N-doped graphene for electrocatalytic O$_2$ and CO$_2$ reduction

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The electrocatalytic CO$_2$ reduction reaction (CO$_2$RR) and oxygen reduction reaction (ORR) are important approaches to realize energy conversion and sustainable development. However, sluggish reaction kinetics severely hinders the practical application of devices related to these reactions. N-doped graphene (NG) with unique properties exhibits great potential in catalyzing the CO$_2$RR and ORR, which is attributed to the electron redistribution. In this review, we start from the fundamental properties of NG, especially emphasizing the changes caused by N doping. Then the synthetic methods are summarized by classifying them into top-down strategies and bottom-up strategies. Subsequently, the applications of NG in the ORR and CO$_2$RR are discussed and the effects of electronic structure on the electrocatalytic activity are highlighted. Finally, we give our own perspective on the future research direction of NG in the applications of the ORR and CO$_2$RR.

1. Introduction

To mitigate the global climate problem and achieve carbon neutrality, the conversion and storage of renewable energy highly demand advanced electrochemical technologies, most of which involve various electrocatalytic reactions. For instance, the oxygen reduction reaction (ORR) is a critical electrochemical process in metal-air batteries and fuel cells.$^{1-4}$ The electrocatalytic CO$_2$ reduction reaction (CO$_2$RR) is especially an effective strategy to re-balance the carbon cycle and to harvest value-added chemical products.$^{5-7}$ However, these electrocatalytic processes have been severely limited by the sluggish reaction kinetics, which necessitate efficient electrocatalysts with good selectivity and durability. Although the state-of-art noble metal-based electrocatalysts have been used in these reactions,$^{8,9}$ the scarcity accompanied by high cost hampers the large-scale applications of these sustainable energy technologies. It is highly desirable to discover and design cost-effective electrocatalysts.

Among many non-precious-metal materials, graphene consisting of sp$^2$-hybridized carbon atoms arranged in a honeycomb lattice has been intensively studied as a potential candidate,$^{10}$ due to the outstanding electronic conductivity, large specific surface area, and good mechanical properties arising from the unique two-dimensional (2D) structure.$^{11,12}$ As a spark, it also inspires the exploration of other 2D materials and broadens the 2D material family, including transition dichalcogenides (TMDs), MXenes, hexagonal boron nitride (h-BN), etc.$^{11-16}$ Graphene can be built into different nanostructures, such as graphene quantum dots,$^{17}$ graphene nanoribbons,$^{18}$ and graphene nanomeshes$^{19}$ as well as recently emerged “magic-angle graphene,”$^{20}$ demonstrating various novel properties,$^{21}$ which attract the intensive interest of researchers around the world. Moreover, heteroatom doping into graphene also plays an important role in the modulation of fundamental properties.$^{22}$ Among various heteroatoms, the doping of nitrogen (N) atoms usually leads to strong charge polarization in the carbon plane due to the remarkable difference of electronegativity between nitrogen ($\chi = 3.04$ on the Pauling scale) and carbon ($\chi = 2.55$),$^{23}$

Since the pioneering work of N-doped carbon nanotubes used for the electrocatalytic ORR,$^{24}$ N-doped carbon materials gained special attention for electrocatalysis applications.$^{25}$ In particular, N-doped graphene (NG) has been placed at the center of the spotlight, due to the unique characteristics and advantages of the structure and properties, including the induced charge redistribution and defects by N doping.$^{26}$ Moreover, NG has been extensively explored as an electrocatalyst for the CO$_2$RR to play an ever-increasingly important role in realizing carbon neutrality and ecosystem protection.$^{27-29}$
Therefore, it is of significance to review the recent advances of NG applications in these reactions and to figure out the opportunities and challenges in the future.

In this review, we primarily start from the fundamental properties of NG followed by the synthesis methods. Then, we summarize the applications of NG in the ORR and CO$_2$RR and emphasize the effects of the electronic structure on the electrocatalytic activity. Based on the specific reaction pathway, we try to construct the correlation between the physicochemical properties and electrocatalytic activity and to elucidate the electrocatalytic reaction mechanism. Finally, we will provide an outlook on the future research direction and give our own perspective on NG in the applications of the ORR and CO$_2$RR.

