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## Solution-phase synthesis of graphene nanoribbons: a review on polymerization strategies

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A promising approach to opening the band gap of intrinsic graphene is through the synthesis of graphene nanoribbons (GNRs)—quasi-one-dimensional (1D) cutouts of the graphene sheet whose electronic and physical properties are governed by edge structure, chemical modifications, and backbone doping or pore integration. GNRs can be synthesized using both top-down and bottom-up approaches, with solution-phase synthesis emerging as a particularly attractive bottom-up method due to its advantages in diverse edge functionalization, possibilities in gram-scale production and characterization using various spectroscopy techniques, as well as exploration in practical applications. Solution-phase synthesis of GNRs typically involves a polymerization step followed by cyclodehydrogenation to form the final structure. This review provides an overview of the different polymerization strategies employed in GNRs solution-phase synthesis, highlighting their recent advancements and unique advantages. The Diels–Alder reaction enables the synthesis of GNRs with considerable lengths (up to 600 nm) and tunable edge functionalization. Suzuki polymerization facilitates the creation of porous GNRs while preserving heterostructures for post-synthetic modifications, whereas Yamamoto polymerization allows for the efficient production of laterally extended GNRs with varying bandgaps. Additionally, we discuss alternative cyclodehydrogenation methods, including photochemical approaches, for transforming polymer precursors into GNRs. By consolidating recent progress in solution-phase polymerization techniques, this review aims to provide valuable insights for the development of next-generation GNRs with tailored properties for electronic and optoelectronic applications.

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# 1. Introduction

Carbon, the sixth most abundant element in the universe, which comprises approximately 18% of human body weight, is remarkably versatile, existing in various allotropes such as diamond, graphite, graphene, carbon nanotubes, and fullerenes (Fig. 1).<sup>1</sup> In graphene, each carbon atom forms three Sigma ( $\sigma$ ) bonds, with the fourth electron contributing to a delocalized network of pi ( $\pi$ ) bonds. These  $\pi$  bonds arise from the overlap of perpendicular p-orbitals, creating a continuous electron cloud across the graphene sheet. This delocalized  $\pi$  system is responsible for graphene's exceptional electrical conductivity, high charge carrier mobility, and optical transparency.<sup>2,3</sup>

Graphene had been theoretically studied for over sixty years before its first isolation by Geim and Novoselov in 2004, *via* micromechanical cleavage<sup>4</sup> which has since opened possibilities in fundamental and applied research, notably spintronic and electronic applications.<sup>5</sup> However, owing to the intrinsic zero bandgap behaviour of graphene originating from the convergence of the Dirac cones at one point, graphene for semiconducting purposes poses a challenge. The solutions to open a bandgap in graphene span from dopant adsorption<sup>6</sup> to applying external electric fields<sup>7</sup> however to form reproducible structures with tuned bandgaps the structures and geometries of these graphene materials must be precisely engineered.<sup>8</sup>

Carbon-based nanostructures come in many forms yet share a similar quality which opens bandgaps compared to that of 2D graphene, structural confinement. These come in the form of graphene quantum dots (GQDs) and graphene nanoribbons (GNRs). The properties of both are mainly governed by their size and edge configurations.<sup>9</sup> GNRs show promising applications in the fields of spintronics, field-effect transistors (FETs), nanoelectronics and optoelectronics. They can be regarded as a quasi-one-dimensional cut-out of a graphene lattice.<sup>10</sup>

Depending on how the graphene lattice is cut, various edge structures and widths of these ribbons can be created (Fig. 2), each with distinctive properties depending on these qualities.<sup>9</sup>

GNRs synthesis can be split into two types: top-down and bottom-up methods.<sup>10</sup> Top-down methods include breaking down a graphene lattice to form the ribbon structure. These methods include nanofabrication,<sup>11,12</sup> sonochemical,<sup>13</sup> nanowire lithography,<sup>14,15</sup> unzipping carbon nanotubes (CNTs)<sup>16,17</sup> and various controlled nanocutting/etching methods.<sup>18–20</sup> Bottom-up synthesis uses precise molecular precursors and builds them up into the desired GNRs. These methods consist of on-surface synthesis (OSS), where metallic substrates are used as a platform to aid the synthesis of GNRs usually under ultra-high vacuum (UHV) conditions and in-solution synthesis which uses organic molecular precursors and various cross-coupling/polymerization synthetic techniques to afford polymer precursors, followed by a cyclodehydrogenation step to 'graphitize' the polymer precursors.<sup>10,21,22</sup>

While top-down methods of GNR synthesis have provided a variety of lengths and promising properties, such as high on-off ratios (up to  $10^6$ ), applicable for FET devices.<sup>23</sup> Most GNRs prepared by top-down methods possess undefined edge structures and little control over changing the edge structures on an atomic scale.<sup>10</sup> Bottom-up OSS of GNRs allows for conformation of theoretical predictions of GNRs by use of scanning probe techniques. However, a significant challenge in OSS of GNRs is the strong coupling between the GNRs and the metal substrate. Additionally, following synthesis, the GNRs remain adhered to the metal surface along with by products, making their removal and utilisation in devices challenging, thereby reducing practical application.<sup>24–29</sup> On the other hand, solution-phase synthesis of GNRs allows for gram-scalability and edge-functionalisation of GNRs opening doorways for unique electronic and magnetic properties.<sup>10,30,31</sup> In general, the route to solution-phase synthesis of GNRs follows two basic steps: (1) polymerization of rationally designed monomers and (2) planarization of the polymer precursors to afford the GNR (Fig. 3). The first step generally utilizes reactions such as Diels-Alder, Suzuki and Yamamoto polymerizations and the second step is an oxidative cyclodehydrogenation using oxidants such as  $\text{FeCl}_3$ ,  $\text{AlCl}_3$ , DDQ/TfOH,  $\text{Cu}(\text{OTf})_2$  which 'graphitizes' the polymer.<sup>32</sup>

This review focuses on the bottom-up, solution-phase synthesis of GNRs, categorized by the polymerization techniques employed, discussed chronologically while highlighting recent advancements. By examining the various polymerization methods, this review aims to provide a comprehensive overview of how these reactions produce a diverse range of GNRs. It begins by discussing the structure and properties of GNRs, followed by an overview of synthesis approaches. The core of the review concerns the key polymerization reactions used in GNRs synthesis, concluding with a brief discussion of the Scholl reaction and its alternatives such as topochemical polymerizations and photochemical methods including cyclodehydrochlorination (CDHC).



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Fig. 1 Schematic representation of some carbon-based allotropes.

## 2. Structure and properties of GNRs

As mentioned in the previous section, depending on how a graphene lattice is ‘cut’, GNRs with various edge geometries and widths can be created, each with distinct properties.<sup>9</sup> Many edge geometries are possible for GNRs (Fig. 2). Zigzag and armchair GNRs (ZGNRs and AGNRs, respectively) were amongst the first theoretically studied edge geometries.

Fujita *et al.* first studied ZGNRs and AGNRs using the tight-binding approach and found that AGNRs show semiconducting behavior and found a ‘peculiar’ flat band for ZGNRs. Upon applying the Hubbard-model within the mean field approximation, it was found that a band gap opening occurs in ZGNRs due to spin polarized electron–electron interactions.<sup>33</sup> Further theoretical investigations using more complex first-principle calculations have further confirmed these initial results<sup>34,35</sup> and built on them. For example, AGNRs can be split into three distinct families based on their width: semiconducting  $N = 3p$ ,  $N = 3p + 1$  and metallic  $N = 3p + 2$  (where  $N$  refers to the width of the AGNR as defined in Fig. 2). ZGNRs spin polarization is predicted to be ferromagnetic along a given edge and antiferromagnetic across the width of the ribbon.<sup>36</sup>

Both theoretical predictions have been either directly or indirectly confirmed *via* experimental investigations.



Fig. 2 Different ways a graphene lattice can be cut producing (a) variety of edge features and (b) conventional numbering methods of different sized GNRs.

G. Z. Magda *et al.* used OSS synthesis of a variety of AGNRs and ZGNRs and found an inverse proportionality between bandgap and width for AGNRs with good theoretical agreement on separation of AGNRs into their respective classes (Fig. 4a). The researchers indirectly confirmed the presence of edge magnetism in ZGNRs by assessing the electronic structure. It was concluded that only the presence of magnetic edges explains both the bandgap and observed semiconductor-to-metal transition (Fig. 4b), supporting the presence of room temperature stable magnetic order in precisely engineered zigzag edges.<sup>24</sup>

Aside from zigzag and armchair edges, other edge geometries also exist with their own distinct properties. Notably, cove and fjord edge GNRs are of particular interest due to their curved conformations leading to helically twisted ribbons.<sup>37,38</sup> These properties prove ideal as reduced  $\pi$ – $\pi$  stacking interactions of these curved GNRs lead to increased dispersibility in organic solvents thereby leading to highly liquid processable GNRs allowing more facile characterization by solid state nuclear magnetic resonance (ss-NMR), Fourier-transform Infrared spectroscopy (FT-IR), Raman spectroscopy, UV/vis spectroscopy and matrix assisted laser desorption/ionization time of flight mass spectrometry (MALDI-TOF MS).<sup>39</sup> The non-planarity arises from the steric repulsion between hydrogen atoms in the [4]-helicene and [5]-helicene precursor shapes.<sup>40</sup>

Other factors affecting GNR properties include chemical modification. These range from backbone nitrogen doped or boronated GNRs<sup>41–43</sup> to edge functionalization such as chlorination.<sup>44</sup> The introduction of non-hexagonal rings<sup>45</sup> and nanopores<sup>46</sup> also proves an effective way to engineer properties. Above all, presence of multi-edge systems giving rise to chiral GNRs<sup>37,38,47–51</sup> are amongst the most effective way to produce GNRs with unique properties, as will be discussed vastly in this review.



