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Russell Kai Liang Tan  
*Iowa State University*, kailiang@iastate.edu

Sean P. Reeves  
*Iowa State University*, spreeves@iastate.edu

Niloofar Hashemi  
*Sharif University of Technology*

Deepak George Thomas  
*Iowa State University*, dgthomas@iastate.edu

Emrah Kavak  
*Iowa State University*

*See next page for additional authors*

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Graphene as Flexible Electrode: Review of Fabrication Approaches

Russell Kai Liang Tan 1#, Sean P. Reeves 1#, Niloofar Hashemi 2, Deepak George Thomas 1, Emrah Kavak 1, Reza Montazami 1, Nicole N. Hashemi 1*

1 Department of Mechanical Engineering, Iowa State University, Ames, IA 50011, USA
2 Department of Materials Science and Engineering, Sharif University of Technology, Tehran, Iran

# R.K.T and S.P.R. contributed equally.
* E-mail: nastaran@iastate.edu

1 Abstract

In recent years, the technological advancement of supercapacitors has been increasing exponentially due to the high demand in electronic consumerism products. As so, researchers have found a way to meet that demand by fabricating graphene. As developments are made toward the future, two big advancements to be made are large-scale fabrication of graphene and fabricating graphene as a flexible electrode. This would allow for use in larger products and would allow for manipulation of the unique properties of graphene to accommodate superior design alternatives. While large scale production is still mentioned, this review is specifically focusing on different methods used to fabricate graphene as a flexible electrode. Various fabrication methods, such as Hummer’s method, chemical vapor deposition, epitaxial growth, and exfoliation of graphite oxide, used to fabricate graphene in such a way that allows flexibility and utilization of graphene’s mechanical and electrical properties are discussed. Additionally, a section on environmentally-friendly fabrication approaches is presented and discussed.
3 Introduction

Since the discovery of graphene and its electrical and mechanical potential over ten years ago, it has been studied and manipulated in the attempt to further technology for the future. Graphene, which are carbon atoms bonded in a honeycomb lattice structure in sheet form with the thickness of one atom, has many factors that go into fabrication, as well as serving as a multi-functional material. It has the capabilities to assist in transistors, electrodes, sensors, and many more applications. The purpose of this study is
to focus on graphene used as an electrode with flexible characteristics. The goal alongside that purpose is for researchers to use this as a resource to assist in their fabrication of graphene, while moving away from past ways that restrict an electrodes potential to be more flexible, more powerful, and significantly smaller. Indium tin oxide (ITO) has been commonly used in applications of transparent electrodes. ITO does not have characteristics to be used as a flexible transparent electrode. This is because of its film brittleness and low infrared transmittance. These characteristics, along with the scarcity of indium and high overall cost prevents the development of future applications (1). Because of graphene’s incredible mechanical and electrical properties, as well as its size, it has great potential to further application development. One of graphene’s outstanding properties is the thermal conductivity (κ) due to the high levels of phonon-dominated on the surface, which can be increased with the switch-on phenomenon and subtracting the defect levels in graphene (2, 3). With research and development reaching a global scale, this review contains compilation and analysis of different fabrication methods and techniques used to fabricate graphene as a flexible electrode. The information provided does not describe the only ways to fabricate graphene as an electrode, but are common methods taken with successful results. In taking the steps toward moving away from past methods and furthering graphene for industrial application, there are a few challenges that are faced. Some of these challenges include large-scale production inability, production costs, and producing it with flexible characteristics. Most synthesis practices continue for a long time and result in high cost and poor scalability in real-world application (4). For example, for optoelectronic and electrochemical devices, inorganic semiconductors are considered for research and exploration of micro- and nanoscale phenomena and learning the importance of size and dimensionality (5). Due to their low work functions, high aspect ratios, and high mechanical stability, the essential nanostructure design of various inorganic semiconductors offers exceptional features for a wide range of systems used in electronic applications. There have been many advances in application for graphene in physics, whereas the application in chemistry, biochemistry and biosensing needs development. The positive environmental impact that graphene can have is significant too. For example, with a constant increase in freshwater resource contamination, graphene can
assist in the desalination of seawater into purified water through capacitive deionization and the graphene mixture used as the positive electrode (6).

Graphene usage for sensors has been reported numerous times in applications such as strain-sensors or efficient human-motion detection, whereas for electrodes, graphene’s usage has mainly been reported on detection of small biologically relevant analytes. Batteries are also a great example of where graphene is becoming extremely beneficial, allowing an alternative to Lithium Ion. With the inspiration of metal-air batteries, which relies on its open structure to generate electricity using oxygen gas, in-situ graphene-like carbon nanosheets adopted a similar strategy (7). It has great potential in point-of-care medical diagnostics, skin-substrate electronics, HMI’s, and many more applications (8). More possibilities include computational framework with terahertz operating speeds for compact all-carbon spin logic circuits and drastically improved in power delay products (9). It also makes for improved use in electronic devices like touch panels, p-n junction materials, wearable technology and hydrogen storage materials (1, 10-12). The possibilities are endless once the fabrication of graphene as electrodes becomes more controllable.

4 Fabrication Methods

There are plenty of ways that graphene can be fabricated, especially as research on graphene’s multi-use capabilities reach various applications at an exponential rate. For the case of graphene fabrication as a flexible electrode, there are four methods that are commonly used today. These methods are the modified Hummer’s method, exfoliation of graphite oxide, chemical vapor deposition, and epitaxial growth. Each fabrication method can be approached from different angles. These approaches may include, but are not limited to, hybrid solutions, using different elements, or manipulating material mass or volume. Other methods to make notice of are facile photochemical strategies to rapidly construct porous graphene frameworks and template methods using a triblock copolymer as a template for fabrication (13).
Table 1: A summary of graphene fabrication property outcomes. ✓ denotes that the fabrication method provides this property. ✗ denotes that the fabrication method does not provide this property.

<table>
<thead>
<tr>
<th></th>
<th>Electrical Conductivity</th>
<th>Strength</th>
<th>Elasticity</th>
<th>Large Surface Areas</th>
<th>Chemically Inert</th>
<th>Transparency</th>
<th>Short Fabrication Time</th>
<th>Large Output Volume</th>
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<tbody>
<tr>
<td>Hummers Method</td>
<td>✓</td>
<td>✓</td>
<td>✗</td>
<td>✗</td>
<td>✓</td>
<td>✓</td>
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</tr>
<tr>
<td>Exfoliation of Graphite Oxide</td>
<td>✓</td>
<td>✓</td>
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<td>✗</td>
<td>✓</td>
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<td>CVD</td>
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<tr>
<td>Epitaxial Growth</td>
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4.1 Modified Hummer’s Method

Hummer’s Method is a very common method in the process of fabricating graphene. Graphite Oxide is created mainly by the chemical oxidation of graphite. It is originally synthesized by an addition of potassium chlorate to a graphite/HNO₃/H₂NO₃ mix (14). These are known as the Staudenmaier and Brodie methods. Hummer’s method was later developed, in which KMnO₄ and NaNO₃ were dissolved in concentrated H₂SO₄ to allow for a more effective oxidation for the graphite to produce graphite oxide. The cons with both the Staudenmaier and Brodie methods were the long turnover rate of 4 days, and the consumption of toxic and harmful gases (15, 16). The difference between the two methods is that Hummer’s method only takes a few hours.

Modifications have been made to the Hummer’s method to address problems that have risen, like adding a peroxidation step, or removing NaNO₃ from the method and finding a replacing substance due to its toxic gas generation. Each version of Hummer’s method was modified in different ways and it depends on the user as to what modifications can be made. To fabricate graphene, Hummer’s method usually
takes place since the graphite must be oxidized (unless a unique direction is taken like the R2R approach explained later).

One example of a general Hummer’s method consists of graphite powder stirred and chilled with H₂SO₄ and NaNO₃ (17). During stirring, KMnO₄ was added in aliquots over 2 hours, followed by room temperature cooling after the stirring process is completed. Then, 35 °C was applied for 30 minutes, followed by adding the solution into a flask with deionized water. The temperature was again increased to 70 °C, and the time to 15 min. The new solution was added to a larger volume of deionized water, where H₂O₂ was added to remove the KMnO₄. Deionized water repetition was necessary until a neutralized pH was achieved. GO was then present and was dried in a vacuum for 2 days at 60 °C.

Like previously mentioned, each Hummers method could be modified in certain ways. An example of a modified Hummers method is as follows.

Firstly, a PANI/GN composite was acquired. To do that, an aqueous GN dispersion (1.0 mg mL⁻²) was prepared using a modified Hummers method. A solution of 0.4 mL aniline monomer (solvent: 2 M HCl, 25 mL) was dropped into the as-prepared GN suspension (25 mL); afterwards the mixture was sonicated for 0.5 hours, mixed with HCOOH and then strongly mechanically stirred for 0.5 hours. Subsequently, 50 mL of 240 mg ammonium persulfate (solvent: 1 M HCl) was added dropwise into the above mixture and stirred in an ice-water bath for 2 hours to form a PANI/GN suspension. The next step was to synthesize PANI/GN to BC to form a film like electrode. Firstly, the gel-like white BC pellicles were washed thoroughly with deionized (DI) water, then cut into small slices, and subsequently pulped with a mechanical homogenizer at the speed of 10,000 r min⁻¹ to obtain the BC suspension (0.7 mg mL⁻¹). The resultant BC suspension (200 mL) was drained on a nitrocellulose filter membrane (porous size of 0.22 μm) to get a uniform BC paper by vacuum filtration. Next, the obtained PANI/GN dispersion was decanted onto the prepared BC paper to form a flexible film, and the fabricated film was washed several times with DI water. After the wet film was dried at 55 °C for 8 hours and then peeled off to get the freestanding PANI/GN/BC paper. Three different abbreviations, namely PANI.(L/M/H)/GN/BC represented the various concentrations of PANI for 0.2, 0.3
and 0.4 mL. An in-situ polymerization and vacuum filtration strategy was employed to coat a uniform layer of PANI onto the GN surface, followed by pouring into BC paper to form the flexible and freestanding film. Since the abundant groups of BC and its excellent physical properties, PANI/GN composites could strongly bind with the BC paper via hydrogen bonding and electrostatic interaction. The mixture was fully immersed and utilized the porous network of BC to obtain large mass loadings (18, 19).