2. Fundamental properties

Graphene is an sp$^2$-hybridized carbon layer with zero bandgap. Heteroatom doping into graphene, especially N doping, can effectively tailor its electronic structure and other intrinsic properties, broadening the application fields.$^{39}$ There are usually several N species in N-doped graphene (NG), including graphitic N, pyridinic N, pyrrolic N, amide N and N oxides as shown in Fig. 1a. The successful doping of N atoms into the graphene lattice can be revealed by the scanning tunneling microscopy (STM) image (Fig. 1b) and simulated STM image computed from the local density of states for graphitic doping (Fig. 1c).$^{31}$ The introduction of an N dopant at different locations leads to different electronic states. For example, the formation of the lone pair at pyridinic N makes the N atom negatively charged (Fig. 1d).$^{32}$ And the spatial propagation of the localized $\pi$ states can be seen in the STM simulations at a negative bias of $-1.0$ V, as shown in Fig. 1e and f.$^{32}$ The charge transfer caused by N doping induces net positive charge at adjacent C atoms, which facilitates the attraction of electrons from the anode and the ORR.$^{33}$ The charge redistribution also changes the chemisorption mode of the O$_2$ molecule from the usual end-on adsorption (Pauling model) to side-on adsorption (Yeager model),$^{24}$ which could effectively weaken the O–O bonding in parallel diatomic adsorption. With the development of synthetic methods and advanced characterization techniques, the understanding on the fundamental properties of NG has been deepened gradually.

The band structure and density of states (DOS) calculated by the ab initio method and tight-binding (TB) model indicate an obvious bandgap opening, as shown in Fig. 2a & b.$^{34}$ When different concentrations of N dopants are randomly distributed in one sublattice, the bandgap ($E_g$) ranges from 45, 110, 340 to 550 meV for various N concentrations corresponding to 0.5, 1, 4, and 8 at% (Fig. 2b).$^{34}$ Experimentally, compared to that of pristine graphene, the Raman spectrum of NG shows an obvious D$^*$-band, which is attributed to the intravalley double resonance scattering processes (Fig. 2c).$^{35}$ And a more intense D-band is also shown, which probably originates from the elastically scattered photo-excited electron created by the large number of N atoms embedded in the graphene lattice before emitting a phonon. There is also an obvious downshift

![Fig. 1](image_url) (a) Scheme of NG with different N species. (b) STM image of monolayer NG and (c) its simulation by DFT calculations. (d) Equilibrium geometry of pyridinic N calculated by DFT (left panel) and schematic representation of $p_z$ orbitals in pyridinic N (right panel). (e) Simulated STM images (constant-current mode) for pyridinic N graphene at a bias voltage of $-1.0$ V. The N atom is placed at the center of the image. (f) DFT equilibrium geometry and isosurface plot of electron density at $1.5 \times 10^{-3}$ electrons/Å$^3$ integrated from the Fermi level to $-0.7$ eV. (b and c) Reprinted with permission from ref. 31, copyright © 2011 AAAS. (d–f) Reprinted from ref. 29 with permission from ref. 32, copyright © American Physical Society 2012.
of the G-band and 2D band caused by the movement of the Fermi level.