Fig. 3 A generalized schematic of bottom-up solution-phase synthetic steps towards GNRs.





Fig. 4 Bandgap measurements from tunnelling spectra: (a) AGNRs exhibit an inverse relationship between bandgap and ribbon width, consistent with theoretical classifications. (b) ZGNRs display band structures influenced by edge magnetism, showing a sharp semiconductor (antiferromagnetic) to metal (ferromagnetic) transition. Theoretical results use the Hubbard model within a mean-field approximation. Error bars reflect thermal broadening and substrate interactions.<sup>24</sup> Reproduced from ref. 24 with permission from [Springer], copyright [2014].

### 3. Bottom-up solution-phase synthesis of GNRs

#### 3.1. Polymerization methods

**3.1.1.  $A_2B_2$  Diels–Alder polymerization.** Amongst the first reactions utilized to afford the polymer precursors for planarization of GNRs was the  $A_2B_2$  Diels–Alder reaction. In 2000, Shifrina *et al.* investigated the Diels–Alder reaction between 1,4-bis(2,4,5-triphenylcyclopentadienone-3-yl) benzene (**1.0**) and 1,4-diethynylbenzene (**1.1**) (Scheme 1) using different concentrations of the monomeric building block and different reaction times.

Regioisomeric branched polyphenylenes (**1.2**) with  $M_w$  in the range of 12–120 kg mol<sup>-1</sup> were obtained and subjected to cyclodehydrogenation reactions using copper(II) trifluoromethanesulfonate and  $AlCl_3$  producing a dark brown product (**1.3**) which was studied by FT-IR and Raman spectroscopy as NMR analysis was not possible due to the products insolubility in standard solvents.<sup>52</sup> Later in 2003, the same group replaced

the bisdienophile from 1,4-diethynylbenzene (**1.1**) to 1,4-diethynyl-2,5-di(4'-*tert*-butylphenyl) benzene (**1.4**) (Scheme 1). The polymer precursor (**1.5**) ( $M_w \sim 62.5$  kg mol<sup>-1</sup>, polydispersity index (PDI)  $\sim 2.5$ ) from this  $A_2B_2$  Diels–Alder reaction was subjected to cyclodehydrogenation by  $FeCl_3$  affording an AGNR (**1.6**) with kinked structure which as well as the previous spectroscopic techniques, also underwent UV/Vis spectroscopy to confirm its increased conjugation compared to the polymer precursor.<sup>53</sup>

Despite the use of the AB Diels–Alder reaction being the more popular choice in GNR synthesis due to its ability to afford long (> 200 nm) gulf edged GNRs,<sup>54</sup> as will be mentioned later in this review. The  $A_2B_2$  Diels–Alder reaction was recently used to synthesize and characterize multi-edged GNRs. In this 2022 paper,<sup>38</sup> Yang *et al.* successfully synthesized curved GNRs (cGNRs) with hybrid cove–armchair–gulf edge structures. To test the efficiency of the Scholl reaction, the synthesis of model compounds 11,11'-diethynyl-5,5'-bichrysene (**1.8**) and 7-di-*tert*-butyl-9,11-bis(4-(*tert*-butyl) phenyl)-10*H*-cyclopenta[*e*]pyren-10-



Scheme 1 Synthetic routes towards GNRs **1.3** and **1.6** using  $A_2B_2$  Diels–Alder polymerization. Reproduced from ref. 52 with permission from [ACS], copyright [2000] Reproduced from ref. 53 with permission from [ACS], copyright [2003].



one (**1.7**) via Diels–Alder reactions yielded **1.9** in 80%. This was subsequently planarized using a Scholl reaction with  $\text{FeCl}_3$ . MALDI-TOF MS revealed a 50/50 mixture of products, a fully and partially cyclized **1.9**. To improve cGNR yield and efficiency, a pre-fused approach was used (Scheme 2a). **2.1** was synthesized from **2.0** which had an integrated bichrysenene unit and additional *t*-Bu groups for solubility. This underwent a complete cyclization under Scholl conditions at room temperature in 2 hours.

Finally, an  $\text{A}_2\text{B}_2$  Diels–Alder reaction between **2.0** and **2.2** yielded precursor polymer **2.3** ( $M_w \sim 108.3 \text{ kg mol}^{-1}$ , PDI  $\sim 1.9$ ), which was planarized into cGNRs **2.4** averaging 71 nm in length (Scheme 2b). FTIR, Raman, and solid-state NMR characterized the product. UV/Vis spectroscopy showed an optical band gap of 1.61 eV, matching theoretical values, and terahertz spectroscopy revealed a charge carrier mobility above  $2 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ , highlighting a potential for single-GNR nanoelectronics.

Using a similar model compound approach as well as assistance in rational design using density functional theory (DFT) calculations, Feng *et al.* in 2024<sup>45</sup> optimised the  $\text{A}_2\text{B}_2$ -type Diels–Alder polymerization between dibenzocyclooctadiyne (**2.6**) and dicyclopenta[*e,l*]pyrene-5,11-dione derivative (**2.5**). A subsequent selective Scholl reaction of the obtained ladder-type polymer (LTP) precursor **2.7** ( $M_w \sim 30.4 \text{ kg mol}^{-1}$ , PDI  $\sim 1.15$ ) produced wavy GNRs (wGNRs) **2.8** of lengths up to 30 nm which incorporate periodic eight-membered rings into its

backbone (Scheme 3). Similarly to the cGNRs, the non-planar geometry of wGNR (**2.8**) efficiently prevents the inter-ribbon  $\pi$ – $\pi$  aggregation, leading to photoluminescence in solution. Using these properties, the researchers fabricated the first GNR-based organic light-emitting electrochemical cells (GOLECs). The cells showed fast response and short turn on times of under 10 s. The current density–voltage–luminance characteristics a peak luminance of  $120 \text{ cd m}^{-2}$  with high stability (Fig. 5).

**3.1.2.  $\text{A}_2\text{B}_2$  Suzuki polymerization.** The  $\text{A}_2\text{B}_2$  type Diels–Alder polymerization reaction provides a simple way to produce GNRs with unique multi-edge structures and innovative designs without the use of a metal catalyst.<sup>38,45,52,53</sup> However, the high temperature requirements for this method leave room for improvement.<sup>10</sup> In 2008, Yang *et al.* developed an  $\text{A}_2\text{B}_2$  Suzuki coupling reaction between the bis-boronic ester (**2.9**) with diiodobenzene derivative (**3.0**) at  $120^\circ\text{C}$  which successfully produced the polyphenylenes (**3.1**) in 75% yield.<sup>55</sup> This reaction was amongst the most sterically hindered Suzuki couplings at the time. The polyphenylene **3.1** ( $M_w \sim 13.9 \text{ kg mol}^{-1}$ , PDI  $\sim 1.2$ ) was subjected to Scholl reaction conditions with  $\text{FeCl}_3$  providing linear AGNRs (**3.2**) of lengths up to 12 nm (Scheme 4). The bulky 3,7-dimethyloctyl groups as sidechains increased the processability of AGNRs causing excellent dispersibility in common organic solvents. Both UV/vis absorption spectroscopy and MALDI-TOF MS studies revealed the formation of the AGNRs with an optical band gap of 2.2 eV revealed. Various



**Scheme 2** (a) Synthesis of model compounds **1.9** and **2.1**; (b) synthetic route toward cGNR (**2.4**) via  $\text{A}_2\text{B}_2$  Diels–Alder polymerization. Reproduced from ref. 38 with permission from [Wiley], copyright [2022].



