

# Bottom-up solution synthesis of narrow nitrogen-doped graphene nanoribbons†

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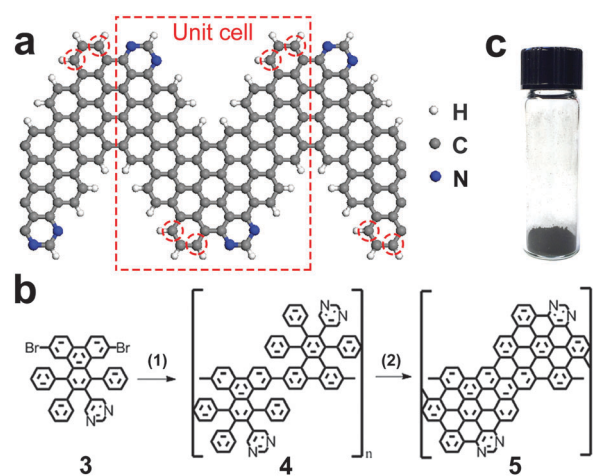
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**Large quantities of narrow graphene nanoribbons with edge-incorporated nitrogen atoms can be synthesized via Yamamoto coupling of molecular precursors containing nitrogen atoms followed by cyclodehydrogenation using Scholl reaction.**

Graphene nanoribbons (GNRs), narrow strips of graphene with high aspect ratios, attract a great deal of attention because of their interesting electronic and magnetic properties.<sup>1</sup> According to the theoretical studies, these properties strongly depend on the width of GNRs as well as their edge structure and termination, and are also very sensitive to the edge disorder.<sup>2–5</sup> Thus, it is very important to precisely control the structure of GNRs on the atomic scale. Numerous top-down fabrication approaches have been developed to produce GNRs from graphite, graphene or carbon nanotubes, as discussed in the recent review papers,<sup>6–8</sup> but the resulting ribbons typically have relatively large and non-uniform widths and disordered edges. Narrow atomically precise GNRs could be synthesized via bottom-up approaches that employ decomposition of molecules inside carbon nanotubes,<sup>9,10</sup> or rely on the surface-assisted<sup>11–14</sup> or solution polymerization<sup>8,15–22</sup> of molecular precursors followed by cyclodehydrogenation of the resulting polymers. While surface-assisted bottom-up synthesis of GNRs yields small quantities of ribbons on metallic substrates in ultrahigh vacuum conditions that are ideal for scanning tunnelling microscopy (STM) characterization, solution-based approaches are preferred for the bulk synthesis of GNRs, so these two groups of methods could be considered as complimentary.

Electronic properties of GNRs could be further tuned via their doping with heteroatoms, such as boron or nitrogen.<sup>23</sup> This possibility has been extensively studied theoretically,<sup>24–30</sup> but only a few experimental attempts to synthesize nitrogen-doped GNRs (N-GNRs) by bottom-up approaches have been reported.<sup>14,20</sup> Bronner *et al.* prepared chevron-like N-GNRs by the surface-assisted approach on Au(111) substrate and performed their spectroscopic characterization.<sup>14</sup> Kim *et al.* synthesized N-GNRs with a different structure in solution via Suzuki coupling of molecular precursors, one of which was dibrominated pyrazine, and cyclodehydrogenation of the resulting polymers.<sup>20</sup> Here we demonstrate that high-quality chevron-like N-GNRs could be synthesized via Yamamoto coupling of molecular precursors containing nitrogen atoms followed by cyclodehydrogenation via Scholl reaction (Fig. 1).

The structure of the N-GNR synthesized in this work is shown in Fig. 1a. Since the unit cell of this ribbon contains four nitrogen atoms, we refer to it as “4N-GNR”. The monomer molecule 3 used as a precursor for the 4N-GNR synthesis is not symmetric relative to the plane perpendicular to the direction



**Fig. 1** Synthesis of 4N-GNRs. (a) Schematic of 4N-GNR. (b) Reaction scheme used in this work: (1) Yamamoto coupling, (2) Scholl reaction; see text and ESI† for details. (c) Optical photograph of a 4N-GNR powder.