A high demand for on-body wearable medical apparatus and implantable devices also used Hummers method in its experiment (20). The research described a flexible graphene electrode based on graphene paper (GP) supported 3D monolithic nanoporous gold (NPG) scaffold (NPG/GP), with modifications made to provide a high density, well dispersed, ultrafine binary PtCo alloy nanoparticles through ultrasonic deposition. The first step was to fabricate a freestanding graphene oxide (GO) via modified Hummer’s Method, which was included in the author’s recent works (21). Freestanding GO paper was fabricated starting with dropping a well dispersed GO solution (5mg mL-1) onto commercial printing paper surface (A4). The GO solution was then spread uniformly across the commercial printing paper via rolling multiple times. The sample was allowed to air dry to obtain a uniform GO paper. A reduction process for GO paper was carried out by soaking it in HI solution (45 wt%) for 10 minutes at room temperature. During this process, bubbles emitted from the cellulose papers allowed the thin film to peel by itself. Then, the rGO paper was rinsed with a saturated sodium bicarbonate solution, water and methanol followed by vacuum drying at 80°C. The rGO paper was then coated with a layer of PANI via in situ polymerization. Generally, the PANI nanofiber network was synthesized with a two-step method on a three-electrode system in a 1M HCl electrolyte and 0.3 M aniline monomer. Nucleation of PANI was performed with a constant feed of voltage at 0.8 V for 1 minute at room temperature. The following step allowed nanowires to grow under a constant current condition with current density of 2 mA cm-2. The resultant PANI layer was then covered with an ultrathin GO film through dip-coating method. The last step was to take the aforementioned
sample and soak it in HI solution for a few minutes at room temperature to produce rGO/PANI/rGO paper. The next step was to fabricate freestanding NPG/GP by immersing a piece of Ag-Au leaf (1 cm x 1 cm) into concentrated nitric acid to float the leaf at air-liquid interface. After the leaf was observed to float freely on the solution surface, the glass slide was removed. Then, the leaf was dealloyed for 6 hours in the dark at 30 C. Subsequently, the leaf was removed from the acidic solution using a glass slide and rinsed with deionized water by allowing it to float for three times (15 minutes for one time). The NPG film was then transferred to a freestanding GP, and 10 microL Nafion solution was dropped on the surface of the NPG to obtain a freestanding GP supported NPG (NPG/GP). To get a freestanding PtCo/NPG/GP, the NPG/GP was allowed to undergo PtCo alloy nanoparticles electrochemical deposition under ultrasonic irradiation (45 W) (22). The aforementioned process was performed in 0.2 M of Na2SO4 containing 5 mM H2PtCl6·6H2O and 15 mM CoCl2·6H2O using cyclic voltammetry technique at a range of -1.0 to 0.0 V and scan rate of 250 mV s⁻¹. The resultant nanohybrid paper electrodes was then cleaned carefully with distilled water and dried at room temperature.

Aerogels and graphene were both interesting materials that were highly sought after for research purposes. One experiment incorporated graphene aerogels into low-cost 3D N-doped biomass-derived carbon framework as a potential flexible energy storage (23). The 3D structural framework potentially boosts the energy storage device with superb electrical conductivity, chemical stability and flexibility (24, 25). Firstly, GO was fabricated using modified Hummer’s method. 2 g of graphite was added into 120 mL of sulfuric acid (98%) in an ice bath and cooled to 0 °C under stirring. Then, 1 g of NaNO3 (AR) was added and stirred for 30 minutes. Subsequently, 6 g of KMnO4 (AR) was slowly added to the solution and stirred for 2 hours to maintain the temperature below 20 C. Next, 150 mL of deionized water was added, followed by 200 mL of H2O2 (38%) dropwise. The resulting mixture was then washed with deionized water and HCl (5%) to obtain an aqueous GO dispersion. The next step was to synthesize N-doping cotton-derived carbon frameworks (NCCF). In this step, the
use of raw cotton as-is condition was sufficient without any pre-treatment. Urea was mixed with a piece of cotton (mass ratio of 4:1) in a quartz boat, followed by carbonizing at 800 °C for 1 hour at the rate of 5 °C min-1 in N2 atmosphere. The high temperature N2 atmosphere allowed the decomposition of urea to react with the carbonized NCCF. The resulting product was further treated with ultrasonication for 1 hour under the presence of HNO3/H2SO4 (v/v = 1/3). The finalized sample was then cleaned with deionized water and dried at 90 °C overnight. The final step was to synthesize NCCF-rGO. A piece of NCCF was cut from the acid treated cotton according to its natural texture. Next, Cu foil and NCCF piece was immersed into acidified GO solution (3 mg mL-1, 0.001 M HCl) to obtain the gel of NCCF. The thin piece of NCCF was to be strongly bound to the Cu foil for 6 hours, allowing interfacial gelation of GO to be self-assembled into hydrogels of the Cu foils. After that, the resulting sample was carefully cleaned with deionized water to remove the unreacted GO solution. After the aforementioned step, the sample was immersed into 10 fold diluted HCl solution to detach the gel of the NCCF from Cu foil followed by washing it with 20 fold diluted aqueous HCl and deionized water respectively to remove any remaining chemical residues. Subsequently, the sample was freeze-dried to obtain aerogel-like dried product, followed by annealing it at 300 °C for 1 hour under N2 atmosphere.

Figure 1: Preparation of 3D NCCF-rGO electrodes via facile synchronous reduction and assembly process (23).
The usage of polypyrrole (PPy) was shown by how researchers experimented on fabricating hybrid paper electrode with rGO (using NaBH4), in situ polymerization of PPy and cellulose (26). The experimenters used in situ polymerization of PPy and chemical reduction with NaBH4 to reduce graphene oxide. Paper proved to be an ideal base for flexible supercapacitor fabrication, since paper could easily synergize with a wide range of electrochemically active substance such as electronically conducting polymers (ECPs) (27-31). Polypyrrole (PPy), polyaniline (PANI) and poly-(3,4-ethylenedioxythiophene) (PEDOT) were examples of ECPs that possessed high pseudocapacitance, quicker redox switching, light weight, high conductivity and low costs crowning these materials as high tier class materials for high-performance supercapacitors (32-37). The first part of fabricating PPy/cellulose papers required the base cellulose paper with dimensions roughly at 40 mm x 40 mm to be soaked in four different concentrations of pyrrole monomer at 0.05, 0.15, 0.45 and 1.35 M for 2 hours. Subsequently, an aqueous solution of iron (III) chloride hexahydrate (FeCl3·6H2O) was added slowly into each individual concentrations of pyrrole monomer to act as an oxidant for in situ polymerization process for 3 hours. The molar ratios of the ferric iron/pyrrole are 0.1, 0.3, 1 and 3 respectively. The resulting 16 PPy/cellulose papers were placed into 1 M HCl for 30 minutes and then rinsed with distilled water, followed by drying in a vacuum oven at 50 °C for 24 hours. The next step was to fabricate GO through modified Hummer’s method. 2 g of graphite powder was mixed into 80 mL of sulphuric acid and 4 g of sodium nitrate, under magnetic stirring in an ice bath. 8 g of Potassium permanganate was added to the resulting suspension under intense agitation. The temperature of the mixture was then brought down to 0 °C for 3 hours, followed by stirring at 40°C for 2 hours. Then, 200 mL of deionized water was added into the resultant mixture, followed by its heating to 98°C for 30 minutes. Then, 30 mL of H2O2 (30%) was added into the solution, followed by rinsing it with 200 mL of 5 wt% HCl at deionized water. The final product would be graphite oxide, which would be immediately subjected to a facile ultrasonication treatment that exfoliated the graphite oxide to GO. To fabricate rGO/PPy/cellulose (RPC) papers, PPy/cellulose papers with the lowest sheet resistance were picked and separated in three different concentrations of GO
dispersions (0.1, 0.5 and 2.5 g L-1). GO dispersions was prepared under ultrasonication for one hour (500 W) in an ice bath. The resulting GO/PPy/cellulose papers was reduced with 0.15M of aqueous NaBH4 solution, then stirred for 12 hours. The resulting RPC papers were then cleaned with deionized water to remove any impurities on the surface, followed by drying at 50°C for 24 hours. The assembly of the all-solid-state flexible laminated symmetric supercapacitor (SSC) is fairly simple, involving just pressing two RPC papers dipped in H3PO4/PVA gel electrolyte together followed by transferring it to a fume hood at room temperature to allow excess water to evaporate. The aforementioned gel electrolyte was prepared by adding 6 g of H3PO4 and 6 g of PVA in 60 mL of deionized water under magnetic stirring at 85 °C until the solution becomes transparent.

In an experiment that investigated a graphene/polyaniline@carbon cloth composite as a supercapacitor via a simple one-step electrochemical co-deposition process, it was realized that graphene oxide to graphene/polyaniline composite must be prepared to make the supercapacitor achievable (38). The synthesis of GO through the oxidation of graphite flakes was done via modified Hummer’s method. The resulting GO dispersion was placed in a centrifuge at 4000 rpm for 30 minutes to remove remaining aggregates followed by dialysis to purify salt impurities for 1 week. The GO dispersion was then adjusted to 0.36 mg mL-1 for later experiments. To remove any grease and impurities, a carbon cloth (1 cm x 2 cm) was soaked in acetone and distilled water followed by sonication for 30 minutes. The aforementioned process was repeated several times followed by drying process in vacuum at 60 C. Then, a transparent solution was prepared by adding 135 microl of aniline into 10 ml of distilled water and sonicating it until transparent in room temperature. The electrolyte for electrochemical co-deposition was created by adding 1 ml of 1.0 M Na2SO4 and the previous sonicated solution into 15 ml of GO dispersion, to be further sonicated again for 10 minutes. A three-electrode system was conducted using carbon cloth, Pt foil and Ag/Agcl as the working electrode, counter electrode and reference electrode respectively. Three different samples needed a slight modification in settings for cyclic voltammetry (CV) cycles, namely GP@cc (the sample before this), PANI@cc
and rGO@cc with voltage ranges at -1.2 to 0.8 V, 0 to 0.8 V and -1.2 to 0 V, respectively. All samples used a scan rate of 2 mV s-1. After each cycle, the samples were cleaned with distilled water and repeated multiple times to remove the absorbed monomer or salts.