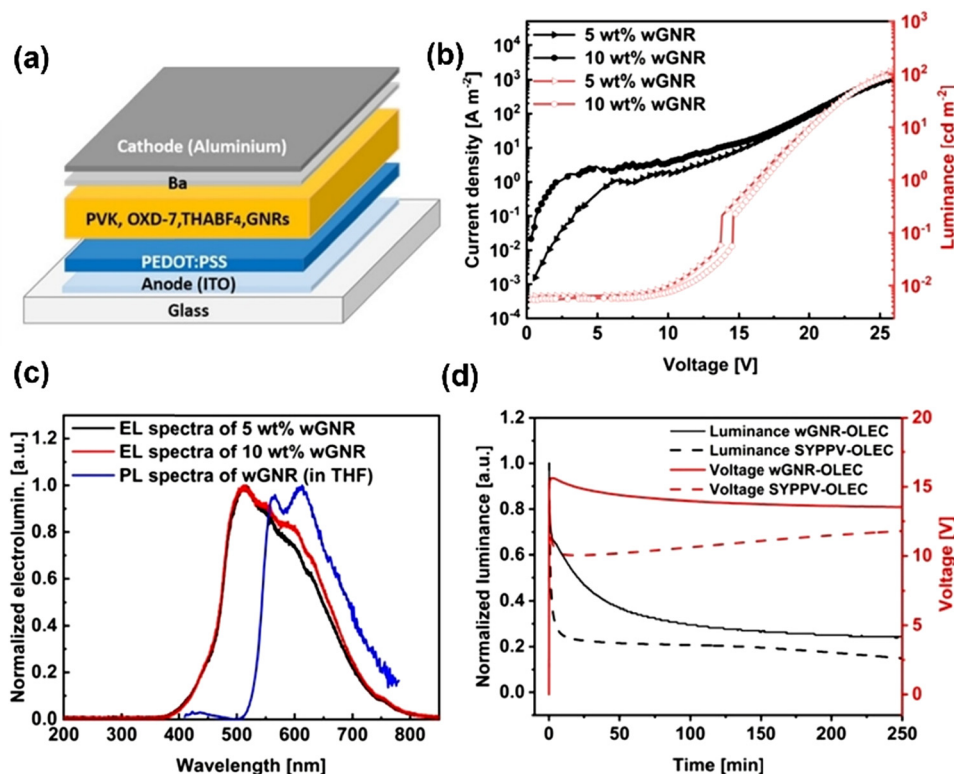


**Scheme 3** Synthesis of wGNRs (**2.8**) via  $A_2B_2$  Diels–Alder reaction between **2.5** and **2.6**. Reproduced from ref. 45 with permission from [Wiley], copyright [2024].

microscopic studies of these novel structures showed a high tendency to self-assemble opening a gateway into the use of GNRs in FETs and optoelectronic devices.

The high functional group tolerance of the Suzuki reaction,<sup>56</sup> allows for band gap engineering in GNRs synthesis by addition of heterostructures in the Suzuki precursors. In 2018, using a model study synthesis, Li *et al.* synthesized two unsymmetrical edged GNRs containing benzothiadiazole and benzotriazole groups.<sup>49</sup> The bispinacol borates (BPin) (**3.3**) were reacted with triaryl dibromide **3.4** under Suzuki coupling conditions using  $Pd(PtBu_3)_2/K_3PO_4$  as the catalyst-base combination to form the desired polymer precursors **3.5** (Scheme 5a). Subsequently, precursor **3.5** of similar molecular weights ( $M_w \sim 37.4 \text{ kg mol}^{-1}$ ,

PDI  $\sim 1.73$ ) were cyclodehydrogenated using a 2,3-dichloro-5,6-dicyano-1,4-benzoquinone/triflic acid (DDQ/TfOH) protocol forming **3.6** (Scheme 5a). Once the AGNRs (**3.6**) were isolated as black powders, they were found to have good dispersibility in THF, chlorobenzene, and *o*-dichlorobenzene. Thus, allowing for characterization by FTIR, Raman, UV-Vis spectroscopy, X-ray photoelectron spectroscopy (XPS), MALDI-TOF MS, atomic force microscopy (AFM), and scanning tunnelling microscopy (STM). Narrow optical band gaps of 1.03 eV and 1.09 eV for **3.6** were extracted for the *S* and *N*-butyl functionalised AGNRs, respectively. Due to the survival of such heterostructures *via* the Suzuki reaction, post-functionalisation of **3.7** was possible *via* a directed C–H borylation using  $BCl_3$  followed by  $ZnPh_2$  producing **3.8**

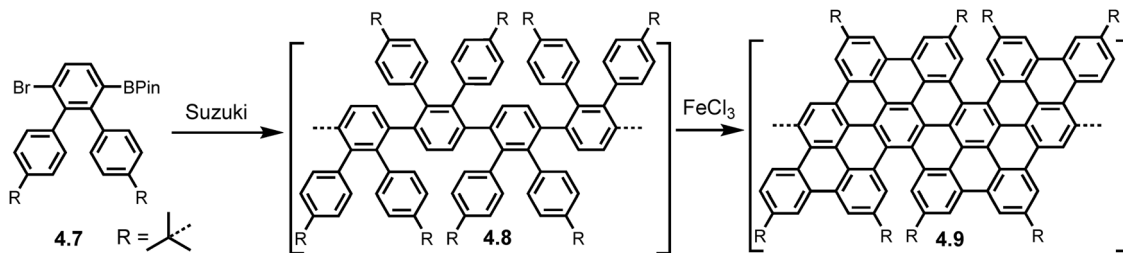


**Fig. 5** GOLEC fabrication with **2.8** as emissive layer. (a) Schematic representation of the GOLECs; (b) current density and luminance against voltage for GOLECs with various weight fractions of **2.8**; (c) EL spectra for GOLECs with 5 wt% and 10 wt% **2.8** and PL spectra of **2.8** in solution with THF as solvent; (d) luminance and voltage against time for GOLECs with 5 wt% **2.8** with a current density at  $100 \text{ A m}^{-2}$ .<sup>45</sup> Reproduced from ref. 45 with permission from [Wiley], copyright [2024].









**Scheme 8** Synthesis of fjord edge GNRs (**4.9**) from the AB Suzuki polymerization and subsequent Scholl reaction of **4.7**. Reproduced from ref. 37 with permission from [ACS], copyright [2021].

studies suggesting a similar conformation for the fGNR supported also by DFT calculations. Photoconductivity analysis revealed an intrinsic charge carrier mobility of  $100 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ , higher than values for other cGNRs<sup>38</sup> placing these highly distorted GNRs as a potential candidates for nanoelectronic devices.

GNR heterojunctions combine the distinct topological features of the individual ribbons allowing for further electronic bandgap engineering. Zhang *et al.* reported a novel chain-growth polymerization strategy that allowed for formation of a GNR heterojunction with  $N = 9$  (where  $N$  refers to the number of pairs of atoms per GNR unit cell) armchair and chevron GNRs segments (9-AGNR/cGNR).<sup>47</sup> The controlled Suzuki catalyst-transfer polymerization (SCTP) between 2-(6'-bromo-4,4''-ditetradecyl-[1,1':2',1''-terphenyl]-3'-yl) boronic ester (**5.1**) and 2-(7-bromo-9,12-diphenyl-10,11-bis(4-tetradecylphenyl)-triphenylene-2-yl) boronic ester (**5.3**), followed by the Scholl reaction of the obtained block copolymer (**5.4**) with controlled  $M_w$  of  $26.1 \text{ kg mol}^{-1}$  (PDI  $\sim 1.45$ ), led to the formation of the 9-AGNR/cGNR (**5.5**) (Scheme 9) which was unambiguously validated by FT-IR, Raman, and UV/Vis spectroscopies and estimated to be 14 nm in length.

Lee *et al.* also reported SCTP type reactions to synthesise AGNRs of a controlled length. **5.6** was subjected to optimised SCTP conditions using a RuPhos-Pd catalyst system giving

polymer (**5.7**) in an excellent 85% yield ( $M_n \sim 29.8 \text{ kg mol}^{-1}$ , PDI  $\sim 1.14\text{--}1.39$ ). This alkyne containing precursor underwent mild alkyne benzannulation forming 5-AGNRs (**5.8**). Subsequently, Scholl conditions were applied to **5.8** using a DDQ/TfOH protocol to laterally extend the conjugated framework and form the 11-AGNR **5.9** (Scheme 10).<sup>58</sup> These innovative SCTP type reactions prove to be a useful method in controlling the length of GNRs and allow for milder conditions to be used such as alkyne benzannulation.

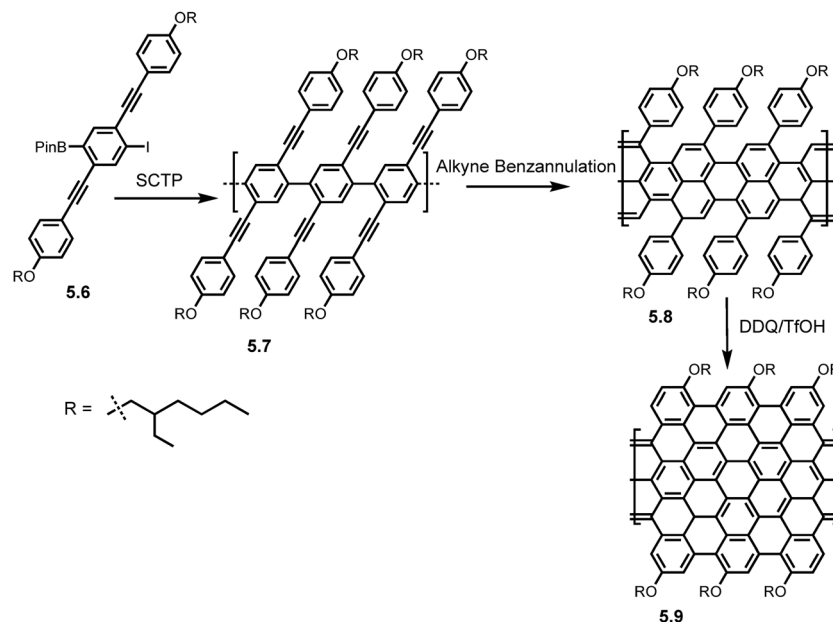
**3.1.4. AA Yamamoto polymerization.** The  $A_2B_2$  type Diels-Alder and Suzuki polymerization require two monomers for reactions to form GNRs making them stoichiometrically sensitive. This limits its ability to form consistent high weight polyphenylene precursors due to possible inefficient reactions.<sup>59,60</sup> On the other hand, the AA type Yamamoto coupling requires only one dihalogenated monomer thereby reducing this effect. The monomer can be of high molecular weight allowing for facile lateral extensions of GNRs as well of the incorporation of multiple edge structures within one monomer.