Different N configurations in NG have different influences on the electrocatalytic performance. Theoretical studies reveal that the doped N located at the zigzag edge of graphene can activate the neighboring edge carbons toward the ORR, while for a given zigzag edge, the presence of pyridinic N significantly reduces the inherent ORR activity of graphene.\textsuperscript{36,37} Moreover, Wang et al. studied the nitrogen configuration and its location (armchair or zigzag edge) in NG by using theoretical calculations and presented the K-edge X-ray absorption spectra (XAS) and emission spectra (XES) as well as the final (initial)-state wave functions (Fig. 2d).\textsuperscript{38} Experimentally, the characteristic peaks can be measured and deconvoluted by X-ray photoelectron spectroscopy (XPS) and X-ray absorption near-edge structure spectroscopy (XANES) as shown in Fig. 2e and f. The high-resolution XPS spectrum of N1s can be fitted into four peaks at around 397.7, 399.2, 400.5 and 402.5 eV, which are assigned to pyridinic N, pyrrolic N, graphitic N and oxygenated N, respectively (Fig. 2e).\textsuperscript{39} As shown in Fig. 2f, the N K-edge XANES spectrum of NG changes with the annealing temperature from 200 to 600 °C, indicating the corresponding variation of N species.\textsuperscript{40} Especially, the N5 peak at ~406 eV in GPA-200 is ascribed to the $\sigma^*$ excitation of C-N bonds, while N4 at ~402 eV can be attributed to the attachment of urea species. Moreover, amino induced small features at about 399 eV (N1 and N2) can also be observed. With the increase of temperature, the N4 peak gradually decreases and a strong N2 peak emerges. When the temperature reaches 500 °C, the N1 peak and N3 peak ascribed to the pyridinic type and graphitic type, respectively, are observed. And the graphitic N species become dominant with the N3 peak increasing greatly at 600 °C. These results provide significant guidance for the identification of N species in NG experimentally synthesized by different methods.

3. Synthetic methods

Generally, the synthesis of graphene can be classified into two main strategies: top-down and bottom-up methods.\textsuperscript{41,42} Top-down strategies refer to cutting or tearing graphite into graphene, including mechanical exfoliation, chemical ablation, electrochemical oxidation, plasma treatment, etc. Bottom-up methods realize the synthesis of graphene from molecular precursors containing a certain number of conjugated carbon atoms or from the gas-phase vapor growth. To realize the substitution or doping of N atoms into the graphene lattice, several treatment methods are usually employed.

3.1 Top-down strategies

3.1.1 Thermal annealing. Thermal annealing is a simple method to prepare NG by heat-treating a mixture of graphene/graphene oxide and N-containing sources. For example, Dai's group developed N-doped reduced graphene oxide (rGO) via thermal annealing of GO in an NH\textsubscript{3} atmosphere, which can occur at a temperature as low as 300 °C.\textsuperscript{43} They also found that the doping process is accompanied by the reduction of GO,
where the oxygen level decreases from ~28% in GO down to ~2% upon thermal annealing with NH$_3$ gas at 1100 °C. Oxygen groups in GO were found to be responsible for reactions with NH$_3$ and C–N bond formation. The sample with the highest N-doping level was annealed at 500 °C and the N level decreases with the increase of temperature. Besides NH$_3$ gas, other N-containing sources were utilized to react with GO to form NG at an elevated temperature, such as urea, polypyrrole, and melamine. The N content in NG can reach as high as about 15 at%.$^{44}$

3.1.2 Hydrothermal method. Different from thermal annealing, the hydrothermal method is another facile way to obtain NG by heat-treating a mixture of GO and N-containing precursors in solution at different temperatures.$^{45}$ N-containing precursors include polypyrrole, NH$_4$HCO$_3$, NH$_4$H$_2$O, urea, etc. In a typical process, GO dispersed in solution is mixed with a certain amount of N-containing precursor, such as urea, under sonication. Afterwards the solution was sealed in a teflon-lined autoclave and maintained at an elevated temperature, e.g., 180 °C for hours. After cooling to room temperature, the final product is washed with distilled water and ethanol, and dried at about 60 °C.