One team (39) experimented on fusing graphene and Polyaniline (PANI) composite film together to further study about graphene and nanostructured pseudocapacitance material (40, 41). Graphene is known to have a stable but low capacitance due to graphene adopting the electric double-layer capacitor mechanism that aids in energy storage (42-44). PANI is well known to be a good candidate for supercapacitor with pseudocapacitance materials because of characteristics such as environmental stability, doping/dedoping chemistry and a special conducting mechanism (45-47). Preparation of materials for this experiment is simple, all other reagents mentioned would be AR grade and aniline to be distilled before use. GO was obtained via modified Hummer’s method. The first step was to fabricate freestanding 3D-rGO film. GO solid was dispersed with ultrasonication for 100 minutes and centrifuged at 400 rpm to obtain approximately 1.25 mg mL-1 of GO dispersion. Subsequently, 0.4 mol of CaCl2 and 2.5 mL NH3. H2O was added to the GO dispersion respectively after CaCl2 was dissolved completely. The mixture was stirred for 10 minutes and then placed in a CO2 atmosphere at a rate of 1 L min-1. The mixture was allowed to react for 80 minutes, then left to rest overnight. Then, one third of the resultant suspension was allowed to undergo vacuum filtration to obtain a GO-CaCO3 film. The resultant film was then reduced in hydrazine vapor at 40 °C for 12 hours, cleaned with diluted hydrochloric acid to remove CaCO3 particles on the surface and finally washed with deionized water and ethanol to get freestanding graphene film. The process to prepare 3D-rGO/PANI composite film allows 3D-rGO/PANI nanocomposites to synthesize with 3D-rGO film via dilute polymerization. First, 20 mL of 1 mol L-1 aqueous HClO4 solution was prepared and poured into a reaction vessel within an ice bath. 5 mL of ethanol was added to the 3D-rGO film and placed next to the reaction vessel to improve the wettability of the film. After the solution cooled down, aniline monomer was added and stirred for 10 minutes to obtain a
homogeneous mixture. (NH₄)₂S₂O₈ (APS) was prepared in 15 mL aqueous HClO₄ solution and precooled (ratio of aniline/APS = 1.5) to act as the oxidant. The oxidant was then added into the monomer solution. The polymerization process started and was sustained for 24 hours in an ice bath. A composite film was obtained by cleaning the sample with deionized water and ethanol, then drying it in room temperature.

Another research team studied reduced graphene oxide based off fiber springs to enhance the flexibility and regenerative capabilities of a supercapacitor (48). Stretchable electronic devices recently have been a strong interest in the modern electronics market (49-55). Fiber was incorporated into the supercapacitor as it plays an important role to help boost structural strength that makes it flexible, lightweight and soft at a low cost (56-58). The combination of fiber and graphene creates graphene fibers, which showcases both high thermal and electric conductivity, superb stiffness and elasticity, and strong stability (59-62). In practical applications, the graphene fibers would undergo high-stressed environments that pushes the reliability and stability of the supercapacitor. Therefore, the graphene fibers could potentially suffer from mechanical damage, and eventually breakdown (63, 64). Self-healing was applied to the graphene fiber supercapacitor to heal damaged fibers, allowing it to regain mechanical and structural properties (65). Not only can those materials heal
and restore electrical and structural characteristics of mechanically damaged supercapacitors, it can also prevent structural fractures from happening (66-68). GO was prepared with oxidized graphite powder using modified Hummer’s method. Multi-walled Carbon nanotubes (MWCNT) and sodium dodecyl sulfate (SDS) was added into 5 mg/ml of aqueous GO solution with a ratio of GO/MWCNTs of 2:1, 1:1 and 1:1.5. The mixture was ultrasonicated to allow even dispersion, followed by addition of Vitamin C particles and ultrasonicated once more for 5 minutes. The resulting mixture was injected into 1.5 mm diameter pipes and sealed both ends with polydimethylsiloxane (PDMS). GO was reduced after heating in an oven at 90 °C for 6 hours and the remaining wet fibers were dried. Partially dried rGO/MWCNT fibers were the base for electrodeposition of PPy according to literature in 1, 1.5 and 2 mins (69). Polypyrrole (PPy) was deposited to rGO/MWCNT fibers using 5% (v/v) pyrrole monomer, 0.2 M NaClO4 with a constant voltage of 0.8 V. PPy/rGO/MWCNT fibers were physically modified into a spring shaped structure, then followed by coating with PVA-H3PO4 gel electrolyte and dried under vacuum at room temperature. Then, the springs were placed parallel and coated with the same electrolyte before and dried under vacuum. The supercapacitor was finally coated with polyurethane (PU) and dried in air.
Figure 3: Schematic diagram of the highly stretchable and healable fabrication process. a) The twisting process of rGO fibers into springs. b) Preparation procedures of the rGO/MWCNT PPY-decorated supercapacitors (48).

Using other elements to promote supercapacitor performance is a good approach (70). This group discussed on utilizing nickel cobalt sulfide to promote supercapacitor performance by uniformly dispersing the ncs on the surface of graphene via one-step solvothermal method with poly(acrylic acid) (PAA) additive. Nickel-cobalt sulfides are binary metal sulfides from transition-metal sulphides (TMSs) group was recently considered for their rich redox reactions compared to both single component sulfides (NiSx or CoSx) (71). Downsides of Nickel-cobalt sulfide reported were insufficient capacitance under high rate because of the disadvantageous ion transportation length and low electronic conductivity (72, 73), and poor electrochemical stability from long-term cyclic that creates mechanical stress and strain that eventually causes a large change in volume and structural failure (74, 75). The combination of Nickel-Colbalt sulfide nanoparticles on graphene (Ni-Co-S@G) delivers outstanding energy densities without sacrificing capacitance or electrochemical stability, since a
simultaneous utilization of the active material and contented conductivity can be achieved (76-78). All chemicals used in this experiment were AR grade. GO was synthesized with modified Hummer’s method to obtain a GO suspension. The one-step solvothermal synthesis method was used to fabricate Ni-Co-S@G nanocomposite. Ni(CH₃COO)₂·4H₂O, Co(CH₃COO)₂·4H₂O and PAA (Mw=~2000, Macklin) with molar ratio of 1:2:2:8 was dissolved in 35 ml 0.5 mg ml⁻¹ GO suspension in ethylene glycol (EG) at 80 °C under vigorous stirring for 2 hours. Then, thiourea was added to the solution to undergo hydrolysis. The resulting solution was transferred into a 50 ml Teflon-lined stainless steel autoclave at 200 °C for 10 hours. The solution was placed in a centrifuge at 5000 rpm to remove the undecorated particles. The remaining samples were cleaned with deionized water and ethanol for multiple times. Ni-Co-S@G/ACC electrode was fabricated the same way as Ni-Co-S@G except adding a piece of ACC into Teflon-lined stainless steel autoclave before solvothermal.

Figure 4: Schematic fabrication process detailing one-step solvothermo method to obtain Ni-Co-S@G and Ni-Co-S@G/ACC electrodes as (Strategy I) and (Strategy II), respectively (70).

One incredibly influential role graphene plays is its ability to act as an electrode in capacitive deionization experiments. Hummers method can assist in the fabrication of this electrode. One example described GO prepared by the Hummers’ method and the powder then ultrasonicated into 3 mg/mL DI (79). To prepare what was termed GO@PS microspheres using polystyrene, 100mL of the mix was placed into a prepared aqueous suspension of 50 mL PS
microspheres under magnetic stirring. Ultrasonication was carried out for 2-3 hours after. Next, vacuum filtration with a millipore filter was conducted with the resulting mix. The spheres and sheets were peeled from the filter and dried at degrees C. The resulting composites were annealed at 550 degrees C for an hour. Next, under N₂, the same was done at 900 degrees C and a heating rate of 1 degree C/min. This reduces the GO into graphene and the PS spheres are taken away. The final resultant material is termed 3DMGA. As expected, there are many other ways to fabricate graphene as an electrode for the improvement of capacitive deionization, including sulfonic and amine function groups on 3D graphene, as well as dual-template prepared carbon composites (80, 81).

An example discussed on the usage of graphene-coated hollow mesoporous carbon spheres (GHMCSs) as electrodes for capacitive deionization (CDI). A template-directed method was used to fabricate GHMCSs, where phenolic polymer coated polystyrene spheres were used as templates. As a result, the graphene composite formed a hierarchically porous nanostructure, with individual hollow mesoporous carbon spheres uniformly distributed across graphene sheets. To fabricate GHMCSs, phenolic polymer coated polystyrene spheres (PF@PS) was synthesized. PS spheres procedures were previously reported with a diameter of 180 nm (82). 150 mg of PS spheres were dispersed into 50 mL of deionized water, followed by 20 mL of aqueous phenol solution (0.188 g) and 10 mL of hexa-methylenetetramine (0.14 g). The solution was gently stirring for 15 minutes and transferred into 100 mL Teflon-lined autoclave at 160 C for 4 hours. PS@PF spheres was obtained after centrifugation and cleaned with deionized water and pure ethanol. The next procedure was to synthesize GO with PF@PS nanospheres. GO was prepared via Hummer’s method. A 3 wt% aqueous suspension of the PF@PS nanospheres were obtained by re-dispersing them into deionized water. 20 mL of GO dispersion was added into 6g of aqueous suspension of PF@PS under magnetic stirring conditions. The solution was ultrasonicated for 1-2 hours to obtain a homogeneous colloidal suspension of PF@PS and GO hydrosol. The assembly of PF@PS nanospheres and GO sheets was achieved by vacuum filtration using a milipore filter. The resulting GO-PF@PS composite from vacuum filtration was peeled off from the filter and air-dried overnight at 40 C. Then, the composite was annealed in a tunular furnace at 150 C for 1 hour, followed by 900 C for 2 hours under N₂ atmosphere at a heating rate of 1 C min⁻¹. GHMCSs was obtained from the thermally reduced composite film of
GO and carbonized PF, while PS microspheres was removed and HMCs was simultaneously formed.

![Figure 5: An illustration of the fabrication process of graphene coated hollow mesoporous carbon spheres (GHMCSs) (83).](image)

**4.2 Exfoliation of Graphite Oxide**

Exfoliation is among the four most common ways to fabricate graphene and can be broken down into many different types. The general method is carried out just as the title describes, it is the exfoliation of graphite layers. Mechanical Exfoliation using scotch tape was the method used to fabricate the first ever sheet of graphene, but lacks the potential for large scale production within a time frame. A contradicting example consists of an economical method to produce high-quality graphene films similar to the scotch tape method in a large scale via a three-roll mill exfoliation (84). One big difference with this study is the ability to continuously exfoliate graphite using the three-roll mill system, with the first and third drum rolling at the same direction and the second drum’s rolling reversed. Rolling at a constant velocity, the rolled graphite will run in an inverted S curve. An adhesive was first prepared by dispersing 2.0 g of polyvinylchloride (PVC) into 50 ml of dioctylphthalate (DOP) in a magnetic stirrer at 250 °C for 30 minutes. The resulting adhesive was then carefully placed in between the feed and the center of the rolls. 1.0 g of pre-dried natural graphite at 100 °C for 24 hours was carefully dispersed on the adhesive after the rolls started to move to attain maximum surface contact. After exfoliation process has been carried out for 12 hours, the resulting materials was collected and steeped in alcohol to remove DOP from the product. After that, the sample was placed in the muffle at
500 °C for 2 hours to burn away the PVC resin layer to obtain a purely graphene product.