In 2012, Schwabz *et al.* produced monomer **6.0** through a multistep synthesis, enabling its AA Yamamoto polymerization using bis(cycloocta-(1,5)-diene) nickel (0) cycloocta-(1,5)-diene ( $\text{Ni}(\text{COD})_2$ ) to form polyphenylene precursor **6.1** (Scheme 11).<sup>61</sup>



**Scheme 9** Synthetic route toward 9-AGNR/cGNR (**5.5**) using subsequent SCTP of **5.1** and **5.3**. Reproduced from ref. 47 with permission from [Wiley], copyright [2023].





**Scheme 10** Alkyne benzannulation of polymer precursor **5.7** to form 5-AGNR **5.8** and subsequently 11-AGNR **5.9**. Reproduced from ref. 58 with permission from [ACS], copyright [2023].

Precursor **6.1** had a weight ( $M_w$ ) of up to  $52 \text{ kg mol}^{-1}$  (PDI  $\sim 1.2$ ) corresponding to 21–24 repeating units of **6.0** leading to GNRs of lengths up to 30 nm, which at the time was superior to the widely used  $A_2B_2$  Suzuki polymerization. The subsequent Scholl reaction using  $\text{FeCl}_3$  as an oxidant yielded the GNR (**6.2**). These GNRs were found to be soluble/dispersible in *N*-methyl-2-pyrrolidone (NMP) allowing for characterization by MALDI TOF, FT-IR and Raman spectroscopy. UV/Vis spectroscopy extracted a narrow optical band gap of 1.12 eV matching well with theoretical calculations of 1.09 eV. Due to the lateral extension of GNRs *via* the AA Yamamoto reaction, this led to high dispersibility allowing for facile liquid phase processing which offers benefits in characterization but also in device fabrication. M. El Gemayel *et al.* fabricated thin-film FET devices using a co-deposited GNR/poly(3-hexylthiophene)

(P3HT) blend.<sup>62</sup> The device showed a 24% increase in charge-carrier mobility compared to a pristine P3HT device. The on/off ratio remained robust upon addition of the GNR, and the photo response of the transistor was doubled compared to the pristine FET.

Monomer **6.3** substituted with *N*-*n*-hexadecyl maleimide units was subjected to AA Yamamoto conditions using  $\text{Ni}(\text{COD})_2/\text{COD}$  yielding polymer precursors (**6.4**) ( $M_w \sim 15.6\text{--}186 \text{ kg mol}^{-1}$ , PDI  $\sim 1.2\text{--}1.5$ ). Intramolecular oxidative cyclodehydrogenation using  $\text{FeCl}_3$  as the Lewis acid and oxidant, converted the polymer precursors into armchair edged GNR-AHMs (**6.5**) with uniform width (1.7 nm) and lengths of 6, 11, and 58 nm, respectively (Scheme 12). The bulky AHM side groups caused excellent dispersibility (concentrations of up to  $5 \text{ mg mL}^{-1}$ ) in many organic solvents such as tetrahydrofuran (THF). Loss of extended



**Scheme 11** Synthetic route toward GNR (**6.2**) *via* AA type Yamamoto polymerization of monomer **6.0**. Spectroscopic studies on single GNRs were conducted by Huang *et al.* in 2018.<sup>21</sup> Reproduced from ref. 61 with permission from [ACS], copyright [2012].





**Scheme 12** Synthetic route to GNR (**6.5**) via AA type Yamamoto polymerization of monomer **6.3**. Reproduced from ref. 21 with permission from [ACS], copyright [2018].

tail in Near Infrared (NIR) region (compared to previously synthesised GNRs) during UV/Vis experiments confirm that dilute dispersions in THF ( $<0.1 \text{ mg mL}^{-1}$ ) consist mainly of nonaggregate ribbons (Fig. 6), exhibiting high fluorescence quantum yield (9.1%) and long fluorescence lifetime (8.7 ns). This dispersibility allows resolving the real-time ultrafast excited-state dynamics of the GNRs, which displays features of small, isolated molecules in solution. This successful synthesis and edge functionalisation using the AA Yamamoto coupling reduces the limitations of solution-based synthesis, allowing for improved characterization of individual GNRs, which is typically difficult compared to OSS methods.

The AA type Yamamoto polymerization can be used to form multi-edge cGNRs much like the  $A_2B_2$  Diels–Alder.<sup>38</sup> Niu *et al.* reported the first synthesis of a cGNR with cove, armchair and usually unstable, zigzag edges.<sup>51</sup> This feature is of particular interest due to the survival of intrinsic zigzag properties in chiral GNRs.<sup>28</sup> Following the multi-step synthesis of monomer **6.6**, AA type Yamamoto polymerization afforded **6.7** ( $M_w \sim 20.8 \text{ kg mol}^{-1}$ , PDI  $\sim 1.15$ ) which was subjected to Scholl reaction conditions using  $\text{FeCl}_3$  producing the cGNR (**6.8**) (Scheme 13) with a narrow width of 0.7 nm and average length of 23 nm. Along with FT-IR, Raman MALDI-TOF MS, UV/Vis spectroscopy revealed a bandgap of 1.22 eV and incredibly, by

THz photoconductivity analysis, a record high intrinsic charge carrier mobility of  $600 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$  was calculated.

Incorporation of periodic zig-zag segments was later executed by Wang *et al.* in 2021.<sup>50</sup> Following the synthesis of the structurally flexible S-shaped monomer **6.9**, AA Yamamoto polymerization was carried out using  $\text{Ni}(\text{COD})_2$  as a catalyst, producing the ‘snake-like’ polymer precursor **7.0** ( $M_w \sim 29.8 \text{ kg mol}^{-1}$ , PDI  $\sim 1.2$ ). Upon ‘graphitization’ using  $\text{FeCl}_3$  as Lewis acid and oxidant the multi-edged cGNR (**7.1**) was formed (Scheme 14) with an average length of 20 nm. The successful synthesis of the cGNR (**7.1**) was confirmed by FT-IR, Raman and ss-NMR investigations. UV/Vis spectroscopy revealed a narrow bandgap of 0.99 eV and THz photoconductivity analysis calculated a charge carrier mobility of  $20 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ . Interestingly DFT simulations show an ‘up-down’ curved topology for the ribbon.

Recent theoretical studies into the use of porphyrin units in GNR backbones have been of interest due to their enhanced delocalization and spin ground states making them useful for molecular wires<sup>63</sup> or spintronic applications.<sup>64</sup> Efforts in OSS to functionalize GNRs with porphyrins has been successful with the intrinsic properties of the porphyrin remaining intact.<sup>65</sup> However, Chen *et al.* managed to successfully synthesize the first solution-made porphyrin-fused GNR (PGNR)<sup>66</sup> by utilizing the AA Yamamoto polymerization of **7.2** yielding **7.3**



**Fig. 6** (a) Absorption and (b) PL spectra of GNR-AHMs (**6.5**) in THF (0.03 and  $0.003 \text{ mg mL}^{-1}$  for absorption and PL measurements, respectively).<sup>21</sup> Reproduced from ref. 21 with permission from [ACS], copyright [2018].





**Scheme 13** Synthetic route toward multi-edged GNR **6.8** via AA type Yamamoto of monomer **6.6**. Reproduced from ref. 38 with permission from [Wiley], copyright [2022].

( $M_w \sim 132.7 \text{ kg mol}^{-1}$ , PDI  $\sim 2.46$ ) in 93%, followed by an oxidative cyclodehydrogenation using a DDQ/TfOH protocol forming **7.4** (Scheme 15). The PGNRs (**7.4**) were characterized by ss-NMR, FT-IR, Raman and XPS. **7.4** functionalised with the dodecyl aryl group was found to exhibit a charge carrier mobility of  $>400 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$  via THz photoconductivity analysis, an optical bandgap of 1.0 eV was extracted from absorption spectra. The charge transport behaviour of **7.4** was investigated by fabricating single nanoribbon devices (Fig. 7) which showed FET behaviour at room temperature and clean single-electron-transistor behaviour at millikelvin temperatures. The non-differential conductance (NDC) regions showed a minimum voltage below 10 mV indicating low power operation possibilities.

