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Title: SUPERCAPACITOR ELECTRODES INCLUDING GRAPHENIC CARBON PARTICLES

Abstract: Supercapacitor electrodes comprising active charge supporting particles, graphenic carbon particles, and a binder are disclosed. The active charge supporting particles may comprise activated carbon. The graphenic carbon particles may be thermally produced. The electrodes may further comprise electrically conductive carbon.

FIG. 1
SUPERCAPACITOR ELECTRODES INCLUDING
GRAPHENIC CARBON PARTICLES

CROSS REFERENCE TO RELATED APPLICATION

[0001] This application claims the benefit of U.S. Provisional Patent Application Serial No. 62/073,298 filed October 31, 2014, which is incorporated herein by reference.

FIELD OF THE INVENTION

[0002] The present invention relates to the use of grapheme carbon particles in supercapacitor electrodes.

BACKGROUND OF THE INVENTION

[0003] A large demand exists for high-power energy resources for use in various products such as portable electronic devices and electric vehicles. Supercapacitors offer a promising alternative to conventional capacitors and batteries for such uses. Compared with conventional capacitors, the specific energy of supercapacitors can be several orders of magnitude higher. In addition, supercapacitors are able to store energy and deliver power at relatively high rates beyond those accessible with batteries.

SUMMARY OF THE INVENTION

[0004] An aspect of the invention provides a supercapacitor electrode comprising: a conductive foil substrate layer; and electrode coating layers on opposite sides of the conductive foil substrate layer, wherein each electrode coating layer comprises: active charge supporting particles; grapheme carbon particles; and a binder. A supercapacitor comprising such an electrode is also provided.

[0005] Another aspect of the invention provides an electrode coating comprising: from 50 to 95 weight percent active carbon particles; from 1 to 10 weight percent thermally produced grapheme carbon particles; and from 1 to 15 weight percent binder.
**BRIEF DESCRIPTION OF THE DRAWINGS**

[0006] Fig. 1 is a partially schematic side view of a supercapacitor in the form of a test coin cell including electrodes comprising grapheme carbon particles in accordance with an embodiment of the present invention.

[0007] Fig. 2 is a partially schematic side sectional view of a supercapacitor electrode in accordance with an embodiment of the present invention comprising a conductive foil layer with electrode coating layers on opposite sides thereof.

[0008] Fig. 3 is a graph illustrating linear charge and discharge characteristics of a supercapacitor test coin cell comprising electrodes with grapheme carbon particles as the active component of the electrode coating layers.

[0009] Fig. 4 is a graph of specific capacitance versus total coating loading for electrodes including activated carbon particles alone or in combination with grapheme carbon particles.

[0010] Fig. 5 is a graph of applied voltage versus time for electrodes including activated carbon particles and different types of grapheme carbon particles.

[0011] Fig. 6 is a graph of applied voltage versus time for electrodes including activated carbon particles and different types of grapheme carbon particles.

[0012] Fig. 7 is a Nyquist plot for electrodes including activated carbon particles alone or in combination with grapheme carbon particles.

**DETAILED DESCRIPTION**

[0013] Fig. 1 schematically illustrates a supercapacitor in the form of a test coin cell 10 including a stainless steel case 12 with two parts separated by an insulator ring, electrolyte 14, separator 16, spacers 18 and electrodes 20. In accordance with embodiments of the invention, one or more of the electrodes 20 may include grapheme carbon particles. The other components of the test coin cell capacitor 10, such as the electrolyte 14, separator 16 and spacers 18 may be made of any suitable conventional materials, as known to those skilled in the art.

[0014] As schematically shown in Fig. 2, a supercapacitor electrode 20 includes a conductive foil layer 22 having electrode coating layers 24 and 26 applied on opposite sides thereof. The conductive foil layer 22 may be made of any suitable conductive material such as aluminum or the like. The conductive foil layer 22 may
have any suitable thickness, for example, from 5 to 25 microns, or from 10 to 20 microns. In a particular embodiment, the conductive foil layer 22 comprises aluminum foil having a thickness of about 15 microns.