Figure 6: Depiction of the exfoliation of natural graphite with the three-roll mill system (84).

One big problem with graphene is that graphene sheets tend to stack on top of each other, which causes a decrease in surface area. 2D and 3D porous graphene architectures were fabricated using KOH and CO2 activation with EGO (exfoliated graphene oxide). All materials in this experiment were used without purification. GO was synthesized by modified Hummer’s method (85). EGO was prepared by placing 1g of dried GO in a quartz tube and purging with N2 for a half hour to remove excess air. The tube was relocated to a vertical furnace at 800 °C for thermal exfoliation and was allowed time for it to cool to room temperature under N2 flow. To create KOH-activated graphene, the chemical activation of EGO was used utilizing KOH as an activating agent. Dry EGO was mixed with deionized water and KOH was added. The mixture required 12 hours of stirring, and was then vacuum-dried at 65 °C. The resulting gray solid was then added to a crucible, where a N2 glow was added to the tube for 1 hour to get rid of the air. Next, there would be a steady temperature increase to 800 °C. The product was then washed with deionized water, and then vacuum-dried marking the end of the KOH experiment. The product behaves differently based on the mass of KOH and EGO used. For the CO2-activated
graphene, the process remained the same until the N2 flow. After the N2 stream, the temperature was increased and the gas stream was changed to CO2 with no change of other variables. After CO2 activation, it was cooled to room temperature under N2. The final product left a black spongey powder. The product was time dependent and not mass dependent. With fluctuation in the KOH/EGO mass ratio or in the time of CO2 activation, activated graphene materials with a high specific surface area of 2518 m$^2$/g and a high specific capacitance of 261 F/g would be obtained. Key factors that influence capacitance would include microporosity, specific surface, and surface wettability. A brief example of fabricating graphene using exfoliation paired with a template method for the use of capacitive deionization is as follows. Since chemical or thermal reduction of graphene exhibits lower electro-absorptive capacity from agglomeration, standard exfoliation was carried out to obtain graphene. The graphene was combined with microporous carbon (GE/MC) using a template method. This template method consists of a triblock copolymer F127 as a template and resol as a carbon precursor by a direct triblock-copolymer-template method. The resultant composites showed the mesoporous structure to be well structured with the addition of graphene. Specific surface area, pore size distribution, and pore volume were all improved through this method (13).

In this example, the author discussed about interconnecting carbon fibers (CFs) in carbon cloth (CC) via In-situ electrochemically exfoliated graphene. Carbon cloth are mechanically flexible woven carbon fibers, which are popular for their electrical conductivity, flexibility, chemical stability and porosity. The issue presented on carbon cloth exhibits a very low surface area due to micro-sized gaps between the fibers during production. By interconnecting CFs, it showed significant amount of improvement in specific capacitance, surface area and conductivity (86).

4.3 Chemical Vapor Deposition

Chemical vapor deposition (CVD) is a fabrication method used to produce high-quality thin filmed graphene, while maximizing its performance. It involves heating materials, in this case carbon, to a vapor and/or reducing the atmospheric pressure to coat a substrate. This substrate is a metal to act as a catalyst in the graphene growth
It is the most commonly used fabrication method for graphene. The process is quite simple and has the potential to coat large surface areas. Electrochemical deposition, is also used for graphene fabrication, but is not nearly as commonly used since it may require a multi-electrode setup and because CVD is much more simple (88).

This report described the production and fabrication methods used for devices that utilizes graphene, as well as the flexible electronic devices that are enabled by graphene (89). Both chemical vapor deposition (CVD) combined with transfer techniques and chemical exfoliation (CE) have advantages for application on substrates like wafers, polymers, textiles, and even human skin. The CVD approach is known to grow large-area and high-quality graphene films using a carbon gas source. Common metals include Ni, Fe, Co, Pt, and Cu. To grow graphene film, there are 3 main stages: the diffusion of carbon into the metal film at a specific temperature, the stripping of carbon out of the thin metal film during the cooling process due to the reduction in solubility, and the formation of graphene layers on the surface. A relationship was found that a relatively high solubility of carbon atoms is projected to produce thick graphene films, as well as the other way around, having low solubility produce monolayer films. Other CVD methods like rapid thermal CVD method involves hydrogen-free, low-temperature conditions, which enables faster and larger production of films over a 400mm x 300mm surface area. Regardless of the method, substrate pre-treatments, polishing, and annealing are crucial for suppressing nucleation site density and even producing monolayer graphene. Another important factor for flexible electronic devices is graphene patterning techniques (direct patterned growth from a metal substrate). Another method called plasma enhanced chemical vapor deposition (PECVD) has benefits in contrast to CVD, like direct growth on a plastic substrate or nanostructures. However, CVD is still more effective, especially at high temperatures. This report includes more detailed methods and transfer methods.
An experimental example utilized flexible transparent graphene electrode to improve the performance of perovskite solar cells due to the increase in demand to use perovskite solar cells as reliable portable power source (4). CVD-grown graphene transfer method and copper foil was used to create a single layer graphene-deposited PEN substrates. The CVD grown single layer thick graphene film was deposited on the copper foil, forming a graphene/copper foil composite. PMMA (poly(methylmethacrylate)) was spin coated on the graphene/copper foil composite and heated on a hot plate at 180 °C for 1 minute. The foil was then etched after placing it in an aqueous solution of ammonium persulfate (H8N2O8S2). The resulting sample was placed on a UV-zone treated PEN-film and submerged in acetone solution to remove the PMMA. The graphene/PEN sample was then dried in a nitrogen blower. To increase the electrical contact, each sample of Cr, Au and Al was deposited consecutively through a rectangular metal aperture mask with vacuum thermal evaporator. A MoO3 layer was deposited at a rate of 0.1 A dot s-1 to adjust the surface properties. The sample was then annealed at 110 °C for 10 minutes and cleaned with acetone, isopropyl alcohol and deionized water for 15 minutes respectively, followed by UV-ozone treatment for 20 minutes. While during the whole fabrication process, it is crucial to maintain the flatness of the flexible substrate to ensure the properties of the highly efficient flexible solar cells. Glass/PDMS (polydimethylsiloxane) was used as a rigid support to achieve constant flatness. The graphene/PEN substrate or ITO was placed on the support for the whole duration of the fabrication process. Once the process was completed, the device was peeled off from the support to be characterized. Perovskite solar cells utilizing transparent graphene as a conducting anode produced a highly reliable and efficient results compared to the ITO counterparts.

Another research team looked into the utilization of graphene foam electrodes loaded with sulfur (90). Generic Lithium Ion batteries (LIBs) at its best was unable to meet high energy density and long life applications in the near future (91). Although Lithium Sulphur batteries (LiS) was a strong contender for the next generation of batteries with much higher capacity and energy density with a two-electron reaction,
sulfur loading in the cathode is inadequate (92). Utilizing graphene foam ties the robust properties of graphene and porous network for high sulfur loading brings flexible Li-S battery electrodes. Ni foam grown graphene via CVD was reported in previous works (93, 94). Briefly, the Ni foam was heated up in a horizontal tube furnace at 1000 °C with H2 atmosphere and annealed for 15 minutes to clean and eliminate the thin oxide layer on the surface. CH4, with a concentration of 29 vol% in all total gas flow was introduced into the reaction tube with H2 and Ar at ambient room pressure for 30 minutes. Subsequently, the resulting sample was cooled to 100 degrees C/min under Ar and H2 atmosphere. After the CVD growth process, a thin layer of graphene/poly(dimethylsiloxane) (PDMS) was coated onto the graphene surface on Ni foam by dipping it into a dilute PDMS solution for 30 minutes. The sample was then cured at 80 °C for 4 hours. The Ni substrate layer was etched away by placing it in a 3 M HCl solution at 80 °C for 12 hours to obtain PDMS/GF composite. A homogenous sulfur slurry was prepared by mixing sulfur and conductive carbon black at 10 wt% PVDF acting as a binder dissolved in N-methyl-2-pyrrolidone (NMP) to produce PDMS/GF electrode. The slurry was carefully casted into the PDMS/GF composite with a controlled sulfur loading to obtain S-PDMS/GF electrode. The electrodes were ready to be used after drying it in a vacuum drying oven at 60 °C for 24 hours.
Figure 7: Schematic of the synthesis procedure to fabricate PDMS/GF and S-PDMS/GF electrodes (90).