**3.1.5. AB type Diels–Alder polymerization.** The synthesis of long GNRs ( $>100 \text{ nm}$ ) is inherently difficult using transition-metal (TM) catalysed reactions due to termination by loss of functional groups.<sup>67</sup> Therefore, the Diels–Alder polymerization is a desired route to this target. Use of the  $A_2B_2$  Diels–Alder polymerization affords GNRs with lengths up to 71 nm.<sup>38</sup> However, stoichiometric inefficiency limits this reaction.<sup>59,60</sup>

In 2014, Narita *et al.* subjected the monomer building block (**7.5**), which consists of a cyclopentadienone as the conjugated diene and an ethynyl group as the dienophile, to AB type Diels–Alder conditions affording polymer (**7.6**) which had an  $M_w$  between 24–620  $\text{kg mol}^{-1}$  (PDI  $\sim 3$ –14), higher than any reported weights using Yamamoto or Suzuki reactions at the time. The polymer precursor (**7.6**) was finally ‘graphitized’



**Scheme 14** Synthetic route toward GNR **7.1** with periodic zigzag segments via AA type Yamamoto polymerization of monomer **6.9**. Reproduced from ref. 50 with permission from [ACS], copyright [2022].





**Scheme 15** Synthesis of PGNR (**7.4**) from monomer **7.2** using AA type Yamamoto polymerization. Reproduced from ref. 66 with permission from [Springer], copyright [2024].

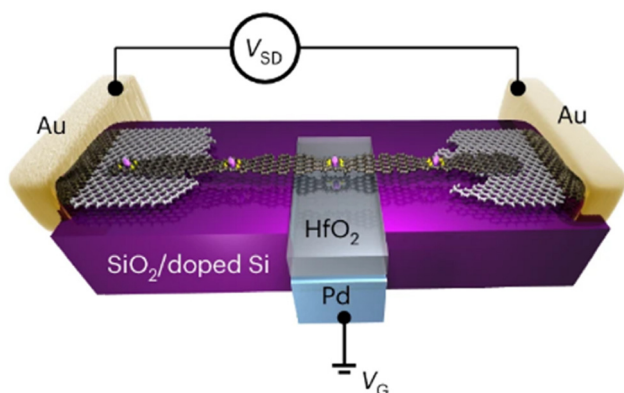
using Scholl reaction conditions with  $\text{FeCl}_3$  affording the GNR (**7.7**) with lengths  $> 200$  nm (Scheme 16a).<sup>54</sup> Model dimer and trimer compounds were subjected to ‘graphitization’ and analysed by MALDI-TOF MS to confirm the successful formation of the GNR (**7.7**). **7.7** was fully characterised by FT-IR, Raman and ss-NMR spectroscopy. THz probing found the intrinsic charge carrier mobilities as large as  $15\,000\text{ cm}^2\text{ V}^{-1}\text{ s}^{-1}$  along with a large optical bandgap of 1.88 eV from absorption spectra. The strong aggregation of the GNRs made characterization difficult however the presence of dodecyl chains in monomer **7.5** improved their dispersibility allowing for characterization, nonetheless.

Installation of different functional groups on the periphery of such GNRs is feasible by modification to the starting monomer. To diminish the bandgap, the same researchers shortly after performed a simple lateral phenyl extension to produce monomer **7.8** which was heated to 260–270 °C to induce AB type

Diels–Alder polymerization forming wider ( $\sim 2$  nm) polymer precursors **7.9** with a largely improved  $M_w \sim 230\text{--}550\text{ kg mol}^{-1}$  and PDI of 2.8–3.8. Post oxidative cyclodehydrogenation afforded **8.0** with lengths up to 390 nm (Scheme 16b).<sup>68</sup> Similarly, the GNRs were fully characterized and UV-vis spectroscopy eluded a smaller bandgap of 1.24 eV.

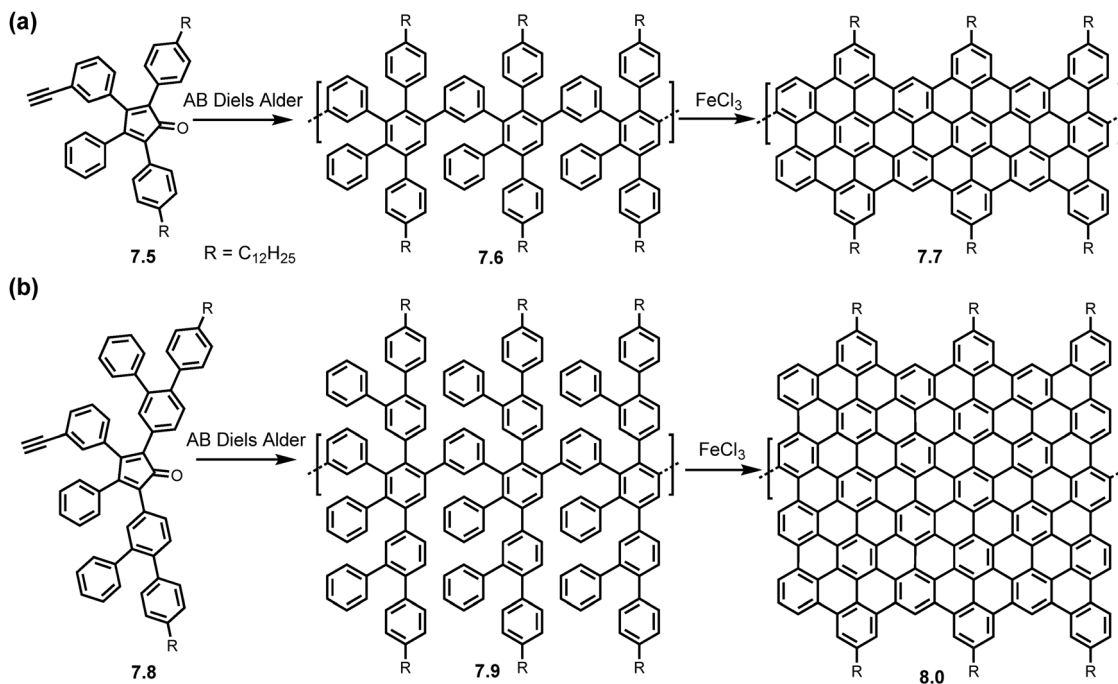
Götz *et al.* used methoxy-functionalized monomer **8.1** to yield polymer **8.2** ( $M_w \sim 31\text{--}54\text{ kg mol}^{-1}$ , PDI  $\sim 1.5\text{--}1.7$ ) which was planarized to methoxy-functionalised GNR (**8.3**) by oxidative cyclodehydrogenation using  $\text{FeCl}_3$  (Scheme 17). Theoretical simulations predicted a reduced bandgap of **8.3** compared to a hydrogen passivated parent (Fig. 8) due to geometric distortion caused by the methoxy groups along with the electron donating nature of the functionality. This was verified by the fingerprint of band shrinkage seen in the red shift of the UV/Vis spectra. THz analysis revealed a 25% increase in photoconductivity due to reduced effective mass of charge carriers in **8.3**.<sup>69</sup>

The modifications on the periphery of these monomers are not limited to simple lateral extensions. Keerthi *et al.* successfully synthesized GNRs with anthraquinone (AQ) and naphthalene/perylene monoamide (NMI/PMI) units achieved through Suzuki coupling of polyphenylene precursors with bromo groups, validated by MALDI-TOF MS, FTIR, Raman and XPS.<sup>30</sup> AB Diels–Alder polymerization of monomer **8.4** produced polymer precursor **8.5** which was subjected to Suzuki coupling conditions ( $\text{Pd}(\text{PPh}_3)_4/\text{K}_2\text{CO}_3$ ) with three boronic esters, 2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)anthracene-9,10-dione (AQ-BPin), *N*-(2-ethylhexyl)-4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)naphthalene-1,8-dicarboximide (NMI-BPin), and *N*-(1-heptyloctyl)-9-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)perylene-3,4-dicarboximide (PMI-BPin), producing functionalised polymer precursors **8.6** (A = AQ, NMI, PMI) with  $M_w$  as high as  $161\text{ kg mol}^{-1}$  and PDI  $< 3$ . Subsequent intramolecular oxidative cyclodehydrogenation using  $\text{FeCl}_3$  as oxidant afforded the GNRs (**8.7**) (Scheme 18), with possible lengths of up to 1000 nm.



**Fig. 7** Scheme of the electronic devices, where a single strand of PGNR (**7.4**) bridges two graphene electrodes (separated by a nanogap) connected to gold pads. A palladium gate is deposited on the  $\text{SiO}_2/\text{Si}$  surface and covered with a 10-nm-thick  $\text{HfO}_2$  dielectric.<sup>66</sup> Reproduced from ref. 66 with permission from [Springer], copyright [2024].





**Scheme 16** Synthesis of (a) GNR (7.7) and (b) laterally extended GNR (8.0) via AB type Diels–Alder reaction. Reproduced from ref. 54 with permission from [Springer], copyright [2014]; Reproduced from ref. 68 with permission from [ACS], copyright [2014].