3.1.3 Microwave method. Microwave reduction is a rapid method to reduce GO to pristine graphene via one- to two-second pulses of microwaves,$^{46,47}$ because GO can absorb microwaves easily and reach a very high temperature (500 °C) in a few seconds. Pure graphene is a dielectric material with an sp2 bonded carbon network, which contains abundant delocalized π electrons. In an external electric field, these electrons start moving in the direction of the field and generate an electric current. As these charge carriers cannot keep up with the change of the phase in an alternating field, the produced energy is dissipated in the form of heat.$^{48,49}$ Both GO and rGO are polar materials due to the presence of oxygen functional groups. In the presence of an electromagnetic field, the alternating electric field forces the dipoles to align themselves in phase with the field, thus generating stretching and bending vibrations, which subsequently are dissipated in the form of heat energy.$^{50}$ During the microwave-reduction process, N doping can be achieved in an NH$_3$ atmosphere. Typically, graphite oxide is placed in a three-neck flask purged with NH$_3$ at a certain rate for several minutes through a mass flow controller. Then the flask was placed in a microwave oven under high power for tens of seconds with the flow of NH$_3$ in the whole process. Finally, NG can be obtained through the microwave-thermal expansion procedure.$^{51}$ In addition, microwave-assisted hydrothermal heating is a combined method to obtain NG by mixing the GO and N sources in a microwave reactor.$^{52}$

3.1.4 Plasma treatment. Plasma treatment is an effective approach to modify the material surface, which can be used for the incorporation of heteroatoms in graphene. Plasma is a gas of charged particles, which can be generated by heating a gas or by applying strong electromagnetic fields.$^{53}$ These charged particles make the plasma electrically conductive. The N$_2$ ions from N$_2$ plasma initiated by a high bias voltage are able to generate defects and create covalent bonds with pre-existing defect sites.$^{44}$ For the typical synthesis of NG, graphene is firmly attached onto a glassy carbon electrode (GCE) and then was placed in a plasma chamber, which was backfilled with a nitrogen atmosphere at a specific pressure. The plasma power and treatment time can be tailored for a targeted composition or structure.$^{54}$ By using synchrotron-based scanning photoemission microscopy (SPEM), Choi and coworkers found that N doping takes place not only at the active edges but also at the basal plane (Fig. 3a).$^{55}$ Moreover, with the increase of treatment duration, at basal planes the pyrrole-N portion increases but the graphite-N portion decreases, and the pyrrole-N portion appears...
higher at edges than at basal planes. N configuration in graphene is highly affected by the vacancy defect species in graphene, because the plasma can stimulate various vacancies, such as divacancies and monovacancies.\textsuperscript{37} Moreover, the thermal spike caused by prolonging implantation time or increasing implantation energy can lead to the conversion from pyrrolic-N to pyridinic-N.

3.1.5 Arc-discharge method. Arc discharge between two graphite electrodes under a relatively high pressure of hydrogen can yield graphene flakes in a water-cooled stainless steel chamber.\textsuperscript{38} When \( \text{NH}_3 \) is used as one of the buffer gases, the generated graphene sheets are doped by N atoms on a large scale, which could be applied to successive production. In a typical process, two electrodes were both pure graphite rods with a diameter of 8 mm, and the current was held at 120 A.\textsuperscript{39} As the rods were brought close enough, discharge occurred and plasma formed. The anode was vaporized in a He and \( \text{NH}_3 \) mixed atmosphere of 760 Torr. To keep a constant distance of 1 mm between the two electrodes, the cathode was kept rotating with the reduction of the anode.

There are also other methods used for synthesizing NG, which are not so popular as the abovementioned methods, for example, supercritical reaction (Figure 3b),\textsuperscript{60} pulsed laser irradiation (Fig. 3c),\textsuperscript{61} and flame treatment (Fig. 3d).\textsuperscript{62} The supercritical reaction is readily employed to prepare NG at a relatively low temperature of about 310 °C. In a typical procedure, few-layer graphene sheets (2 mg) were dispersed in acetonitrile (25 mL), and the mixture was transferred to a corundum-lined autoclave (50 mL).\textsuperscript{60} The autoclave was heated to 310 °C and an environment of supercritical reaction fluid was maintained within the autoclave. After a certain reaction time, N-doped graphene sheets can be harvested after centrifugation and rinsing. Laser irradiation can also provide enough heat for direct \textit{in situ} growth of NG on an insulating substrate and does not need a complicated transfer process. By virtue of flame treatment, Pan’s group prepared NG with dominant pyridine-N from amine plus ethanol flames.\textsuperscript{62} During the process, a Si substrate with a Ni film was inserted into the flame at 5.5 cm above a lamp for 1 min. A brown combustion product, \textit{i.e.}, NG, was observed on the substrate when it was withdrawn from the flame.