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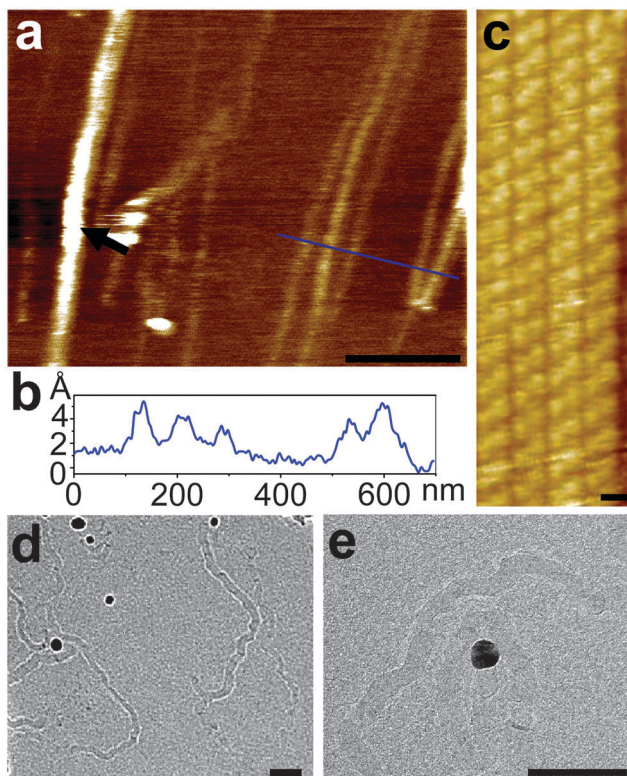
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of coupling (Fig. 1b), so as the result, the positions of nitrogen atoms in the ribbon edge could vary, *i.e.* the nitrogens could also occupy the positions highlighted with red circles in Fig. 1a instead of the ones shown in the schematic. However, the N:C ratio of 1:20 in this ribbon should be the same regardless of how the monomer molecules couple with each other.

The synthesis and characterization of the precursor molecule **3**, as well as the reactions used to convert monomer **3** to polymer **4** to 4N-GNR **5** (Fig. 1b) are described in detail in the ESI.† About 100 mg of 4N-GNRs was obtained in a single reaction (Fig. 1c). Previously, we have demonstrated that a similar procedure could be used to obtain >1 g of chevron-type GNRs (nitrogen-free analogues of 4N-GNRs shown in Fig. 1a) in a single reaction,<sup>21</sup> so we believe that the synthesis of 4N-GNRs could also be scaled up to at least a gram scale. Though not soluble in any solvents that we tried, these ribbons could be suspended by sonication in, for example, toluene or mesitylene, and then deposited on different substrates for microscopic and spectroscopic characterization.

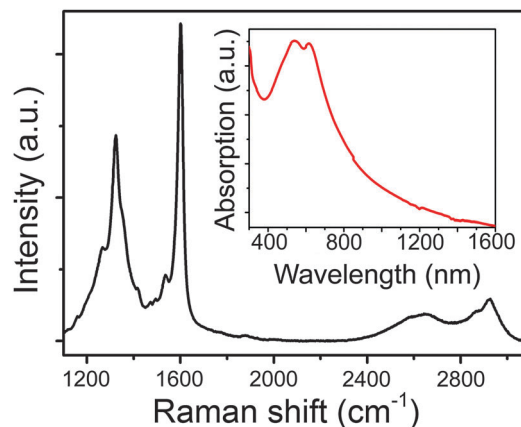
It was previously demonstrated that when chevron-type GNRs are deposited from dispersion to a substrate, they could be found in a form of elongated self-assembled nanostructures, referred to as “GNR nanobelts”.<sup>21</sup> These nanobelts are only one carbon atom thick and comprise multiple GNRs arranged in a side-by-side fashion.<sup>21</sup> We deposited 4N-GNRs on a variety of substrates (mica, Au(111), copper TEM grid with ultrathin carbon support film) and observed similar nanobelts. Fig. 2a shows an atomic force microscopy (AFM) image of 4N-GNRs deposited on mica.



**Fig. 2** Microscopic characterization of 4N-GNRs. (a) AFM image of 4N-GNRs deposited on mica. Scale bar is 400 nm. (b) Height profile along the blue line shown in (a). (c) STM image of 4N-GNRs deposited on a Au(111) single crystal. Scale bar is 1 nm. (d, e) TEM images of 4N-GNRs. Scale bars are 50 nm.

image of 4N-GNRs that were dispersed in toluene by sonication, deposited on freshly cleaved mica and dried prior to imaging. Observed in this image are high-aspect-ratio nanobelts that are several  $\mu\text{m}$  long. The alignment of 4N-GNR nanobelts likely happens in the contact angle between a droplet of 4N-GNR dispersion and a substrate during the solvent drying. As shown in the height profile measured along the blue line in Fig. 2a, the majority of these nanobelts have heights  $< 5 \text{ \AA}$ , which corresponds to the thickness of a single layer or graphene (Fig. 2b). In fact, in multiple AFM images we observed only one structure with a thickness ( $t$ ) larger than  $5 \text{ \AA}$ , which is indicated by the black arrow in Fig. 2a. This structure ( $t \sim 1 \text{ nm}$ ) is likely a rare stack of 2–3 nanobelts. Fig. 2c shows an STM image of the fragment of a 4N-GNR nanobelt on Au(111), which illustrates the side-by-side arrangement of individual ribbons that are over 40 nm long. As we indicated in the previous study of solution-synthesized chevron-like GNRs, it remains unclear if these nanobelts exist in solution or form directly on a substrate by capillary forces during the solvent evaporation.<sup>21</sup> 4N-GNR nanobelts were also imaged by transmission electron microscopy (TEM), see Fig. 2d and e. The nanobelts found on the TEM grids are 8 to 20 nm wide, which corresponds to 8–12 4N-GNRs arranged in a side-by-side fashion, and up to  $1 \mu\text{m}$  long.