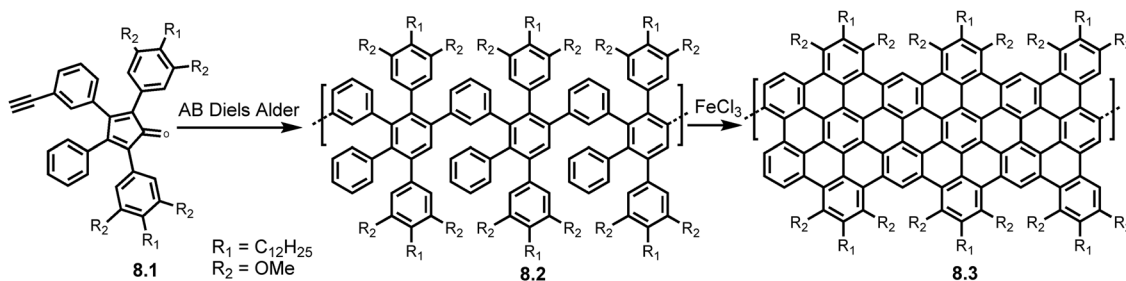
However, these vast lengths could have also been due to end-to-end tangling of individual ribbons. Interestingly, drop casting the three GNRs (8.7) (A = AQ, NMI, PMI), revealed a self-assembling behaviour. The PMI functionalised GNR (8.7) formed a fascinating rectangular network providing evidence that edge functionalisation of GNRs can be used to tune their supramolecular assemblies.

More sophisticated edge functionalization was later developed by Slota *et al.* where GNRs functionalized with spin-bearing nitronyl nitroxide (NIT) radicals (8.9) were synthesized from the AB Diels–Alder polymerization. A non-cyclized polyphenylene precursor (8.8) with NIT radicals was also synthesized as means of a control (Fig. 9). Using electron paramagnetic resonance (EPR) spectroscopy, the researchers manipulated the spin states to measure the spin-dynamics and spin-environment interactions. Spin coherence times were measured and found to be in a promising range of microseconds at room temperature, ideal for spintronic applications and decoherence channels were

identified.<sup>31</sup> The survival of the brominated monomer through the AB Diels–Alder polymerization in this case, opened the possibility of edge functionalization and direct manipulation and observance of magnetism in GNRs for the first time.

The functionalization opportunities provided by the AB Diels–Alder polymerization has expanded GNR applications. Rogers *et al.* demonstrated that GNRs 9.0 effectively supported gold nanoparticles (AuNPs) for electrocatalytic CO<sub>2</sub> reduction (Fig. 10). The GNR-AuNP interaction reduced the onset potential by several hundred millivolts, increased catalytic activity, and improved nanoparticle dispersion, enhancing the electrochemically active surface area. The GNR-AuNPs also exhibited over 100-fold greater stability and performance compared to amorphous carbon supports. Edge functionalization of the GNRs with ester groups tuned the catalytic environment, shifting the rate-determining step and altering the reaction mechanism.<sup>70</sup>

Other innovative uses include the inclusion of additional ‘capping’ monomers to form GNR heterostructures for



**Scheme 17** Synthesis of methoxy-functionalised GNRs (8.3) via AB type Diels–Alder polymerization of monomer 8.1. Reproduced from ref. 69 with permission from [RSC], copyright [2022].





Fig. 8 DFT band structures of GNR-OMe (**8.3**) ( $R = \text{OMe}$ ; blue, right) and GNR-H ( $R = \text{H}$ ; orange, left). The two band structures are aligned to the vacuum level, which is set to zero.<sup>69</sup> Reproduced from ref. 69 with permission from [RSC], copyright [2022].

applications in next generation molecular diodes (Fig. 11)<sup>71</sup> and bulky edge functionalization has led to pure carbon based GNRs to be used as single electron transistors.<sup>72</sup> Also, advancements in 'clickable' GNRs have been made for functionalisation with fluorescent dyes for facile imaging,<sup>73</sup> or for successful application in biosensor interfaces (Fig. 12).<sup>74</sup>

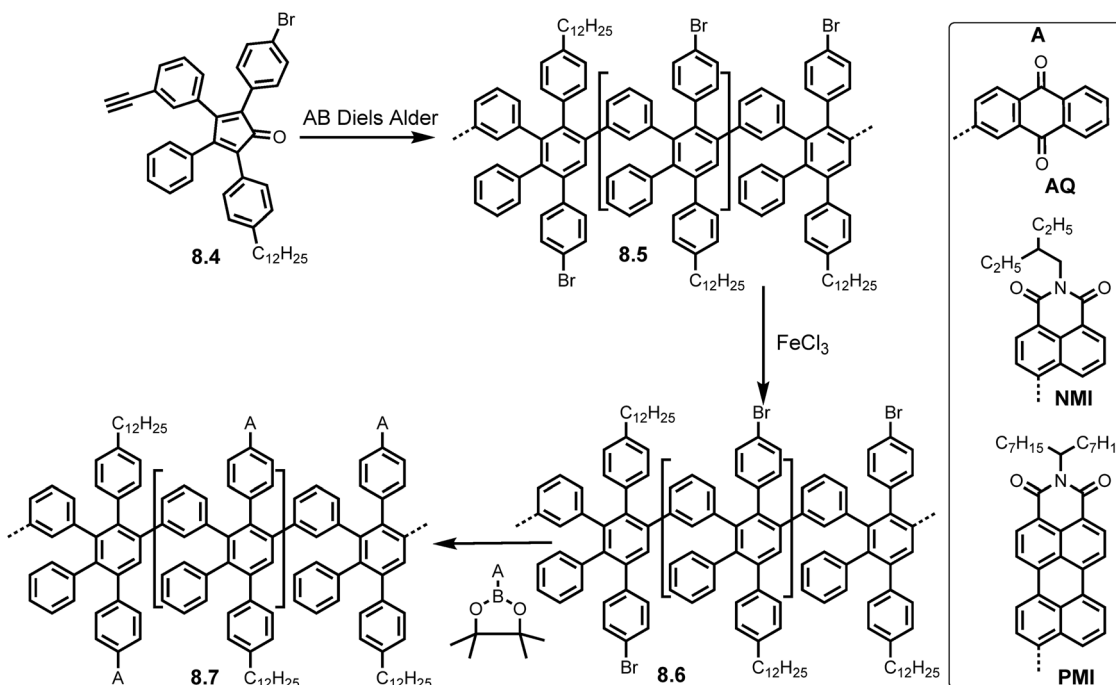
Manipulation of the cyclopentadienone monomer is also possible (Scheme 19) as demonstrated by Veber *et al.* in 2020

where modifications to a borated AB Diels–Alder monomer (**9.4**) via Suzuki coupling with **9.5** formed an acetal protected cyclopentadienone monomer **9.6** and thus an acetal functionalised polyphenylene precursor (**9.7**). Acid catalysed deprotection yielded the aldehyde functionalised precursor which was subjected to Scholl reaction conditions to yield the aldehyde functionalised GNRs (**9.8**). Subsequent Lewis acid catalysed interfacial polymerization caused the growth of Covalent Organic Framework GNRs (COF-GNRs) using imine cross links (Fig. 13). These COF-GNRs showed 2D self-assembly and crystals gave access to a large surface area ( $>10^5 \text{ nm}^2$ ).<sup>75</sup>

Outside the realm of  $\text{AB}/\text{A}_2\text{B}_2$  Diels–Alder polymerisations, the Diels–Alder reaction can be used post-polymerization using Ring Opening Alkyne Metathesis (ROAMP) techniques.<sup>76</sup> Kugelgen *et al.* initiated ROAMP of the rationally designed alkyne **9.9** using catalyst **10.0** and alkyne **10.1** to terminate. This repeating alkyne polymer **10.3** underwent a Diels–Alder benzannulation to form the GNR precursor polymer **10.4**. A  $\text{FeCl}_3$  Scholl condition was applied forming the desired cGNR **10.5** with lengths up to 68 nm. The ROAMP method proves useful due to the structural control in functionalized end groups and controlled lateral extension by taking advantage of precise Diels–Alder reactions (Scheme 20).

### 3.2. Alternative planarization techniques

So far, the planarization techniques discussed have been using Scholl reactions. The Scholl Reaction is traditionally an oxidative cyclodehydrogenation reaction using  $\text{FeCl}_3$  as a Lewis acid and oxidant. The mechanism of this reaction is not definitively known due to the dual behaviour of  $\text{FeCl}_3$  however it is



Scheme 18 Synthetic route to peripherally modified GNRs (**8.7**) with AQ, NMI and PMI electron withdrawing groups. Reproduced from ref. 30 with permission from [ACS], copyright [2017].