3.2 Bottom-up strategies

3.2.1 Chemical vapor deposition (CVD). The CVD method is a popular approach to obtain large-area and high-quality graphene and heteroatom-doped graphene. For the synthesis of NG, the C and N precursors are introduced onto the surface of a hot substrate (\textit{e.g.}, Cu or Ni) to realize the formation of C–C and C–N bond networks at a high temperature (>800 °C).\textsuperscript{63,64} In the synthesis process, Cu or Ni not only serves as the substrate, but also functions as a catalyst to tune the growth of graphene. To lower the temperature, plasma-enhanced CVD (PECVD) and thermal CVD were also employed to grow NG (Fig. 4a).\textsuperscript{65,66} During the CVD process, generally, methane (\( \text{CH}_4 \)) is used as a carbon source, while \( \text{NH}_3 \) is adopted as the N source. Besides gas-phase precursors, liquid or solid organic molecules are also employed as precursors, such as pyridine,\textsuperscript{60} acrylonitrile,\textsuperscript{67} hexaphenylborazine\textsuperscript{68} and pentachloropyridine.\textsuperscript{69} The structure and quality of the resulting NG films were correlated with the functional groups of the precursor molecules and gas phase composition. For example, amine functional groups in methylamine can lead to a direct growth of graphene even without additional H\(_2\) in the gas phase, which indicates that the intermediate species (\textit{e.g.}, amine radicals) in the gas phase can significantly enhance the growth kinetics of graphene.\textsuperscript{70}

3.2.2 Solvothermal method. The Wurtz-type reductive coupling (WRC) reaction is also a bottom-up method for the preparation of high-quality graphene by a solvothermal reaction of tetrachloromethane (CCl\(_4\)) with metal potassium (K).\textsuperscript{71} Deng \textit{et al.} synthesized two types of NG by utilizing CCl\(_4\) or cyanoenic chloride as the carbon source and lithium nitride as the nitrogen source in a stainless steel autoclave filled with N\(_2\) gas at 250 ~ 350 °C for several hours (Fig. 4b).\textsuperscript{72} The products were washed sequentially with 18 wt% HCl aqueous solution, water, and ethanol, followed by drying at 120 °C for 12 h. Pentachloropyridine was also employed as the C and N source to react with K in an autoclave.\textsuperscript{72} Our group also used CCl\(_4\) as the carbon source and an appropriate amount of pyrrole as the nitrogen source to react with metallic K at 200 °C for 6 h and finally obtained NG with a certain N content.\textsuperscript{73,74} The doping level can be readily controlled by the amount of N-containing precursors for the synthesis.

Besides these two typical bottom-up methods, electrochemical bottom-up synthesis or bottom-up self-assembly was also developed to synthesize NG, strictly speaking, N-doped...
graphene quantum dots or graphene nanoribbons. Tian et al. used o-phenylenediamine as the reactant in pure water as the electrolyte and applied a constant voltage of 500 V between two high-purity Pt sheets for more than 1 hour at 25 °C at a stirring rate of 1000 rpm. The colorless solution gradually turned yellow indicating the formation of NG with a size from 2 to 6 nm. Sinitskii and coworkers synthesized high-quality chevron-like N-doped graphene nanoribbons via Yamamoto coupling of molecular precursors containing nitrogen atoms followed by cyclodehydrogenation via the Scholl reaction.

4. Electrocatalysis applications

4.1 Oxygen reduction reaction

The oxygen reduction reaction (ORR) is a critical reaction occurring at the cathode of fuel cells and metal-air batteries. Due to the sluggish reaction kinetics, precious Pt-based electrocatalysts are commonly used to boost the reaction, which usually accounts for almost half of the cost of fuel cells. To replace the commercial Pt-based electrocatalysts, many noble-metal-free materials have been developed, among which carbon-based materials are promising alternatives, especially N-doped carbon materials. As one of the N-doped carbon materials, NG not only holds the advantages of carbon materials (e.g., good electrical conductivity), but also possesses the unique features of graphene (e.g., high specific surface area and charge redistribution), which exhibits a highly promising prospect in the future.

