the on-surface synthesis.⁶₃,⁶₄ The 5 min exposure to air is a good compromise between quality and decoupling of graphene from its substrate, which represents a significant advancement to the state-of-the-art on N-graphene synthesis. The analysis corresponding to this material is shown in Figure 5. The panel a shows the C 1s spectrum of the N-graphene on the oxidized Cu(111) with its corresponding deconvolution. Compared to graphene on Cu(111) (Figure 3c), a shift toward lower binding energies (0.1 eV) is observed, which is the first hint toward a low...
Figure 5. (a) XPS C 1s deconvoluted core level spectrum recorded on N-doped graphene grown on oxidized Cu(111). The inset shows the N 1s core level spectrum with the corresponding deconvolution. (b) Experimental LEED pattern of polycrystalline graphene on oxidized Cu(111) acquired at 94.2 eV at nearly normal incidence. The dashed curved arrow indicates the position of the graphene ring. (c) Simulated LEED pattern. The visible spots in the experimental pattern have been highlighted. The white arrows point at one of the Cu(111) and Cu2O spots. The ARPES image of N-graphene on oxidized Cu(111) (d) and Cu(111) (e), acquired with a photon energy of 40.8 eV (He II) along the ΓK direction. The white dotted line shows the position of the Fermi level.

interaction of the N-graphene with the substrate. This has been reported for graphene on oxidized Cu(111) grown from CVD. Further looking at the C 1s, the response corresponding to N-graphene on crystalline Cu(111), in Figure 3c, is very different from the spectrum of N-graphene on oxidized Cu(111), which has additional components. Besides the peak at 284.3 eV corresponding to sp2-hybridized C atoms on oxidized Cu(111), the minor components at 284.8 and 285.4 eV reflect the presence of carbon in another binding environment, which might be attributed to C–OH and C–O bonds, justified by the formation of molecular species on the surface with O and H atoms that are present after the feedstock’s decomposition. Furthermore, the LEED diffraction pattern in Figure 5b obtained after annealing the C16H6N6 at 1250 K deposited on oxidized Cu(111) shows well-defined spots, suggesting a long-range order over extended zones of the surface. To disambiguate the interpretation of this diffraction pattern and gain deeper insight into the surface structure, we have performed LEED simulations using LEEDpat software (Figure 5c). With this, it was possible to associate the LEED spots to a surface reconstruction described by the matrix \( \begin{pmatrix} 3 & 2 \\ -1 & 2 \end{pmatrix} \) and its three equivalent rotational domains. This surface reconstruction has been interpreted as a distorted Cu(100)-(\( \sqrt{3} \times \sqrt{3} \))R45°-O layer on the Cu(111) substrate. Note that a careful observation of Figure 5b unveils a circular ring typical of polycrystalline graphene just above the 1 × 1 diffraction spots of Cu(111). For the sake of comparison, the graphene grown on the clean Cu(111) was inspected. Also, in this case, the diffraction pattern corresponds to the characteristic ring from polycrystalline graphene combined with the first-order diffraction of Cu(111) (Figure S6 in the Supporting Information).

To further analyze the electronic performance of the N-graphene on oxidized Cu(111), ARPES measurements where performed with a photon energy of 40.8 eV (He II source). The fingerprint of graphene in ARPES on Cu and other metals is well established in the literature. Graphene on Cu(111) shows a Dirac point residing at 0.4 eV below the Fermi level (corresponding to n-type doping). Our N-graphene sample grown on Cu(111) shows the Dirac point residing at 0.43 eV below the Fermi level (Figure 5e). Only for graphene decoupled from the substrate, the intrinsic Dirac cone of pristine graphene can be retained, and this has been shown in the literature with methods such as decoupling by gold intercalation under graphene grown on nickel,77 the growth of polycrystalline samples finding decoupled flakes by microARPES,78 and by actually oxidizing Cu prior to the growth,34 where an additional flat band is observed at \( \sim 0.8 \) eV below the \( E_F \) which originates from the hybridization of the Cu4s and the O2p states. In Figure 5d, the linear dispersion of the \( \pi \) and \( \pi^* \) bands at the vicinity of the Dirac point along the \( \Gamma K \) direction in the Brillouin zone is observed. Note that the Dirac point here is around 0.1 eV below \( E_F \) which is associated to the charge transfer that arises from the N-substitution. We expect distribution of the dopants to be random but it is mainly substitutional, differing from previous studies that correlate the local densities around N impurities versus the spectral densities given by ARPES79 in samples where a larger N concentration can be attained but not necessarily exclusively on direct substitutional fashion. Moreover, this confirms that the presence of copper oxide favors the decoupling of N-graphene from its metallic substrate.

### CONCLUSIONS

In summary, we have proved the feasibility of on-surface synthesis of N-graphene using C16H6N6 as the molecular precursor, which has allowed obtaining a material with nearly exclusive substitutional N, which is in turn decoupled from its...
catalytic surface. Our method has been proven for single-crystalline Cu(111) and polycrystalline copper foils, which represent a technologically relevant low-cost option. We have shown that it is possible to grow N-graphene with \( ~1.4 - 2 \text{ at } \% \) N in substitutional configuration. Further studies on the tunability of this doping level would enhance the applicability of doped graphene grown by this method even more. Our ARUPS measurements show that N-graphene is decoupled from its substrate, opening encouraging prospects for its direct growth on other relatively inexpensive high-\( \varepsilon \)-dielectric materials.

**ASSOCIATED CONTENT**

Supporting Information
The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.jpcc.0c06415.

Molecular structure of the dicyanopyrazobenzothiadiazole molecule; C 1s XPS spectra for mono- and multilayer coverages; C 1s XPS spectra as a function of postdeposition annealing temperature; C 1s and N 1s XPS spectra after annealing at 1250 K; medium-angle ADF-STEM images of N-graphene; and LEED of N-graphene on metallic and oxidized Cu(111) (PDF)

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**Notes**

The authors declare no competing financial interest.

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