[0015] Each of the electrode layers 24 and 26 shown in Fig. 2 may have any suitable thicknesses, typically from 5 to 200 microns, e.g., from 10 to 120 microns, or from 20 to 100 microns. In the embodiment shown in Fig. 2, each of the electrode coating layers 24 and 26 have the same thicknesses. However, the layers 24 and 26 may have different thicknesses in certain embodiments.

[0016] In accordance with embodiments of the present invention, graphene carbon particles are included in the electrode coating layers 24 and 26. Graphene carbon particles have large specific surface area and exceptionally high electronic qualities, making such particles useful in supercapacitor applications. The graphene carbon particles may be combined with active charge supporting particles such as activated carbon and/or transition metal oxides for use in supercapacitor electrodes. Additional electrically conductive particles such as carbon black may optionally be added to such electrode materials. In certain embodiments, the electrode coating layers 24 and 26 include active charge supporting particles, graphene carbon particles, and a binder, as more fully described below.

[0017] As used herein, the term "supercapacitor" means capacitors having capacitance values greater than 1,000 farads at 1.2 volt. Supercapacitor electrodes of the present invention have been found to provide significantly improved capacitive properties such as specific capacitance. Specific capacitance is the capacitance per unit mass for one electrode and is calculated by the following equation:

$$C_{sp} \ (F/g) = \frac{4It}{Vm}$$

where I is the discharge current (A), t is the discharge duration (second), V is the voltage window (V) and m is the total mass (g) of the active material in both electrodes.

[0018] In accordance with certain embodiments, each supercapacitor electrode 20 has a specific capacitance of at least 75 F/g at a current density of 1 A/g for relatively thin electrodes with a mass loading of 1 mg/cm² of active material per electrode, for example, at least 100 F/g, or at least 110 F/g. Furthermore, in accordance with embodiments of the present invention, the supercapacitor electrodes
20 may have a typical specific energy density of at least 1 Wh/kg, or at least 2 Wh/kg. For example, the specific energy density of each electrode 20 may range from 2 to 20 Wh/kg or from 6 to 15 Wh/kg.

[0019] In certain embodiments, the electrode coating layers 24 and 26 of the supercapacitor electrodes 20 include active charge supporting particles comprising activated carbon. The activated carbon particles typically comprise from 50 to 95 weight percent of the electrode coating material, for example, from 60 to 90 percent activated carbon, or from 70 to 85 percent activated carbon. In a particular embodiment, the activated carbon comprises 80 weight percent of the electrode coating material. The activated carbon particles may be of any suitable size, for example, the average particle size of the activated carbon particles may typically range from 0.5 to 50 microns, or from 1 to 10 microns.

[0020] Graphene carbon particles may be added with the activated carbon particles in the conductive coating material in typical amounts of from 0.5 to 20 weight percent of the conductive coating material, for example, from 1 to 15 weight percent or from 2 to 10 weight percent.

[0021] While some commercial supercapacitors have used activated carbon, which has advantages of low cost and high capacitance at low current densities, it suffers from poor retention of specific capacitance at high current densities due to the low electrical conductivity deriving from its highly amorphous nature. This limits the power density of supercapacitors with activated carbon electrodes. In accordance with embodiments of the invention, the superior electrical conductivity and unique structure of graphene carbon particles make it a superior additive for activated carbon by acting as conductive bridges between activated carbon particles, thereby improving the conductivity and power density of activated carbon.

[0022] In addition to the activated carbon and graphene carbon particles, the electrode coating compositions may include a binder in a typical amount of up to 20 weight percent, for example, from 1-15 weight percent, or from 2-10 weight percent. Any suitable binder may be used, such as vinyls, latexes, acrylates, cellulosic binders, and conductive polymers. For example, the binder may comprise polyvinylidene fluoride (PVDF), styrene-butadiene rubber (SBR) latex, sodium carboxymethyl cellulose (CMC), polyaniline, polyacrylonitrile or the like. The binder may be used as
a solid in a powder coating formulation, or in a liquid coating formulation as a dispersion in a solvent or as a dissolved binder material.