An example of a simple fabrication method consists of sandwich-like carbon nanotubes (CNTs)/NiCo2O4 hybrid highly conductive paper (95). The metal compound, spinel nickel cobaltite (NiCo2O4), showcased to have ultra-high theoretical capacitance (96), better electrical conductivity and variety of redox reactions (97). NiCo2O4 is also environmental friendly, low cost and its natural abundance makes it a very attractive for usage with electrode applications (98). Within consist of a layer of conductive CNT buckypaper to house honeycomb-like NiCo2O4 nanosheets and to provide strength that makes it flexible. CVD method using ferrocene and dichlorobenzene as precursors was the method to synthesize CNT buckypaper (99). CNT buckypaper was sliced into approximately 0.9 x 0.7 cm2 rectangular pieces and treated with UV/O3 for 2 hours. Precursor solution was prepared by dissolving 4.17 mM Co(NO3)2·6H2O, 2.06 mM Ni(NO3)2 . 6H2O, and 6.25 mM urea in 30 ml methanol. A piece of as-prepared CNT buckypaper was placed into the precursor solution, followed by sealing the mixture in 50 ml Teflon-lined autoclave. The mixture was hydrothermally treated at 120 °C for about 1 to 6 hours for the fabrication of the composites. After the sample was cooled, it was cleaned with deionized water repeatedly, followed by freeze-drying and annealed in air at 320 °C for 2 hours to obtain CNTs/NiCo2O4 paper-like electrode.
One report discussed about reduced graphene sheet (GR) patched carbon nanotubes (CNT)/MnO2 as a suitable pseudo-capacitive electrode material as those materials which includes conductive polymer and transitional metal oxides has shown to have a much larger electrochemical capacitance and electrical densities (100-102). MnO2 alone possesses the characteristics of high-performance supercapacitors such as a wide potential range, environmental friendly and low cost (103), but lacks low conductivity and the flaking off phenomenon from the cyclic crystal shrinkage/expansion (104). Multiwalled CNTs (MWCNTs) was first synthesized via Chemical Vapor Deposition (CVD). GO was obtained via Hummer’s method and reduced to obtain a highly conductive GR nanosheet by according to the literature (105). The main process to obtain CNT papers was vacuum filtration. Catalyst particles was removed by carefully shearing MWCNTs to pieces (CNT cotton) with high speed shearing followed by immersing it into 5 mol/L of aqueous hydrochloric acid solution for 48 hours. The cleaned MWCNTs were sheared the same way again and dispersed into deionized water this time via ultrasonic treatment with Tween-80 to act as the dispersant. Then, a cellulose filter membrane with pore diameter of 0.45 microm was used to filter the uniformly dispersed CNT solution by vacuum filtration. The resulting sample was cleaned with deionized water several times to ensure the removal of remnant dispersants. The sample was placed and dissolved in acetone to obtain a freestanding CNT paper. The size of the CNT paper is controlled by the size of the filter, which ranges from 40 mm in diameter to A4 size. Electrochemical galvanotactic deposition with a two-electrode model was used to coat CNT papers with MnO2 nanoparticles on the surface. For larger sized CNT papers, the papers were bent into a ring shape in a 500 ml graduated cylinder to act as the working electrode along with a stainless steel rod as the auxiliary electrode. The electrolyte used in this process were 0.6 M MnSO4 and 0.8 M H2SO4. The temperature of this whole process was controlled by a water bath. The composite paper was cleaned with deionized water to remove remnant electrolytes, then immersed it in a 0.05 mg/ml GR solution for a few hours to infuse the GR into the composite paper. The sample was then dried in a vacuum at 60 °C for 4 hours and collected to use. The counterpart of
the asymmetric supercapacitor, flexible CNT/PANI composite paper was prepared with CV electrochemical polymerization in an electrolyte mixture that consist of 1 mol/L H2SO4 and 0.5 mol/L aniline for 300 cycles. A saturated calomel (SCE) was used as the reference electrode and a graphite sheet was used as the auxiliary electrode. The voltage used in this process ranged from -0.2 V to 0.8 V with a scan rate of 100 mV/s.

rGO/CNT is grown on Carbon Fiber (CF) with the combination processes electrophoretic reposition (EPD) of GO onto surface of CF to follow up with floating catalyst chemical vapor deposition (FCCVD) (106, 107). CNT is a great candidate to be utilized as electrode material for supercapacitor fabrications (108-111). To make the aforementioned samples, CF-GO hybrid must be fabricated first. A plain-weave CF sheets (10 x 10 mm²) was acting as the substrate. 200 mg of GO was obtained from modified Hummer’s method with natural graphite powder (112). The GO was dispersed into 100 mL of deionized water and sonicated for 1 hour. Then, CF sheets were connected to the positive and negative terminal of a constant voltage source. The CF sheets were soaked in GO solution with the voltage range of 2.0 to 5.0 V, with a deposition time of 1 minute. Next, CF-rGO-CNT hybrid sample was fabricated. From the author’s previous works (113), CNT grafted CF-rGO was discussed. Briefly, a mixture of concentrated solution containing carbon source/catalyst was prepared. 0.35 g of grounded ferrocene was dissolved into 7 mL of xylene solution and sonicated for 1 hour until the color turned to orange red. Then, CF-GO hybrid was placed into the middle of the furnace under 200 scum argon atmosphere through a tube furnace to ensure no air was left in the system. The furnace temperature was set to 750 °C at the heating rate of 15 °C min⁻¹, with 50 scum hydrogen and 350 scum argon. The final mixture was injected into the furnace with a rate of 0.2 mL min⁻¹ for 30 minutes and addition 5 minutes for reaction. The final product obtained was CF-rGO hybrid that can be directly used as a template for CNT growth. CF-rGO-CNT hybrid growth can be seen in Figure 8.
4.4 Epitaxial Growth

Epitaxial growth is a fabrication method used to produce large scale graphene films as a microfabrication process. It can be grown and can incorporate CVD as a process to deposit graphene on specific substrates. When deposition is involved the main difference between the two is that CVD uses conducting substrates while epitaxy uses an insulator substrate (114). Epitaxial growth is known to use SiC because of its flexibility and carbon content so that when the silicon is removed or melted, carbon (graphene) can be left over depending on how the experiment is carried out. Different commonly used ways of measurement for characterization include STM and LEEM. Fabricating graphene through epitaxial growth is a great option for large-scale application-fitting desires, but a uniform thickness on a large domain holds to be a difficulty to get past, along with substrate bonding affecting the electronic properties of the graphene layers produced. The use of different substrates allows for eliminating these drawbacks. For example, one source sets out use epitaxial growth on ruthenium(0001) to produce arrays of graphene in a layer-by-layer fashion to provide a path towards synthesis for applications in electronics (115).
Because of graphene’s properties, creating N-doped graphene for electrode material in recent studies show that N incorporation in graphene can potentially enhance the electron transfer efficiency for use in electrochemical capacitors. In this experiment, graphene nanowalls (GNWs) and N-doped graphene nanowalls were synthesized on a carbon cloth (CC). Before the synthesis, the CC was brought into a MPECVD chamber. H2 plasma was ignited by supplying 99.999% H2 into the chamber to rid of impurities from the CC at 900 degrees Celsius, 40 Torr, and 1500W for a few minutes. Once H2, CH4, and SiH4 (200:5:1) were added in, plasma was maintained under 1500-2000W at 1300 degrees Celsius for a few hours to allow SiC nanowalls to grow. The supply of CH4 and SiH4 were stopped while the substrate temperature was kept at the same temperature. The layers of graphene were grown covering the SiC nanowalls from surface graphitization under H2 plasma ambient within seconds. The GNWs were functionalized within NH3 by prolonging the process soon after the end of CH4 and SiH4, which is the N-doping. The process describes successful large-scale direct-growth for GNWs on a flexible substrate. The result NGNW showed maximum specific capacitance of 991.6 F/g, an energy density of 275.4 Wh/kg, and a power density of 14.8 kW/kg (116).

Another team of researchers presented a low-cost and simple epitaxial lift-off flexible and transparent single-crystal gold foils electrodes with silicone as the base (117). The interest to move beyond the conventional silicone-based chips to innovate newer electronics such as flexible transparent displays, wearable solar cells, and sensors (118-123). Although numerous breakthrough with ultrathin Si foils was found to support flexible devices and architectures, the typical fabrication process for ultrathin Si foils, vacuum evaporation or sputtering, are known to suffer from polycrystallinity or textured deposit that leads to electron-hole recombination at grain boundaries (124). This research presented a wafer-sized flexible and transparent gold foils. With the help of the epitaxial lift-off method on single-crystal substrates, a free-standing single-crystal foils can be fabricated by dissolving an adhesion layer as the sacrifice (125). Epitaxial lift-off procedure for ultrathin electrodeposited Au single crystal foils
on Si(111) substrate was shown in Figure 9. A method developed by Allongue and co-workers were used to electrodeposit epitaxial Au on Si(111) substrate (126, 127). A single-crystal Si(111) with properties such as 1.15 ohm.cm of resistivity and 0.2° miscut towards (112) was used as the substrate for Au foil growth. 0.1 mM HAuCl4, 1 mM KCl, 1 mM H2SO4, and 0.1 M K2SO4 with a pre-polarized Si electrode at -1.9 potential voltage vs. Ag/AgCl was inserted into the solution to perform the deposition process at room temperature. The pre-polarization of the electrode activated the epitaxial growth of Au and helped with the formation of an amorphous native oxide layer on the substrate. A sacrificial SiOx layer was grown in between Au and Si deposits via photoelectrochemically oxidizing Si under 0.75 potential voltage of light irradiation vs Ag/AgCl in 0.5 M H2SO4 solution. Then, polymer adhesive as the additional support and facilitator for the foil separation process was added onto the surface of Au. The etching process was carried out by using 5% dilute hydrofluoric acid to detach Au foil from the Si substrate. The foil was easily separated from the etched SiOx layer. The remaining Si substrates was etched with 0.6 M of KI and 0.1 M of I2 solution to dissolve any Au particles to ensure a clean Si substrate ready to be reused. The Si substrate can be reused numerous times because of each removal process of the 2 – 3 nm thick of SiOx during fabrication was without extensive roughening.

Figure 9: Depiction of the epitaxial lift-off method on a single-crystal Au foil. a) A pure n-type Si(111) non-oxidized wafer. b) Epitaxial deposition of Au on the Si(111) wafer. c) An insert of SiOx layer in between the Au and Si layer via photoelectrochemical oxidation of Si under irradiation of light. d) A form of adhesive (tape/hot glue) was placed on the Au layer to assist in
the lift-off process. e) Sacrificial SiOx layer was etched to provide easy separation of the Au foil from the Si substrate. f) A successful separation of the Au foil from the Si substrate (117).

Table 2: Pros and Cons of discussed methods.

<table>
<thead>
<tr>
<th>Method</th>
<th>Pros</th>
<th>Cons</th>
<th>Citations</th>
</tr>
</thead>
<tbody>
<tr>
<td>General Hummers</td>
<td>Quick oxidation method, most effective</td>
<td>High consumption of the oxidants and intercalating agents. High cost and poor scalability in real-world application</td>
<td>(17, 128)</td>
</tr>
<tr>
<td>Exfoliation of Graphite Oxide</td>
<td>Simplicity, low cost</td>
<td>Hard to fabricate in a large scale</td>
<td>(84, 85)</td>
</tr>
<tr>
<td>Chemical Vapor Deposition</td>
<td>Large quantity, high quality, Single Layer Graphene (SLG)</td>
<td>Expensive Equipment, Time consuming</td>
<td>(18, 38, 129-132)</td>
</tr>
<tr>
<td>Epitaxial Growth</td>
<td>Large quantity, van der Waals heterostructure,</td>
<td>High Temperature, Inconsistent Uniformity, Effects of substrate bonding</td>
<td>(115, 133)</td>
</tr>
</tbody>
</table>

4.5 Quality Enhancement Techniques

Complementary techniques are done within an experiment to assist the main methods that were earlier described. One or all methods can be involved within an experiment and each offers result that make fabrication more successful or simpler. A compiled set of advantages and disadvantages to using each complementary factor can be found in Tables 2 and 3. Other techniques can be implemented and created in fabrication, like using a Fe-catalyzed glucose-blowing approach with NH4Cl as a clowing agent and Fe as a graphitization catalyst to fabricate graphene-like carbon nanosheets (134). Even by doping graphene with dual atom (N, P) will significantly boost the oxygen reduction reaction (ORR) and oxygen evolution reactions (OER) performance (135).