The ORR is a multi-step reaction proceeding through either a direct 4e pathway or a two-step 2e pathway, which highly depends on the activity of the electrocatalyst. In an acidic medium, the possible reaction pathways are expressed as eqn (1)-(3), while those in alkaline solution are expressed as eqn (4)-(6).

In an acidic medium,

4e pathway: \( \text{O}_2 + 4\text{H}^+ + 4\text{e}^- \rightarrow 2\text{H}_2\text{O} \)  
2e pathway: \( \text{O}_2 + 2\text{H}^+ + 2\text{e}^- \rightarrow \text{H}_2\text{O}_2 \)  
\( \text{H}_2\text{O}_2 + 2\text{H}^+ + 2\text{e}^- \rightarrow \text{H}_2\text{O} \)

In an alkaline medium,

4e pathway: \( \text{O}_2 + 2\text{H}_2\text{O} + 4\text{e}^- \rightarrow 4\text{OH}^- \)  
2e pathway: \( \text{O}_2 + \text{H}_2\text{O} + 2\text{e}^- \rightarrow \text{HO}_2^- + \text{OH}^- \)  
\( \text{HO}_2^- + \text{H}_2\text{O} + 2\text{e}^- \rightarrow 3\text{OH}^- \)

A good selectivity of NG for the 4e reaction pathway is preferred in the application of fuel cells and metal-air batteries. For the reaction pathway in an alkaline medium, there are two

Fig. 5 (a) ORR pathways on NG in alkaline solution, where (I) presents an associative mechanism and (II) is a dissociative mechanism. (b) Free energy diagram for \( \text{O}_2 \) reduction on S1 and S2 under the conditions of 0.04 V and pH = 14. (c) Intermediate structures of the associative mechanism on S1 and S2. The gray, blue, red, and white spheres represent C, N, O, and H atoms, respectively. Reprinted with permission from ref. 80, Copyright © 2012 Elsevier Inc.
different mechanisms, i.e., associative mechanism and disso-
ciative mechanism, as shown in Fig. 5a. A density functional
theory (DFT) study reveals that the associative mechanism
(black line) on two surfaces (S1 and S2) of NG is more ener-
ggetically favorable than the dissociative one (blue line), as
shown in Fig. 5b and c. In S1 and S2, N atoms are separated by
two C atoms and three C atoms, respectively, so S1 possesses
a more concentrated N distribution than S2. The generation of
OOH$^-$ is represented by the red line. The results show that the
rate-determining step (RDS) in the associative mechanism is
the removal of O(ads) species from the active site. And the formation
energy of O(ads) on S1 is always lower (more exothermic) than
that on S2, which is attributed to the higher local concentration
of N atoms on S1 than that on S2.

The real active sites in NG have been an attractive research
topic due to contradictory opinions: carbon atoms adjacent to
the N dopant, graphite-N or pyridine-N, or both. For example,
Sidik and coworkers reported that substitutional N atoms from
the graphite sheet edges are more active than those close to the
edges. Niwa et al. also found that catalysts with more graphite
N exhibit a higher ORR activity than those with more pyridine
N. Huang et al. suggested that graphite N next to the zigzag
edge enhances the O$_2$ adsorption and the subsequent ORR