Fig. 9 Structures of NIT-polyphenylene **8.8** and NIT-GNRs **8.9** synthesized via AB Diels–Alder polymerization. Reproduced from ref. 31 with permission from [Springer], copyright [2018].

generally accepted to follow either a radical cation mechanism or via the formation of an arenium ion.<sup>77</sup> The Scholl reaction is an easy procedure which produces good yields for GNRs.<sup>10</sup> However, in some cases it can lead to a mixture of fully cyclized and partially cyclized products<sup>38</sup> or undesirable ring formation/chlorination.<sup>78</sup> The uncertainty in the reaction mechanism of the Scholl reaction, contributes to variability in efficiency across novel substrates.<sup>79</sup> While FeCl<sub>3</sub> is commonly used, alternative systems like DDQ/TfOH are also effective.<sup>49,66</sup> Oxidant choice is influenced on the substrates electronics: theoretically, electron-rich arenes would react via an arenium ion pathway promoted by strong Lewis acids (e.g., FeCl<sub>3</sub>, AlCl<sub>3</sub>), whereas electron-poor substrates would favour a radical cation mechanism, which can be facilitated by single-electron oxidants (e.g., DDQ) in the presence of strong Brønsted acids like TfOH.<sup>32,80</sup> In this section, alternative pathways to planarization, using alkyne precursors and photochemical methods, are briefly discussed. Whilst some of these methods are not strictly solution-mediated synthesis, they are important to mention to give depth into the variety of methods used in the field.

Other reviews have discussed these methods in more detail<sup>80,81</sup> however such depth is beyond the scope of this review.

**3.2.1. GNRs from alkyne precursors.** Alkyne rich precursors are favourable for the formation of carbon nanostructures owing to their intrinsic thermal instability.<sup>80</sup> Topochemical polymerization is the solid-state photo/heat induced 1,4-addition polymerization of conjugated triple bonds giving rise to a polymer with three cumulated double bonds per repeating unit.<sup>82</sup> In 2016, Jordan *et al.* used monomers containing butadiyne units (**10.6**) and using topochemical polymerization converted these units into polydiacetylenes (**10.7**) set up for further aromatization to afford GNRs (**10.9**) (Scheme 21a) (~1.36 nm in width).

The GNRs (**10.9**) had a bandgap of 1.4 eV. The solid-state reaction only required UV light or heat to proceed with the addition of no other reagents.<sup>83</sup> The same group later synthesized an AGNR (**11.2**) of the 3p + 2 class (Scheme 21b) using this method which would be predicted to have a very low bandgap. Use of Raman, FT-IR, ss-NMR show the target structure was synthesized however the harsh conditions used to perform the cyclization resulted in loss of the bulky side chains, causing analysis and processing to be difficult.<sup>84</sup> Other methods using alkyne precursors include alkyne benzannulation<sup>58</sup> (Scheme 10) however these methods typically still make use of a Scholl reaction as a last step.<sup>80</sup>

**3.2.2. Photochemical methods.** Whilst the use of alkyne precursors removes the need for additional reagents relying solely on heat and/or light, photochemical methods only rely on light removing the requirement of very high temperature reactions and can also be done in solution phase.<sup>80</sup>

The Mallory reaction is amongst the oldest photochemical carbon–carbon bonding reaction.<sup>85</sup> This reaction has been used widely in the synthesis of nanographene's from polyaromatic hydrocarbons (PAHs) however is rarely used in GNR synthesis.<sup>80</sup> Zhong *et al.* used monobromo, dibromo and monostannane derivatives and subjected them to Stille coupling conditions to form perylene diimides (PDIs) which afforded GNRs **11.4** after a Mallory reaction.<sup>86</sup> Scheme 22 shows the synthesis using a dibromo monomer **11.3**.



Fig. 10 (a) GNRs (**9.0**) synthesized (b) as support for AuNPs for electrocatalytic CO<sub>2</sub> reduction.<sup>70</sup> Reproduced from ref. 70 with permission from [ACS], copyright [2017].





**Fig. 11** (a) GNR heterostructures (**9.1**) synthesized for use in molecular diodes (b) Schematic  $I$ - $V$  measurement of GNR-based device (100) (c) Average photoresponsivity of six GNR-based devices on diamond at 573 K and  $-5$  V under vacuum conditions.<sup>71</sup> Reproduced from ref. 71 with permission from [ACS], copyright [2023].

A more popular photochemical method for GNR synthesis is reactions most notably the CDHC. The mechanism is believed to be a photochemical conrotatory electrocyclization followed

by elimination of HCl. This reaction affords very high conversion attributed to the entropic benefit of HCl release.<sup>81</sup> Scheme 23 shows the synthesis of thiophene annulated GNRs



**Fig. 12** Structures of 'clickable' GNRs used for (a) facile imaging via inclusion of fluorescent dyes (**9.2**) and (b) applications in biosensor interfaces (**9.3**). Reproduced from ref. 73 with permission from [ACS], copyright [2018] Reproduced from ref. 74 with permission from [RSC], copyright [2024].





**Scheme 19** Synthesis of COF-GNRs **9.8** using acetal protected cyclopentadienone monomer **9.4**. Reproduced from ref. 75 with permission from [Cell Press], copyright [2020].

(**11.8** And **12.1**) synthesized by Suzuki polymerization methods but with the replacement of the Scholl reaction with CDHC reaction due to the inclusion of chlorinated precursor **11.5**.<sup>87</sup>

## 4. Conclusion and future perspectives

Each polymerization technique discussed in this review offers distinct advantages and remains relevant in current literature for tailoring GNR structures for specific applications (Table 1). The  $A_2B_2$  Diels–Alder reaction, despite being one of the earliest methods, continues to be utilized in recent studies for synthesizing multi-edged GNRs with unique electronic properties

nm.<sup>38</sup> It has also enabled the incorporation of eight-membered rings, opening pathways for photoluminescent properties with successful device applications.<sup>45</sup> However, its broader use has declined due to advancements in other polymerization techniques.

Suzuki coupling remains widely adopted due to its broad functional group tolerance, which facilitates bandgap engineering. It enables both bandgap narrowing through the preservation of heterostructures in monomeric units with post-borylation options<sup>49</sup> and bandgap widening, owing to the stability of defined pores in GNR structures.<sup>46</sup> Unique AB variations of SCTP have also led to the synthesis of weight controlled GNR heterostructures, enabling precise topological engineering for potential quantum applications.<sup>47,58</sup>

The AA Yamamoto polymerization overcomes the stoichiometric inefficiencies of  $A_2B_2$  polymerization<sup>59,60</sup> while retaining the ability to synthesize multi-edged GNRs, including those with typically unstable magnetic zigzag edges.<sup>50,51</sup> This method has also enabled the solution-phase synthesis of highly sought porphyrin GNRs.<sup>66</sup> However, the reliance on TM catalysts in Suzuki and Yamamoto reactions limits the achievable GNR lengths due to functional group degradation.<sup>59,60</sup>

The AB Diels–Alder reaction addresses these limitations, enabling the synthesis of GNRs up to 600 nm in length<sup>54</sup> and offering versatile functionalization options.<sup>38,60</sup> Functional groups can be incorporated into monomeric building blocks<sup>68–70</sup> or retained through halogenated monomers, allowing post-functionalization with electron-withdrawing units for supramolecular assembly tuning,<sup>30</sup> spin-containing radicals for magnetic probing,<sup>31</sup> and fluorescent compounds for facile characterization.<sup>73</sup> These advances have broadened GNR applications, including catalytic enhancements with tunable reaction mechanisms,<sup>70</sup> molecular diode integration,<sup>71</sup> and biosensor development.<sup>74</sup>

Emerging directions include further rational design of monomers incorporating pores and non-hexagonal rings—areas that are still underexplored in the literature. The OSS method has demonstrated multiple examples of boron-containing GNRs *via*



**Fig. 13** Structural model of COF-GNRs with unit cell (black box), orientation displaying lattice plane (01–1) corresponding to distance between linkers (0.7 nm) (top), orientation displaying interlayer packing and lattice planes (001) (orange), (002) (gray), and (105) (blue) corresponding to 2.5, 1.25, and 0.35 nm (bottom).<sup>75</sup> Reproduced from ref. 75 with permission from [Cell Press], copyright [2020].





Scheme 20 ROAMP synthesis of cGNR **10.5** using catalyst **10.0**. Reproduced from ref. 76 with permission from [ACS], copyright [2019].

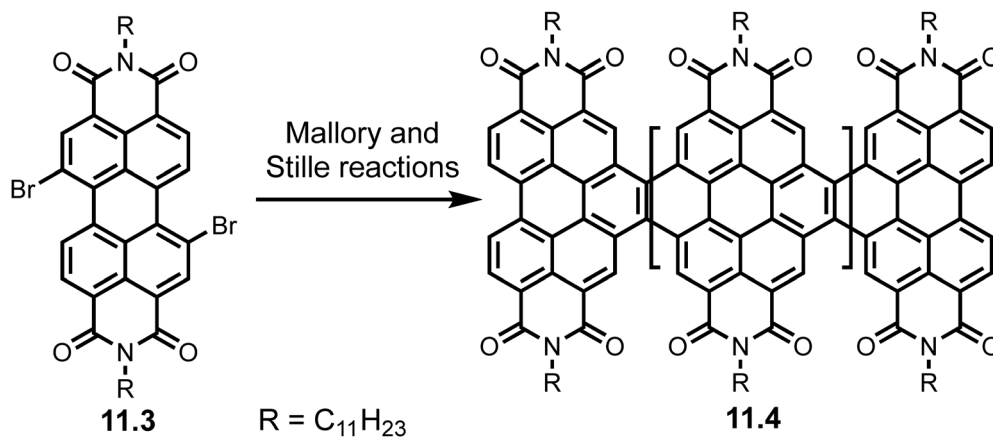
boronated monomers.<sup>43</sup> In contrast, solution-phase synthesis has so far only achieved post-functionalization with boron,<sup>49</sup> highlighting a key challenge: developing monomers with pre-installed boron units to enable precise incorporation into the

GNR backbone. Such strategies would offer fine control over bandgaps and optoelectronic properties, enhancing the potential for device integration. For practical implementation, techniques like SCTP and ROAMP show strong promise,<sup>47,58,76</sup> as



Scheme 21 Synthesis of GNRs from alkyne precursors (a) topochemical polymerization from butadiyne precursors **10.6** and (b) low bandgap AGNR (**11.2**). Reproduced from ref. 83 with permission from [Cell Press], copyright [2016]; Reproduced from ref. 84 with permission from [ACS], copyright [2017].