Raman spectroscopy is very sensitive to the disorder (edges and other structural defects, chemical functionalization, *etc.*) in carbon materials,<sup>31</sup> so we used this method to confirm the high structural quality of 4N-GNRs. Fig. 3 shows the Raman spectrum of 4N-GNRs, where in addition to the most intense lines at  $1324$  and  $1600 \text{ cm}^{-1}$  that are typically referred to as D and G bands, respectively,<sup>31</sup> a series of smaller peaks around the D and G lines could be observed. This fine structure is a direct result of the low dimensionality of a nanoribbon, and the relative intensities and positions of the lines are characteristic for each specific polycyclic aromatic hydrocarbon (PAH).<sup>32</sup> 4N-GNRs synthesized in this work are structurally close to chevron-like GNRs, so this is not surprising that the Raman spectrum of 4N-GNRs is very similar to that of chevron-like GNRs synthesized by the solution approach.<sup>21</sup> Since disordered carbon materials usually show only broad D and G lines without any fine features, the spectrum shown in Fig. 3 indicates the high structural quality of 4N-GNRs.



**Fig. 3** Raman spectrum of 4N-GNRs. The inset shows UV-vis-NIR absorption spectrum of 4N-GNRs suspended in mesitylene by sonication.



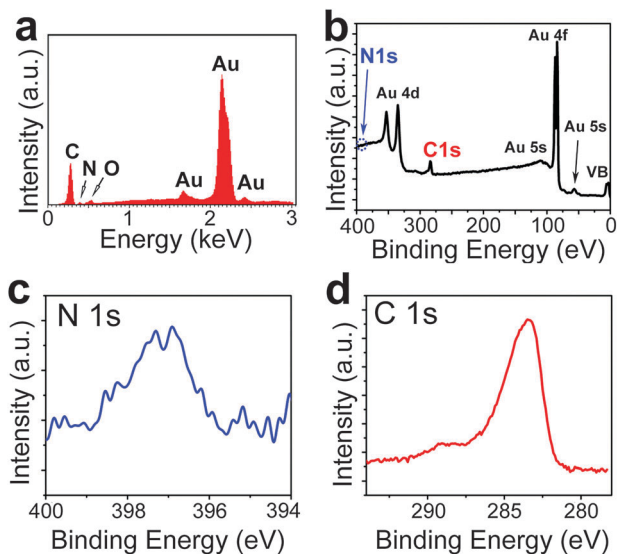


Fig. 4 Spectroscopic characterization of 4N-GNRs. (a) EDX spectrum of 4N-GNRs deposited on a gold substrate. (b–d) XPS spectra of 4N-GNRs deposited on a gold substrate: (b) survey spectrum, (c) N 1s spectrum and (d) C 1s spectrum.

The inset in Fig. 3 shows a UV-vis-NIR absorption spectrum of 4N-GNRs that were suspended in mesitylene by sonication. This spectrum is qualitatively similar to the UV-vis-NIR spectra of other solution-synthesized GNRs, showing a strong absorption in the visible range that gradually decreases with the wavelength increasing toward the infrared region.<sup>19,21</sup> The optical band gap of 4N-GNRs determined from this spectrum is  $\sim 1.6$  eV.

In order to confirm the presence of nitrogen atoms in the structure of 4N-GNRs we characterized the ribbons by energy-dispersive X-ray (EDX) spectroscopy and X-ray photoelectron spectroscopy (XPS). Fig. 4a shows an EDX spectrum of 4N-GNRs deposited on a gold foil. The nitrogen peak is detectable although small, which is not surprising, considering the expected N:C ratio of 1:20. The only foreign peak observed in the spectrum is the low intensity O line, which is likely caused by atmospheric adsorbates. The XPS survey spectrum (Fig. 4b) of the same sample demonstrates only the peaks associated with the GNRs and the gold substrate. XPS N 1s spectrum of 4N-GNRs (Fig. 4c) shows a single peak at 397.1 eV, which is close to pyridinic nitrogen peaks in N-doped graphene.<sup>33</sup> XPS C 1s spectrum (Fig. 4d) contains the main peak corresponding to the C–C bonds and the weaker peak at  $\sim 288.5$  eV likely representing C–N bonds.<sup>33,34</sup>

In summary, we demonstrate that the synthetic approach based on Yamamoto coupling of N-doped molecular precursors and cyclodehydrogenation *via* Scholl reaction could be used to synthesize high-quality 4N-GNRs. The ribbons were investigated by the combination of microscopic (STM, AFM, TEM) and spectroscopic (UV-vis-NIR, XPS, EDX and Raman spectroscopy) techniques. In the future studies, we will investigate if this approach could also be employed for the synthesis of N-GNRs with other structures and N:C ratios.

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