Preparation of Graphene Oxide from Lignin by Gel Combustion Method and Its Performance as Supercapacitor

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Abstract. Carbon materials derived from biomass are drawing increasing attention due to its advantages in economical as well as ecological production. Among the biomass resources to be utilized for that purpose, lignin possesses unique characteristics: it is available in bulk quantity as a low-value by-product from the pulp and paper industries and its chemical structure based on complex aromatic networks provides excellent precursor for grahitic carbon materials such as graphene. The existing techniques to convert lignin to carbon materials require long processing time, complex steps and higher temperatures, thus increasing the production cost and hindering its commercialization. Here an alternate low-cost, one-pot synthesis method utilizing organic solvent gel combustion technique is developed to yield crystalline graphene oxide as an end product. Capacitor performance of the developed graphene oxide was investigated in terms of the parameters involved in the synthesis procedure.

1 Introduction

Demand for environmental friendliness and renewable nature on energy sources have been increasingly prominent in recent years with the heightening concerns about climate neutrality as well as depletion of natural [1-5]. Supercapacitors, a type resources of electrochemical capacitors, have attracted much attention as one of the promising energy storage system for leveling renewable power fluctuations, because of their outstanding properties such as excellent cycle performance, fast charging/discharging capability and high power density [6-9]. The performance of supercapacitors strongly depends on the electrode employed, and carbon materials including graphene, activated carbon, carbon nanotube and so on are among the candidates sought after for the purpose [10-12].

A potential renewable source for the carbon materials is lignocellulosic biomass [13]. In Malaysia, for instance, about 5.4 million hectare of the oil palm agricultural land generates about 80% of the biomass, making it one of the largest contributors of lignocellulosic biomass in the world [14]. On the other hand, crude lignin is obtained in large quantities in the pulp and paper industries, mainly as Kraft lignin and lignosulfonate. Kraft lignin is the largest raw lignin byproduct produced from the pulping process. As a low value compound, it has so far mainly been used as a fuel source in combustion processes and less than 5 % of its world supply has been used for other purposes [15,16]. Therefore, valorization of lignin into value-added chemicals or materials suitable for largescale industrial applications has been the focus of many researches [17].

Lignin is regarded as a starting core for synthesis of functional carbon materials, and numbers of functional carbon materials like carbon fibers and porous carbon with different functionalities can be synthesized from it through thermochemical methods such as pyrolysis or hydrothermal carbonization. Synthesis of amorphous (activated carbon and carbon black) and crystalline (carbon nanotubes/nanofibers and graphene) carbon materials with controllable functionality often involve tedious synthetic process. Furthermore, most of these methods require fossil fuel-based chemical precursors (e.g., CH₄, C₂H₂, and C₂H₄), metal catalysts, and/or complicated apparatus involving high processing temperatures, which are not necessarily environmentally friendly or economically sustainable [16].

Among those carbon materials, graphene has carbon atoms in honeycomb structure and shows interesting characteristics with excellent electrical and mechanical properties [18,19]. It attracts significant attention due to their potential applications in conductors, batteries, catalysts and biosensors. Common production method for graphene is by mechanical exfoliation, epitaxial growth and chemical vapor deposition [20]. The largescale production of graphene from lignin available globally, however, is bottlenecked due to the high cost of its production.

In search of a low-cost process for graphene synthesis from lignin, gel combustion method is chosen in the current study. The citric acid sol-gel combustion method is one of non-alkoxide sol-gel methods, which is efficient for the preparation of nanocrystalline inorganic powders. The attractive features of this method are its ability to synthesize materials with high purity, better homogeneity and high surface area in a single step [21]. Even though the concept of gel autocombustion is universal, the methods to be applied may not be the same for all the materials. Thus in this research, synthesis procedure utilizing gel combustion method will be developed to obtain crystalline carbon material graphene oxide as an end product from Kraft lignin, and their supercapacitance performance will be evaluated based on the synthesis parameters.

2 Experimental Procedure

Lignin purchased from Tokyo Chemical Industry was used as a starting carbon material and A.R. grade citric acid, ethylene glycol, magnesium nitrate purchased from Sigma Aldrich were used as chelating, gelation and combustion agent, respectively. Potassium hydroxide was selected as solvent to dissolve the raw material and other chemical agents.