4.5.1 Vacuum Filtration

Suction (vacuum) filtration is a technique carried out in chemistry recrystallization experiments that gives a greater rate of filtration. Compared to normal filtration, which uses gravity as the force to push the liquid through the filter, vacuum filtration uses a pressure gradient. This allows for a variable rate depending on the pump strength. Vacuum filtration can be applied to any method, since its effects can benefit GO substitutes or solutions in different ways. It can also be implemented in ways to insert nanostructures within graphene sheets, like the example of a filtration method via self-assembly of water dispersible graphene and mesoporous carbon nanosphere.
(MCS) using the MCS to be inserted between graphene sheets by vacuum filtration (136). Overall, it is done to filter out liquid in a solution so that all that is left is graphene.

To reproducibly obtain uniform thin films with enough GO layers over large surface areas, the vacuum filtration method is a good approach. The vacuum filtration method can be used widely to deposit highly uniform single-walled carbon nanotube thin films. As described in the following experiment, as well as in a general sense, vacuum filtration is the filtration of a GO suspension through a commercial mixed cellulose ester membrane, which has an average pore size of 25 nm (137). The membrane pore sizes can vary for each experiment depending on the desired filtration. The liquid passes through the pores, but the GO sheets become lodged.

Another report starts out with GO from applying the modified Hummers method (17). In preparing the rGO films, vacuum filtration was applied to aqueous solutions of the created GO using mixed cellulose ester membranes as the filter (filtration volume can fluctuate depending on the desired film thickness and transparency. This also applies to variance in GO solution concentration). Here it can be seen that transparency and film thickness are key elements in vacuum filtration, along with the pump strength to determine the variable rate of flow.

In an example of a hybrid fabrication method, one team compared the use of carbon black (CB) as a conductive additive for both graphene and reduced graphene oxide (rGO) (138). The graphene/CB mixture acts as a good supercapacitor electrode. The reduced graphene oxide/CB mixture was subjected to vacuum filtration methods to obtain a free-standing, flexible nanocomposite film. Vacuum filtration is essential in obtaining the film, regardless if it is a hybrid material or not. The benefit of using CB is that it not only acts as a conductive additive but it also acts as a spacer which prevents restacking of rGO layers. Preventing restacking is desirable because it means that the layer(s) will reach a larger surface area and higher capacitance. CB amounts must be balanced/regulated though, because too much can cause agglomeration.

### 4.5.2 Roll-to-Roll Production
Roll-to-roll (R2R) production is a unique complementary method and transfer process that falls under CVD. It involves a multistep process with a long and thin copper film for the graphene to be applied to. Roll-to-roll production is usually accompanied by wet-chemical doping of (mainly) monolayer graphene films grown by chemical vapor deposition onto flexible copper films (139). Due to lack of efficient methods for synthesis, transfer, and doping of graphene at the nanoscale, the production of transparent conducting films from graphene are difficult. In the roll-to-roll wet chemical doping process, there are essential steps to ensure success. The films used have sheet low resistances (~125 Ohms) and high optical transmittance (~97.4%). The graphene film was grown on copper foil and attached to a thin adhesive polymer film. Copper was removed by electrochemical reaction with a solution, in the following case it was ammonium sulfate. Finally, the films were separated so that the graphene film can be thermally treated. It is assumed that the graphene was grown on copper foil because of its relatively cheap price point and electrical conductivity. Copper foils with larger grain size yielded higher-quality graphene films. The copper foils were heat-treated to increase the grain size.

An in-depth experimental detail showed a similar approach and explained their four key steps for the R2R encapsulation (140). The first step was R2R growth of a long monolayer graphene film on Cu foil from the CVD method. Second was either coating or transfer of metal nanowires onto a EVA/PET (ethylene vinyl acetate/polyethylene terephthalate) plastic film. The usage of the exact plastic as the film in other experiments was optional. Third, hot lamination of graphene/copper foil onto metal nanowires precoated with EVA/PET plastic was carried out. Lastly, electrochemical bubbling delamination of graphene film from copper foil was done. The process started out with a roll of continuous monolayer graphene film grown on 5 cm x 5 m copper foil by the CVD method. After electrochemical transfer, Cu foil could be preserved and reused for graphene growth. The following figure illustrates the process that is being described.
Three cycles were conducted and after Raman spectra analysis, the graphene R2R grown on the reused copper foil was improved. This was from the surface flattening and grain size enlargement of the foil during high temperature annealing. Next, the metal nanowires were coated on the EVA/PET sheet by using a Mayer rod. The EVA surface was pretreated by air plasma so that its hydrophilicity is improved. Once the CuNWs (copper nanowires) were coated, Cu/graphene film was hot-laminated onto the CuNWs which creates a laminated structure at 100 degrees C. This was done by two rollers giving off the specified heat and mechanical pressure during the lamination. This process reduced air traps, cracks, or any blemishes due to the simultaneous heat and pressure. What was left was a cathode film of Cu/graphene/NWs/ EVA/PET. Lastly, it was stripped away of the lamination by,
what the source calls it, the electrochemical bubbling delamination transfer method. This process involved the laminated film to be partially immersed in water. While immersed, hydrogen bubbles were generated through water electrolysis, which delaminated the graphene film at very low electrolysis voltage. The speed at which the film immersed and emerged can reach up to 2 cm/s. Since the cathode has a negative charge during the delamination process, the films can avoid oxidation and the Cu foil can be reused. The resulting graphene/NW/plastic films proved to have high optoelectronic performance, great corrosion resistance, flexible, and adhesiveness. This film proved to be successful for flexible, transparent electrodes. It offered mass production and low cost, which helped to pave a pathway to future applications. Other R2R experiments may include alterations like expanding the length of the copper foil to 100 meters long in the process (130). Another group fabricates films at 294 mm thin under low temperatures (131). This shows that the R2R complementary factor does not have to be done in critically specific ways. However, the experiment provided by Deng and Hsu does provide an appropriate film length and appropriate steps to fabricate a stable, successful electrode.

4.5.3 Ultrasonication

Ultrasonication is a process that applies sound energy to agitate particles in a sample with frequencies greater than 20 kHz. It reduces small particles in a liquid to enhance stability and uniformity. Two common ways to apply ultrasonication to a solution are by bath and probe units. Even though bath units are more commonly used, they have two drawbacks that decrease experimental repeatability and reproducibility (141). These two drawbacks are a lack of uniformity and a decline of power over time. A lack of uniformity is resulted from ultrasonic baths because only a portion of the volume of the ultrasound source experiences cavitation. Luque-Garcia and De Castro suggested that ultrasonic probes have the upper hand over ultrasonic baths since they focus their energy on a localized sample zone, which allows for more efficient cavitation.
This experiment started out by using Hummers method to oxidize and form the graphite oxide (132). It was then added to a concentrated solution of H2SO4, K2S2O8, and P2O5 with an atmospheric temperature of 80 degrees C. Further experimental processes similar to the Hummers method examples were carried out until the point that the obtained graphite oxide powder was dried and dialyzed in 0.5% graphite oxide dispersion. At this point, a new approach was taken instead of the common next step of exfoliation of graphite oxide. SDBS (sodium dodecylbenzene sulfonate) was added as a surfactant, which facilitated the exfoliation, lowered the surface tension, and allowed for a larger sheet of graphene oxide from ultrasonication. Using Ultrasonication as the exfoliation method allowed for particle agitation to properly exfoliate the solution. It is typical to sonicate 0.1 mg/mL graphite. In a review, sonication was carried out with lithium salt intercalated graphite mixed with water (142). It recorded more than 80% few-layer graphene, where lithium and water react to form hydrogen gas which can further aid exfoliation.

One group utilized fibers as the base for a superconductor material that was capable to show great performance in conductivity and capacitance (143). A flexible fiber-shaped supercapacitor (FSSCs) was recently picked up by researchers due to its potential to be assembled into other structural systems in flexible electronic applications for its tiny volumes, structural flexibility, and high capacitance density (144-148). The ED (electronic deposition) method was previously reported to prepare the nickle-coated CF (carbon fiber) successfully (149). The fabrication of nickle-coated carbon fiber started off treating CF threads with acetone, alcohol and distilled water respectively under ultrasonication for 15 minutes. After that, the CF thread was immersed in 10 g . L-1 of SnCl2 for 20 minutes, followed by cleaning it with distilled water several times and then drying it at 70 C. Subsequently, the CF thread was immersed into 0.25 g . L-1 of activated PdCl1 and 10 ml . L-1 of hydrochloric acid solution for 20 minutes, then repeated the same cleaning and drying process as before. The resulting treated CF thread was then immersed into a homemade plating bath (40 g . L-1 of NiSO4 . 5H2O, 20 g . L-1 of sodium citrate, 10 g . L-1 of lactic acid and 1 g . L-1 of dimethylamine borane in water) at 90 °C for 15 minutes to
undergo electronic deposition method to obtain the metallic structure in the water bath. The next step is to fabricate the pen ink/nickel/CF via facile dip-coating method (150). The procedure required the previously prepared nickel-coated CF to be immersed into pen ink solution (average thickness of in film = 600 nm) for 5 minutes. After that, the nickel/CF was transferred into a hot stage at 60 °C for 2 hours. The desired thickness of pen ink to be coated on Nickel/CF could be controlled via repeating the aforementioned process until desired thickness. Then, the fabricated Pen Ink/Nickel/CF sheet was coated with Ni-Co DHs via ED method in a standard three-electrode system at 25 C. The pen ink/nickel/CF was treated as the working electrode, where else a Pt wire was the counter electrode and the saturated calomel electrode (SCE) as the reference electrode. 100 ml of 0.1 M metal ion solution with Ni^2+/Co^2+ with a 1:2 ratio was the acting electrolyte for the process, with the voltage range from -1.0 V for 10 minutes. The resulting Ni-Co DHs/pen ink/nickel/CF sheet was cleaned with distilled water followed by drying at 60 °C for 30 minutes. The final process involved the assembly of solid fiber supercapacitor. An electrolyte consisting of PVA/KOH gel was prepared with 3 g of dissolved PVA in 20 mL of distilled water at 100 °C under vigorous stirring for 1 hour. Subsequently, 10 ml of 0.3 g . mL-1 KOH solution was added into the previous mixture via the drop method gradually, followed by a continuous stirring at 80 °C for 1 hour. Then, Ni-Co DHs/pen ink/nickel/CF was used as the positive electrode while pen ink/nickel/CF was used as the negative electrode for the ED process consisting of the prepared gel electrolyte for 5 minutes. After the ED process, the resulting samples were taken out to be dried at room temperature. Finally, PVA/KOH gel electrolyte was used to assemble the previously said electrode together, placed in a silicone tube to be protected and heated at 60 °C for 2 hours to evaporate excess water containing in the electrolyte.
4.5.4 Hydrothermal

The hydrothermal method is a method done to substances that are intended to be mixed with graphene to create a hybrid. This method involves using an autoclave and temperature control to crystallize or recrystallize a substance. It can also apply to any main method of fabrication, since the hydrothermal method allows for crystallization and each method can operate with a solid hybrid material. In the following example, fabrication of a graphene/vanadium hybrid supercapacitor electrode took place. The mixtures were designated as VURGO and VSRGO (Vanadium Ultralarge/Small Reduced Graphene Oxide) and were produced through the following process (151). The preparation of UGO started with thermally expanding graphite flakes and synthesizing GO using the Hummers method. The GO was diluted with deionized water and centrifuged at 8000rpm for 40 min, which was when the SGO was obtained. The sediment was spread in deionized water and centrifuged another time, except at 4000rpm. A precipitate was produced, which contained the UGO, and was added to DI water. Usually, ultrasonication is a step that follows this, but it was not
done here specifically because it reduces the lateral size of the GO sheets. The highest yield for UGO was at 8.3 wt%. Modifications should be made to see a higher yield for UGO.