[0023] In certain embodiments, conductive carbon black may be added with the activated carbon and graphene carbon particles in the conductive coating material in typical amounts of up to 20 weight percent based on the weight of the electrode coating composition, for example, from 1 to 15 weight percent, or from 2 to 10 weight percent, or from 5 to 8 weight percent. Typical average particle sizes of the conductive carbon black typically range from 0.5 to 50 microns, for example, from 1 to 10 microns.

[0024] In embodiments in which graphene carbon particles and conductive carbon black are used, the relative weight percentages thereof may be controlled. For example, the graphene carbon particles may comprise from 20 to 80 weight percent of the combined weight of the graphene carbon particles and conductive carbon black, for example, from 20 to 50 weight percent. The conductive carbon black may typically comprise from 20 to 80 weight percent of the combined weight of the graphene carbon particles and conductive carbon black, for example, from 50 to 80 weight percent.

[0025] In certain embodiments, the active charge supporting particles may comprise transition metal oxides such as manganese oxide (MnO$_2$), iron oxide (Fe$_2$O$_3$, Fe$_3$O$_4$), and the like. In this embodiment, the transition metal oxide particles may typically comprise 40 to 90 weight percent of the total weight of the electrode coating, for example, from 60 to 80 weight percent. When such transition metal oxides are used, the amount of graphene carbon particles may typically range from 10 to 60 weight percent, for example, from 20 to 40 weight percent. Typical average particle sizes of the transition metal oxide particles may range from 0.05 to 10 microns, for example, from 0.1 to 2 microns.

[0026] In certain embodiments, the active charge supporting particles, graphene carbon particles, binder and optional conductive carbon black particles are dispersed in a solvent and applied to the conductive foil substrate layer 22 by a coating process and converted into a dry film 24, 26 before being used in a supercapacitor. Conversion from a liquid formulation or a powder formulation into a dry and cured film may be accomplished by any suitable method such as oven heating.
[0027] In accordance with other embodiments of the invention, the grapheme carbon particles may be used as the sole active material of the electrode coatings the amount of grapheme carbon particles present in the supercapacitor electrode material, as a sole active material may typically range from 60 to 98 weight percent, for example, from 80 to 95 weight percent, or from 85 to 90 weight percent.

[0028] As used herein, the term "grapheme carbon particles" means carbon particles having structures comprising one or more layers of one-atom-thick planar sheets of sp²-bonded carbon atoms that are densely packed in a honeycomb crystal lattice. The average number of stacked layers may be less than 100, for example, less than 50. In certain embodiments, the average number of stacked layers is 30 or less, such as 20 or less, 10 or less, or, in some cases, 5 or less. The grapheme carbon particles may be substantially flat, however, at least a portion of the planar sheets may be substantially curved, curled, creased or buckled. The particles typically do not have a spheroidal or equiaxed morphology.

[0029] In certain embodiments, the grapheme carbon particles have a thickness, measured in a direction perpendicular to the carbon atom layers, of no more than 10 nanometers, no more than 5 nanometers, or, in certain embodiments, no more than 4 or 3 or 2 or 1 nanometers, such as no more than 3.6 nanometers. In certain embodiments, the grapheme carbon particles may be from 1 atom layer up to 3, 6, 9, 12, 20 or 30 atom layers thick, or more. In certain embodiments, the grapheme carbon particles have a width and length, measured in a direction parallel to the carbon atoms layers, of at least 50 nanometers, such as more than 100 nanometers, in some cases more than 100 nanometers up to 500 nanometers, or more than 100 nanometers up to 200 nanometers. The grapheme carbon particles may be provided in the form of ultrathin flakes, platelets or sheets having relatively high aspect ratios (aspect ratio being defined as the ratio of the longest dimension of a particle to the shortest dimension of the particle) of greater than 3:1, such as greater than 10:1.