Fig. 7 CO$_2$RR at different standard potentials in aqueous solution at
pH = 7.0, 1.0 atm, and 25°C. Reprinted with permission from ref. 26,
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process to two $\text{H}_2\text{O}$ with a small activation barrier of about 5 kcal mol$^{-1}$, whereas the graphite $\text{N}$ inside the graphene sheet does not contribute to ORR activity. Lai et al. synthesized different NG samples as shown in Fig. 6a by using different $\text{N}$-containing precursors and found that the limiting diffusion current ($j_L$) and the onset potential ($E_{\text{onset}}$) are governed by graphitic-$\text{N}$ and pyridinic-$\text{N}$, respectively (Fig. 6b). Jung’s group found that graphitic $\text{N}$ in particular enables improved ORR activity and becomes pyridinic $\text{N}$ (stage IV) in the next transfer reaction of electrons and protons via the ring-opening of a cyclic $\text{C}$–$\text{N}$ bond as shown in step (3a) in Fig. 6c. And the projected density of states (pDOS) and the orbital shape of $\text{N}$ in stage (IV) are highly similar to those of typical pyridinic $\text{N}$. Recently, Guo et al. tactfully designed pyridinic $\text{N}$-dominated HOPG (pyri-HOPG) and graphitic $\text{N}$-dominated HOPG (graph-HOPG) as evidenced by Fig. 6d and proved that Lewis basic $\text{C}$ atoms neighboring pyridinic $\text{N}$ instead of pyridinic $\text{N}$ themselves are active sites toward the ORR as illustrated by the polarization curves in Fig. 6e.

Although there are unconsistent opinions on the function of different $\text{N}$ species, there is no doubt that the introduction of $\text{N}$ heteroatoms induces a positive charge density on an adjacent $\text{C}$ atom, due to the high electronegativity of $\text{N}$. The redistribution of electrons facilitates the adsorption and subsequent reduction process. Zhang et al. theoretically calculated the atomic charge density and spin density of pristine graphene and NG by constructing the models of $\text{C}_{46}\text{H}_{18}$ and $\text{C}_{45}\text{NH}_{18}$. They pointed out that $\text{N}$-doping would promote the electrons to be easily excited from the valence band to the conduction band, because $\text{N}$ doping introduces an unpaired electron and causes the localized distribution of molecule orbitals. Moreover, spin density is more important than atomic charge density in determining the catalytically active sites. Generally, carbon atoms with the highest spin density are the electrocatalytically active sites. These findings inspire the modulation of carbon-based materials toward a high-performance ORR, such as introducing other types of heteroatom or carbon defects to adjust the electronic structure or create more exposed active sites.
4.2 CO$_2$ reduction reaction

Carbon neutrality has become a global goal, which is desired to achieve net-zero carbon dioxide (CO$_2$) emissions. Enormous efforts have also been made by worldwide researchers. The electrochemical CO$_2$RR is an effective approach to realize the conversion from CO$_2$ to value-added chemical products.$^{87}$ The CO$_2$ molecule consisting of two linear C=O bonds is thermodynamically stable and chemically inert, due to a bond dissociation energy of 750 kJ mol$^{-1}$, which is difficult to reduce.$^{88}$ Moreover, the CO$_2$RR becomes more challenging in an aqueous electrolyte, due to the competition of the hydrogen evolution reaction (HER) and the diversity of products. Generally, the CO$_2$RR involves 2 to 18 electrons depending on different pathways as shown in Fig. 7. Therefore, electrocatalysts with high selectivity and activity toward CO$_2$ are necessary.