In continuing to fabricate VUGO and VSGO, the VO2 was synthesized by the hydrothermal method, which crystallized the substance for high-quality crystal growth and composition control. Using a VO2:UGO weight ratio of 7.3 or using an annealing temperature of 250 °C for 2 hours are two ways to optimize the electrodes in forming a free-standing film. The VUGO mixture was stirred at 70 °C for an hour and then filtered through a cellulose acetate membrane by method of vacuum filtration. Make it known that the transparency and film thickness will depend on the characterizations of the membrane used. The film must be washed with DI and dried several times until it can be removed from the filter paper. The film was annealed at 250 °C for 2 hours in argon gas so that the GO could change to reduced graphene oxide (rGO). VURGO proved a better resultant than VSRGO, because of its porosity and interconnected structure. This example showed that the hydrothermal method used to fabricate a graphene mixture can simply apply to just the hybrid material being added (in this case the VO2).

One group discussed about a freestanding reduced graphene electrode coated with carbon Li4Ti5O12 nanosheets (LTO-C/rGO) to be used in sodium ion batteries (152). Electrochemical performance of nano/microstructured LTO composites and carbon-based materials such as CNT and graphene was focused to be improved. The improvements for the former are the transportation lengths of Na ion and shortens the electron (153-155) where else the latter provides more efficient electron transportation (156-161). Firstly, modified hydrothermal method with annealing was used to fabricate LTO nanosheets (162). To process involved 1.7 ml (5mM) of tetrabutyl titanate, 0.189 g of LiOH. H2O and an estimated amount of GdCl3. 6H2O to be mixed into 20 ml of ethanol under magnetic stirring for 24 hours at room temperature. After that, 25 ml of deionized water was added into the mixture and stir it for 0.5 hours. The resulting solution was then transferred into a 50 ml Teflon-lined stainless autoclave and placed into an oven. The solution was heated to 180 °C for 36 hours to achieve Li1.81H0.19Ti2O5 xH2O (H-LTO). After the hydrothermal process,
white H-LTO precursor was collected at the bottom of the reactor. The aforementioned sample was then washed with ethanol 3 times, followed by heating at 700 °C for 6 hours in a horizontal furnace tube with air to get LTO nanosheets. Next, LTO nanosheets needed to be coated with carbon. H-LTO precursor was then prepared and allocated 200 mg to be dispersed in 10 ml of ethanol under magnetic stirring. Nominal 5 wt% of H-LTO precursor and D(t)-glucose ethanol was dissolved in 5 ml of ethanol and later to be drop-wise added into the previous solution and stirred, followed by slow evaporation at 80 C. The resultant slurry was heated to 700 °C under stirring with Ar/H2 (5%) atmosphere for 6 hours to obtain carbon-coated LTO (LTO-C) nanosheets. The final step was to obtain LTO-C/rGO film by getting GO via oxidized graphene using modified Hummer’s method. LTO-C nanosheets were dispersed into 100 ml of 1 M NH4HCO3, later to be added drop-wise into a solution with dispersed GO in 100 ml of deionized water with magnetic stirring. Resulting black composite that sank to the bottom within 10 minutes was collected with vacuum filtration through a microporous filter (0.45 microm). The black composite was dried at 60 °C for 6 hours, then peeled off and annealed at 700 °C under Ar/H2 (5%) atmosphere for 2 hours. Then, a free-standing LTO-C/rGO film was fabricated.

4.5.5 Substrates

There are two main types of bendable electrodes for LIBs/SCs. One uses nonconductive bendable substrates, such as paper, polymers, and textiles to support the active materials (163-166). The selected materials must meet certain requirements including ability to deform under stress/strain. The other uses conductive supports, which involve CNTs or graphene films with high conductivity. On top of having an ideal elastic modulus and having ionic conductivity, whatever material used for flexible electrode (in this case graphene) should have substrates that do not produce any contaminants during electrochemical reactions and should be totally inactive against any chemicals in the system. High resistances would be a drawback too. Polymer substrates have high resistances, which acts against what is trying to be achieved, so the use of substrates with lower resistances like Cu or Al foil current
collectors is more desirable. The figure below shows the difference between using polymer substrates (purple) and using cellulose paper/cloth substrates (green).

Figure 13: Comparison of cellulose paper (green) and polymer non-conductive substrates (purple) for flexible electrodes (conductive meaning graphene) (163).

### 4.5.6 Comparison Table for Quality Enhancement Techniques

Table 3: A comparison table for quality enhancement techniques.

<table>
<thead>
<tr>
<th>Methods</th>
<th>Pros</th>
<th>Cons</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vacuum Filtration</td>
<td>Large Surface Area (Eda)</td>
<td>Cost of equipment, Time</td>
<td>(17)</td>
</tr>
<tr>
<td>Ultrasonication</td>
<td>Stability and Uniformity</td>
<td>Reduce lateral size of graphene</td>
<td>(151)</td>
</tr>
<tr>
<td>Hydrothermal</td>
<td>Nanometer sized graphene sheets, For hybrid materials</td>
<td>Long hours required</td>
<td>(151, 167)</td>
</tr>
<tr>
<td>Substrates</td>
<td>Can optimize properties</td>
<td>Poor selection can result in lack of properties</td>
<td>(163, 167)</td>
</tr>
<tr>
<td>General Environmental Methods</td>
<td>Utilizing recyclable materials</td>
<td>Time, Money, Does not yield the best characteristics</td>
<td>(4, 168, 169)</td>
</tr>
</tbody>
</table>

### 5 Environmental Impact and Toxicity

While each fabrication method conveys desirable results, some methods do have environmental drawbacks. For example, in Hummers Method, due to the NO₂ and N₂O₄, toxic gas can be generated. Other drawbacks include residual nitrate and low yield. Many attempts have been made to modify this method like replacing the NaNO₃ or adding a
peroxidation step before the KMnO4 oxidation (14). Overall, there is a high demand for an economic, cost-effective, environmentally friendly process to synthesize GO. The researchers tackle these problems to make a modified Hummers method more desirable for practical application. When increasing the amount of KMnO4 and concentrated H2SO4 (with 1/9 H3PO4) in place of NaNO3, the modified Hummers method leads to higher yield, easier temperature control, and less toxicity.

On the renewable/recyclable side, a proposal came up to recycle industrial waste and utilize it as graphene electrodes using coal tar pitch (CTP) as carbon feedstock (170). Dilute coal tar pitch with 8 wt% of quinoline solvent was used. Then the mixture was spin coated at 6000 rpm for 1 minute on a SiO2 (500nm)/Si substrates and baked at 240 °C for 30 minutes. The next step was to deposit the Ni layer (200 nm) on a coal tar pitch film at room temperature. (Magnetron Sputtering System, SNTEK; working pressure 7mTorr; power 50W; Ar flow rate 50 sccm). The resultant film was then annealed using thermal CVD method at 1100 °C for 4 minutes with flowing 50 sccm of AR and 10 sccm of H2, with a combined pressure at 0.3 Torr. Then, the tube was removed from the furnace but maintained the same amount of Ar/H2 flow to cool the film to room temperature. Once cooled, the Ni layer could be etched away and cleaned with deionized water after dipping the sample film into FeCl3 solution for 1 minute (Iron(III) chloride solution 45°Be’).

The process to get a flexible electrode is time consuming and tedious, usually involving high-temperature treatment or the use of toxic chemical treatment (171). Hence, the researchers showed a method that can be fast, scalable and environmental-friendly to fabricate high-performance rGO/cellulose paper supercapacitor electrode. An aqueous suspension of cellulose pulp fibers that consist of virgin pulps and recycled waste pulps from newspapers (0.15 wt%, 200 mL) was mixed with aqueous dispersion of single layer GO sheets that was purchased (GO0TQ2, 1 wt%, width: 10-30 μm) (0.032 wt%, 15.5 mL, GO content: 5 mg) and an aqueous solution of polyethylenimine (PEI) (Average molecular weight: 1800) (1.0 wt%, 0.4 mL) with intervals of 10 minutes in between adding each solution. The resultant solution was then filtered by suction filtration through
a #300 wire mesh for 4 s. After that, the wet paper sample was sandwiched between a hydro-phobic glass and paper towel followed by hot pressing it at 110 °C for 10 minutes to dry the sample (1.1 MPa). The resultant sample was then peeled off from the wire mesh to make GO/cellulose paper composite with diameter of 75 mm and thickness of ca. 100 μm. The GO/cellulose paper then underwent flash reduction at room temperature in air using a Pulse-Force 3300 equipped with xenon flash lamps. The process was carried out 10 times at a frequency of 2 Hz for each surface of the GO/cellulose paper composite.

Figure 14: Depiction of the preparation procedure for the rGO/cellulose paper composite via papermaking and successive flash-reduction processes with paper composite thickness of 100 μm and diameter of 75 mm (171).