**Scheme 22** Synthesis of GNRs (**11.4**) using combination of Stille and Mallory reactions of dibromo monomer **11.3**. Reproduced from ref. 86 with permission from [ACS], copyright [2014].

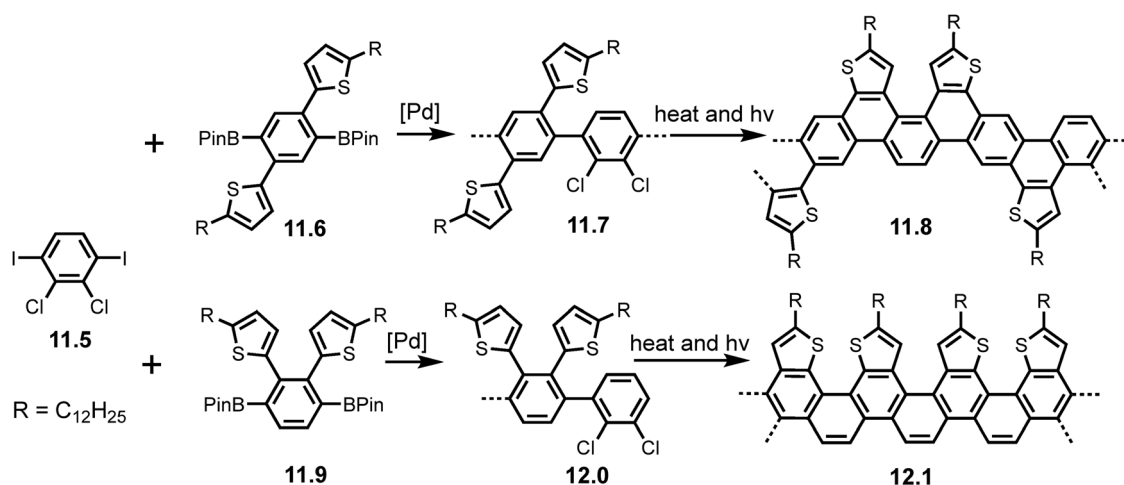
they enable control over GNR length and highly ordered hetero-junction formation—critical parameters for high-performance devices.<sup>88</sup>

One of the main challenges in GNR synthesis remains the ‘graphitization’ step. The Scholl reaction, while widely used and generally high yielding, suffers from significant drawbacks. These include ring rearrangements and unwanted chlorination when using Lewis acid oxidants like  $FeCl_3$ ,<sup>89</sup> as well as degradation of solubilizing groups with  $AlCl_3$ ,<sup>32</sup> resulting in poor solubility and limited post-synthetic characterization. Alternative oxidative conditions, such as DDQ/TFOH, can lead to oxidized/triflated by-products,<sup>32,79</sup> introducing structural defects that negatively impact the intrinsic properties<sup>89</sup> and device performance of the resulting GNRs. Furthermore, for GNRs to be viable at an industrial scale, synthesis methods must deliver gram-scale, defect-free material—something that remains a significant hurdle.<sup>80</sup> Beyond traditional polymerization methods, alternative strategies such as topochemical polymerization using alkyne precursors leverage thermodynamic driving forces from heat or light, offering milder conditions and sustainability benefits.<sup>80</sup>

Notably, photochemical approaches, particularly CDHC, provide a faster and regioselective route for synthesizing GNRs, enabling fine-tuning of structural properties.<sup>80,87</sup>

Furthermore, while GNR mobilities have reached up to  $600 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ ,<sup>51</sup> they remain far below CNTs, which exceed  $50\,000 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$ .<sup>90</sup> This gap stems from various reasons, such as device fabrication, device optimization, contact resistance, defects in GNRs backbone or terminals, GNR aggregation *via*  $\pi$ - $\pi$  stacking, which complicates isolation and hinders reproducible measurements,<sup>91</sup> even with advanced deposition.<sup>92</sup> In contrast, CNTs benefit from defect-free structures and well-established fabrication methods.<sup>93</sup>

Overall, AB Diels–Alder polymerization remains the most promising strategy for GNR synthesis due to its ability to produce long, highly functionalized ribbons which have already led to various device implementation including, molecular diodes, bioimaging, and transistors.<sup>70–74</sup> Whilst other methods remain useful for their respective purposes, SCTP and ROAMP polymerizations also show high potential for device implemented GNRs due to their superior structural control and pathway for milder ‘graphitization’ methods.



**Scheme 23** Synthetic route toward thiophene annulated GNRs (**11.8** and **12.1**) *via* Suzuki polymerization and using CDHC mechanism for aromatization. Reproduced from ref. 87 with permission from [Wiley], copyright [2018].



**Table 1** Summary of the referenced GNRs discussed throughout this review providing information on lengths, molecular weights ( $M_w$  unless stated otherwise), polymerization and graphitization methods. '—' in place where information was not available in original article

GNR	Average length (nm)	Molecular weight of polymer precursor ( $\text{kg mol}^{-1}$ )	Polymerization method	Graphitization method	Ref.
1.3	—	12–120	$\text{A}_2\text{B}_2$ Diels–Alder	$\text{AlCl}_3$	52
1.6	—	62.5	$\text{A}_2\text{B}_2$ Diels–Alder	$\text{FeCl}_3$	53
2.4	71	108.3	$\text{A}_2\text{B}_2$ Diels–Alder	$\text{FeCl}_3$	38
2.8	30	30.4	$\text{A}_2\text{B}_2$ Diels–Alder	$\text{FeCl}_3$	45
3.2	12	13.9	$\text{A}_2\text{B}_2$ Suzuki	$\text{FeCl}_3$	55
3.6–	86	37.4	$\text{A}_2\text{B}_2$ Suzuki	DDQ/TfOH	49
3.8	—	—	—	—	—
4.2	20	33	$\text{A}_2\text{B}_2$ Suzuki	$\text{FeCl}_3$	46
4.6	35	28.8	$\text{A}_2\text{B}_2$ Suzuki	$\text{FeCl}_3$	48
4.9	17	36	AB Suzuki	$\text{FeCl}_3$	37
5.5	14	26.1	AB Suzuki	$\text{FeCl}_3$	47
5.8	—	29.8 ( $M_n$ )	AB Suzuki	Benzannulation	58
5.9	—	—	AB Suzuki	DDQ/TfOH	58
6.2	30	52	AA Yamamoto	$\text{FeCl}_3$	61
6.5	6, 11, 58	15.6–186	AA Yamamoto	$\text{FeCl}_3$	21
6.8	23	20.8	AA Yamamoto	$\text{FeCl}_3$	51
7.1	20	29.8	AA Yamamoto	$\text{FeCl}_3$	50
7.4	—	132.7	AA Yamamoto	DDQ/TfOH	66
7.7	200	24–620	AB Diels–Alder	$\text{FeCl}_3$	54
8.0	390	230–550	AB Diels–Alder	$\text{FeCl}_3$	68
8.3	—	31–54	AB Diels–Alder	$\text{FeCl}_3$	69
8.7	—	161	AB Diels–Alder	$\text{FeCl}_3$	30
8.9	100	—	AB Diels–Alder	—	31
9.0	—	—	AB Diels–Alder	$\text{FeCl}_3$	70
9.1	—	—	AB Diels–Alder	$\text{FeCl}_3$	71
9.2	—	125–278	AB Diels–Alder	$\text{FeCl}_3$	73
9.3	10–371	—	AB Diels–Alder	$\text{FeCl}_3$	74
9.8	—	26 ( $M_n$ )	AB Diels–Alder	$\text{FeCl}_3$	75
10.5	68	14 ( $M_n$ )	ROAMP	$\text{FeCl}_3$	76
10.9	—	—	Topochemical polymerization	Topochemical polymerization	83
11.2	—	—	Topochemical polymerization	Topochemical polymerization	84
11.4	—	—	Stille/Mallory	—	86
11.8	—	9.7 ( $M_n$ )	$\text{A}_2\text{B}_2$ Suzuki	CDHC	87
12.1	—	9.5 ( $M_n$ )	$\text{A}_2\text{B}_2$ Suzuki	CDHC	87

## Conflicts of interest

There are no conflicts to declare.

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