Inspired by the achievements of NG toward the ORR, researchers have also studied the electrocatalytic performance and mechanism of N-doped carbon materials. For example, Kumar et al. reported that N-doped carbon nanofibers (N-CNFs) are able to reduce CO$_2$ to CO at a low overpotential of 0.17 V vs. SHE and exhibit exceptionally high current densities (~13 times higher than that of bulk Ag) and a high FE of 98% for CO formation at −0.573 V vs. SHE.$^{89}$ They proposed that the naturally oxidized carbon atoms with positive charge and later reduced C atoms adjacent to pyridinic-N in N-CNFs were responsible for the CO$_2$RR. Sharma et al. also reported that the presence of graphitic and pyridinic nitrogen significantly decreases the overpotential and increases the selectivity towards the formation of CO.$^{90}$ Ajayan’s group conducted a series of experiments to improve the performance of the CO$_2$RR. Firstly, they synthesized N-doped CNTs (N-CNTs) and obtained good CO$_2$RR performance with an overpotential of −0.18 V and a selectivity of 80%.$^{91}$ They thought that pyridinic N as a catalytic site possesses a low free energy for CO$_2$ activation and a high barrier for hydrogen evolution. Moreover, by assembling NG to 3D foam, they improved the faradaic efficiency for CO production to ~85% at a lower overpotential.
(−0.47 V) and stability for at least 5 h.24 DFT calculations show
that the uphill free energy barrier comes from the first proton-
coupled electron transfer step to form COOH* and drops
dramatically upon introducing dangling N bonds (Fig. 8a & b).
Triple-pyridinic N maximally lowers the barrier of the COOH*
asorption step followed by single pyridinic and graphitic N.
Although COOH* adsorbing on pyrrolic N site is exergonic, the
release of chemisorbed CO from pyrrolic-N is endothermic with
an energy barrier of ~0.6 eV (Fig. 8a). To increase the density of
pyridinic N, they prepared nanometer-size N-doped graphene
quantum dots (NGQDs) as shown in Fig. 8c and d. The NGQDs
show a high total faradaic efficiency of up to 90% for the CO2RR,
and the selectivity for ethylene (C2H4) and ethanol (EtOH and
C2H5OH) conversions reaches 45% (Fig. 8e and f).31 The C2 and
C3 product distribution and production rate for the CO2RR are
well revealed with the development of computational power and
characterization techniques,109 such as in situ/operando Fourier
Transform Infrared Spectroscopy (FTIR) and in situ/operando X-ray
absorption spectroscopy (XAS).110,111 It is even necessary to estab-
lish an “in situ probing map” consisting of complementary aspects
of electrocatalysts to track the dynamic process and identify the
real active species systematically.102 Theoretically, machine
learning is a powerful strategy to almost exhaustively screen NG
with various N species and predict the desirable NG electro-
catalysts.111–116 The combination of high-throughput screening of
big data with purposeful experiments would open a new avenue
for the development of NG used as an electrocatalyst.

Besides being a good candidate for electrocatalysts, NG also
displays structural advantages to anchor single metal atoms due
to the intrinsic defects of NG, thus forming high-performance
single-atomic catalysts (SACs).107–109 For example, Sun’s group
fabricated single platinum atoms on NG nanosheets for
gas evolution by using the atomic layer deposition (ALD)
technique, resulting in the utilization of nearly all the Pt
atoms.118 Fei et al. synthesized a series of atomic 3d metals
embedded in N-doped holey graphene and investigated the
electrocatalytic performance toward the oxygen evolution reac-
tion.114 The synthesis of SACs by using NG or N-doped carbon
materials has become an important strategy.115 The flourishing
of SACs would move forward the understanding of the catalytic
mechanism and development of low-cost catalysts with high
performance toward various catalytic reactions not only limited to
the ORR and CO2RR.

5. Conclusion and outlook

In summary, we review the advances of NG with respect to
fundamental properties, synthesis and electrocatalytic applica-
tions toward the ORR and CO2RR. As a promising alternative
to noble meal catalysts, NG has exhibited promising potential to
use in these two energy-conversion reactions. Because of the
ideal 2D structure, it provides a good platform for exploring the
underlying electrocatalytic mechanism behind the superior
performance by both theoretical calculations and experiments.
But it is still challenging to design and synthesize targeted NG
with good selectivity and activity for the ORR or CO2RR.
Developing advanced synthetic methods is an important
precondition to guarantee a precise synthesis of NG with
specific N species. In this regard, a bottom-up strategy holds
a unique advantage, especially by coupling reaction between
organic molecules with specific N species.98,99

The complexity of N species in NG induces difficulty in
uncovering the underlying electrocatalytic mechanism. Solid
evidence on the electrocatalytic process and mechanism will be
well revealed with the development of computational power and
characterization techniques,109 such as in situ/operando Fourier
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of electrocatalysts to track the dynamic process and identify the
real active species systematically.102 Theoretically, machine
learning is a powerful strategy to almost exhaustively screen NG
with various N species and predict the desirable NG electro-
catalysts.111–116 The combination of high-throughput screening of
big data with purposeful experiments would open a new avenue
for the development of NG used as an electrocatalyst.

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