To obtain reduced graphene oxide, a general method is to use a reducing agent to chemically reduce GO to rGO. Reducing agents usually possess harmful characteristics such as toxicity, flammable and reactivity. A way to harness aqueous extract of *Hibiscus sabdariffa* L was figured out (172) to be used as a reducing agent, which is in contrast,
1.0 g of natural graphite sheets was gathered and added into 100 mL of concentrated sulfuric acid (H2SO4) while stirring it for 1 hour in an ice bath. After that, 4.0 g of potassium permanganate (KMnO4) was slowly added into the solution in a water bath at room temperature for 6 hours. 200 mL of deionized water was then added to dilute the solution followed by adding 100 mL of deionized water and 20 mL of hydrogen peroxide (H2O2) into the solution gradually while stirring. The solution eventually changed color from dark purple to a bright yellow brown solution. The stirring solution was allowed to rest at room temperature for 24 hours. After 24 hours, the solution was then centrifuged and cleaned with deionized water until the pH level is neutral, followed by drying the sample in a vacuum pumping system at room temperature. Next, preparation of the aqueous extract of *Hibiscus sabdariffa* L was executed by purchasing 10.0 g of *Hibiscus sabdariffa* L and adding it into 1.0 L of deionized water. The mixture was then heated to 60 °C for 1 hour. The resulting aqueous extract then could be obtained by filtering the solution using 0.45 μm pore size polypropylene filter papers. To reduce GO, 80 mg of GO suspension was put into 200 mL of *Hibiscus sabdariffa* L aqueous extract. The mixture was then to underwent ultrasonication for 1 hour and followed by subsequently stirring and heating for another hour. After that, the sample was cleaned with deionized water and vacuum dried at room temperature. To get a flexible graphene film electrode, HRGO-R in 0.4 mg mL-1 of deionized water had to be dispersed under ultrasonication followed by filtration and drying at 70 C. Graphene films could be obtained by peeling off 1 μm of HRGO-R. Supercapacitors could also be created by placing two graphene films separated by a filter paper soaked with 1 M H2SO4 that acted as the electrolyte. Then placing two platinum foils acting as a current collected in a sandwich-like structure made the supercapacitor cell.

This source showcases eco-friendly fabrication method of 3D porous structured aerogel that encapsulates characteristics like large specific area, efficient ion-diffusion, superior mechanical integrity and multidimensional continuous pathways for electron transport for high performance energy storage and conversion applications, catalysis and adsorbents for environmental remediation (173-175). The most common method to fabricate
graphene hydrogels in the research field seems to be via hydrothermal method (176-178). The simple fabrication process allows significant savings in industrial terms to produce high-performance electrodes for lithium-ion batteries. The first step was to prepare GO dispersion via modified Hummer’s Method. 3 g of graphite powder, 1.5 g of NaNO3 and 69 ml of H2SO4 were mixed together in a beaker, followed by adding 9 g of KMnO4 and stirred in an ice bath to keep the mixture at 0 °C. The temperature was increased to 35 °C after 30 minutes of stirring. Stirring for an additional 30 minutes, then gradually adding 138 ml of cold deionized water (15 °C) was completed. Then, 213 ml of warm water (80 °C) and 15 ml of H2O2 respectively were added. The resulting mixture was placed in a centrifuge to be separated and washed with HCl-H2O solution (HCl/H2O = 1/10) three times. Finally, the solution was dissolved in deionized water and dispersed homogenously by ultrasonic agitation for several hours. To synthesize SnO2/Graphene aerogels, SnO2 nanocrystal was prepared by dissolving 2.325 g of SnCl4·5H2O in 200 ml of deionized water followed by hydrothermally treated for 160 °C for 16 hours. The resulting precipitate was washed by placing it in a centrifuge three times and re-dispersed in 200 ml of ethanol to produce a suspension. 2.4 ml of SnO2 dispersion was then added drop-wise into 6 ml of GO (2mg ml⁻¹) (mass ratio 1:1). The dispersion was placed in an ultrasonic agitator and stirred for 24 hours to ensure a homogenous dispersion. Subsequently, 120 mg of ascorbic acid powder was added into the mixture and stirred for 30 minutes. The aforementioned mixture was placed into a mold and into a draught drying cabinet at 75 °C for 4 hours to obtain 3D graphene hydrogels. Then, the monolith was taken out of the mold and immersed in deionized water for 24 hours and freeze-dried into graphene aerogel. Finally, the aerogels were heated to 550 °C for 30 minutes to increase the degree of reduction. The preparation of graphene aerogel as electrodes was the final process. The graphene aerogels were heated at 650 °C for 30 minutes with N2 atmosphere. After the heating process, the 3D graphene sheets were used as electrodes without any post-processing.

Another source utilized eco-friendly methods to produce a flexible supercapacitor from wood transverse section slice (WTSS) and reduced graphene oxide (179). The thickness of a native WTSS was cut from Chinese fir to about 180 µm with rotary microtomy,
followed by vacuum drying at 50 °C for 2 hours. Without further purification, graphite power and 85% aqueous solution of hydrazine hydrate was used to create an aqueous dispersion of graphene oxide (GO) nanosheets with Hummer’s method. WTSSs of different areal densities (0.36 mg cm⁻², 0.49 mg cm⁻², and 0.68 mg cm⁻²) was immersed in aqueous GO dispersions of 1 mg mL⁻¹ for 5 min and allowed to dry naturally in room temperature to obtain different GO loadings. The WTSS-GO sheets was then reduced with aqueous solution of hydrazine hydrate at 90 °C for 3 hours. The finalized results were then be rinsed with deionized water and dried in a vacuum at 50°C for 2 hours. The formation of the supercapacitor was created as seen in Figure 15. The assembly was seen to utilize the traditional laminated two-electrode configuration with PVA/H₃PO₄ gel (blue) used as electrolyte and a separator.

This example describes how researchers devised a way to synthesize transparent graphene oxide flexible electrode by mixing GO with the conductive polymer in an aqueous solution (180). First, GO:H2O dispersion (15 μg L⁻¹) was obtained using the chemical route and the conductive polymer from Sigma-Aldrich [3,4-ethylenedioxythiophene]:poly[styrenesulfonate] (PEDOT:PSS) was purchased. Then 3 ml of GO dispersion was mixed with the desired amount of PEDOT:PSS solution volumes via magnetic stirring for 24 hours. The GO dispersion to GO:PEDOT ratio were known to be GO:PEDOT (0.1%), (0.5%), (1.0%), (5.0%), (10.0%) and (20%). Next, Polyethylene terephthalate (PET) was used as substrates. GO:PEDOT films were obtained via the drop casting method over PET substrates and left it to dry at room temperature for 4 hours. After that, it was placed on a hot plate to thermally anneal it at 80 °C at standard room pressure and temperature. Not only are there sustainable ways to
fabricate graphene, but there are environmentally friendly ways to utilize graphene. Like previously mentioned in Hummers’ method and exfoliation experiments, along with epitaxial growth and the ability to fabricate it via CVD using ethanol as a carbon source on Ni foams, capacitive deionization (CDI) is an effective way to treat wastewater and benefit the environment (181, 182). Heavy metal wastewater pollution has become a problem affecting our environment and water purity as metallurgical and other manufacturing plants rise. Utilizing graphene as an electrode with capacitive deionization to separate and recover heavy metal ions and salt ions in wastewater allows for greater treatment and higher electromagnetic interference shielding effectiveness (183, 184). Current strategies in desalination of saline water include ultrafiltration, reverse osmosis, and distillation processes, but with graphene electrode CDI, it would become the most optimum option (181, 185).

### 6 Summary Table

Table 4: A summarized synthesis experimental methods.

<table>
<thead>
<tr>
<th>General Synthesis Method</th>
<th>Experiment</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Modified Hummer’s Method</td>
<td>A flexible polyaniline/graphene/bacterial cellulose supercapacitor electrode</td>
<td>(186)</td>
</tr>
<tr>
<td></td>
<td>Producing large-area, foldable graphene paper from graphite oxide suspensions by in-situ chemical reduction process</td>
<td>(187)</td>
</tr>
<tr>
<td></td>
<td>Silver Fiber Fabric as the Current Collector for Preparation of Graphene-Based Supercapacitors</td>
<td>(188)</td>
</tr>
<tr>
<td></td>
<td>Cu particles induced distinct enhancements for reduced graphene oxide-based flexible supercapacitors</td>
<td>(189)</td>
</tr>
<tr>
<td></td>
<td>Controllable morphology of polypyrrole wrapped graphene hydrogel framework composites via cyclic voltammetry with aiding of poly (sodium 4-styrene sulfonate) for the flexible supercapacitor electrode</td>
<td>(190)</td>
</tr>
<tr>
<td></td>
<td>Assembly of graphene aerogels into the 3D biomass-derived carbon frameworks on conductive substrates for flexible supercapacitors</td>
<td>(23)</td>
</tr>
<tr>
<td>Chemical Vapor Deposition</td>
<td>Superflexible, high-efficiency perovskite solar cells utilizing graphene electrodes: towards future foldable power sources</td>
<td>(4)</td>
</tr>
<tr>
<td></td>
<td>Graphene-Based Flexible and Stretchable Electronics</td>
<td>(89)</td>
</tr>
<tr>
<td></td>
<td>A graphene foam electrode with sulfur loading for flexible and high energy Li-S batteries</td>
<td>(90)</td>
</tr>
<tr>
<td></td>
<td>Flexible, sandwich-like CNTs/NiCo2O4 hybrid paper electrodes for all-solid state supercapacitors</td>
<td>(95)</td>
</tr>
</tbody>
</table>
In conclusion, it is evident that the need for furthering graphene’s capabilities in an applicational sense is strong and that there are many approaches one can take to fabricate graphene as a flexible electrode. The incredible interest in graphene can be linked to its ballistic transport at room temperature combined with chemical and mechanical stability (191). The presence of advancement in portable, wearable, and implantable that go with sensing devices with a flexible electrode can shape the future. Having something with biocompatibility and powerful properties at a nanoscale can surely benefit many areas like biomedical engineering (120, 192-200). With the motivation to supply compressed informational and experimental articles, many main and minor fabrication methods were compiled and gone into detail. This was done so that they could be learned from and manipulatable for researchers to fabricate ideal graphene as flexible electrodes. The modified Hummers method, chemical vapor deposition, and exfoliation of graphite oxide proved as critical methods to use to obtain the desired electrode. Along with that, the complementary factors allow for support in the fabrication process. The final portion that was addressed revolves around remaining environmentally friendly in the fabrication process. As the world takes steps forward on the path toward sustainability, using materials and resources that will not put future generations at risk should always be a priority (201, 202). For this case, different methods were addressed to give opportunity to pursue a “green” fabrication process. This review paper does not contain every possible method and factor to fabricate graphene as a flexible electrode, but solemnly addresses
common methods taken by researchers. Each method is organized and compiled into multiple tables for quick and easy understanding of advantages and disadvantages as well as fabrication potential.

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