Lignin conversion procedure for graphene adopted here is as follows.

- Kraft lignin was kept in an oven at 105 °C for 2 h to remove its moisture.
- Calculated amount of Kraft lignin and magnesium nitrate were added, followed by citric acid, into the solvent and stirred for 1 h with magnetic stirrer to prepare clear solution.
- Ethelene glycol was added into the solution at 1:1 volume ratio.
- The prepared composition was placed on a hot plate at 80 °C and kept stirring to maintain the homogeneity of the mixture.
- Depending on the volume of the initial solution, the gelation time differs.
- Continued the heating even after gelation which will removes the moisture and powder remained at the bottom.
- Once moisture was removed autoignition started and whole gel will be combusted and turn into carbon powder.
- Synthesized samples were washed with hydroxide solution to eliminate magnesium.
- Prepared samples were washed with acetone several times.
- When necessary, potassium hydroxide washing was also performed.

Prepared graphene samples were examined by Raman spectroscopy (Horiba Jobin Yvon HR800), and their capacitance by cyclic voltammetry (PARSTAT 4000, Princeton Applied Research, India.).

3 Results and Discussion

3.1. Appearance of Starting Lignin and Produced Graphene

Figures 1 and 2 show physical appearances of the starting lignin material and the synthesized graphene samples, one as synthesized and the other KOH-washed.



Fig. 1. Starting Kraft lignin.



Fig. 2. Synthesized graphene samples (left: as prepared, right: KOH-washed).

3.2. Raman Spectroscopy

Figures 3 and 4 exemplify Raman spectra of the produced graphene shown in Fig. 2 using laser excitation at 2.41 eV. Prominent features in the spectra are the so-called G band of monolayer graphene appearing at about 1590 cm⁻¹ and the 2D band at about 2700 cm⁻¹. We also see the so-called disorder-induced D band at about half of the frequency of the 2D band (around 1360 cm⁻¹) for the samples disordered or with fair amount of graphene edges.

Referring to Figs. 3 and 4, the above observation in Raman spectra of the prepared sample lies in the same peak ranges of the reported graphene Raman shift [22]. It ensures that the synthesized samples contain graphene family, evidencing that the prepared samples are fully converted into graphene material.



Fig. 3. Raman Spectrum for the as synthesized sample.



Fig. 4. Raman Spectrum for KOH washed synthesized sample.

3.3. Cyclic Voltammetry

Figures 5 and 6 show cyclic voltammograms of the two samples shown in Fig. 2 and their Raman spectra indicated in Figs. 3 and 4. On the left of each figure cyclic voltammograms of the synthesized graphene materials are shown with different sweeping rates, and on the right, capacitance measurement with different amperages. Table 1 summarizes the capacitance data for these two samples at various current.



Fig. 5. Left: Cyclic voltammogram of as-synthesized sample under different scan rates. Right: Specific capacitance at 1-10 A/g.

Capacitance data found in Table 1 indicate that washing of as-synthesized sample by KOH for Mg removal decreases its performance compared to the asprepared sample. It is due to the contribution of Mg metal traces that as-prepared sample exhibits higher performance compared to Mg-removed sample, although the latter should contain potassium ion traces in the sample.



Fig. 6. Left: Cyclic voltammogram of KOH-washed synthesized samples under different scan rates. Right: Specific capacitance at 1-10 A/g.

Table 1. Capacitance performance of the two samples.

Sample	Specific capacitance at following current densities (A/g)					
	1	2	3	4	5	10
As synthesized	192	138	102	90	75	60
KOH washed	117	87	65	60	46	33

4 Conclusions

The present research indicated that lignin can be converted into graphene material in one-pot synthesis using gel combustion method. The prepared samples showed promising performance for the application in the field of supercapacitance.

Further exploration on the graphene oxide doping with various metal dopants will enhance its supercapacitance performance. Those efforts will be helpful in commercializing the lignin biomass conversion process, which will expand the valorization of lignin biomass.

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