[0030] In certain embodiments, the grapheme carbon particles have relatively low oxygen content. For example, the grapheme carbon particles may, even when having a thickness of no more than 5 or no more than 2 nanometers, have an oxygen content of no more than 2 atomic weight percent, such as no more than 1.5 or 1 atomic weight percent, or no more than 0.6 atomic weight, such as about 0.5 atomic
weight percent. The oxygen content of the grapheme carbon particles can be determined using X-ray Photoelectron Spectroscopy, such as is described in D. R. Dreyer et al., Chem. Soc. Rev. 39, 228-240 (2010).

[0031] In certain embodiments, the grapheme carbon particles have a B.E.T. specific surface area of at least 50 square meters per gram, such as 70 to 1000 square meters per gram, or, in some cases, 200 to 1000 square meters per grams or 200 to 400 square meters per gram. As used herein, the term "B.E.T. specific surface area" refers to a specific surface area determined by nitrogen adsorption according to the ASTMD 3663-78 standard based on the Brunauer-Emmett-Teller method described in the periodical "The Journal of the American Chemical Society", 60, 309 (1938).

[0032] In certain embodiments, the grapheme carbon particles have a Raman spectroscopy 2D/G peak ratio of at least 1:1, for example, at least 1.2:1 or 1.3:1. As used herein, the term "2D/G peak ratio" refers to the ratio of the intensity of the 2D peak at 2692 cm⁻¹ to the intensity of the G peak at 1,580 cm⁻¹.

[0033] In certain embodiments, the grapheme carbon particles have a relatively low bulk density. For example, the grapheme carbon particles are characterized by having a bulk density (tap density) of less than 0.2 g/cm³, such as no more than 0.1 g/cm³. For the purposes of the present invention, the bulk density of the grapheme carbon particles is determined by placing 0.4 grams of the grapheme carbon particles in a glass measuring cylinder having a readable scale. The cylinder is raised approximately one-inch and tapped 100 times, by striking the base of the cylinder onto a hard surface, to allow the grapheme carbon particles to settle within the cylinder. The volume of the particles is then measured, and the bulk density is calculated by dividing 0.4 grams by the measured volume, wherein the bulk density is expressed in terms of g/cm³.

[0034] In certain embodiments, the grapheme carbon particles have a compressed density and a percent densification that is less than the compressed density and percent densification of graphite powder and certain types of substantially flat graphite carbon particles such as those formed from exfoliated graphite. Lower compressed density and lower percent densification are each currently believed to contribute to better dispersion and/or rheological properties than grapheme carbon particles exhibiting higher compressed density and higher percent densification. In
certain embodiments, the compressed density of the graphene carbon particles is 0.9 or less, such as less than 0.8, less than 0.7, such as from 0.6 to 0.7. In certain embodiments, the percent densification of the graphene carbon particles is less than 40%, such as less than 30%, such as from 25 to 30%.

[0035] For purposes of the present invention, the compressed density of graphene carbon particles is calculated from a measured thickness of a given mass of the particles after compression. Specifically, the measured thickness is determined by subjecting 0.1 grams of the graphene carbon particles to cold press under 15,000 pound of force in a 1.3 centimeter die for 45 minutes, wherein the contact pressure is 500 MPa. The compressed density of the graphene carbon particles is then calculated from this measured thickness according to the following equation:

\[
\text{Compressed Density (g/cm}^3) = \frac{0.1 \text{ grams}}{n \times (1.3\text{cm}/2)^2 \times (\text{measured thickness in cm})}
\]

[0036] The percent densification of the graphene carbon particles is then determined as the ratio of the calculated compressed density of the graphene carbon particles, as determined above, to 2.2 g/cm³, which is the density of graphite.

[0037] In certain embodiments, the graphene carbon particles have a measured bulk liquid conductivity of at least 100 microSiemens, such as at least 120 microSiemens, such as at least 140 microSiemens immediately after mixing and at later points in time, such as at 10 minutes, or 20 minutes, or 30 minutes, or 40 minutes. For the purposes of the present invention, the bulk liquid conductivity of the graphene carbon particles is determined as follows. First, a sample comprising a 0.5% solution of graphene carbon particles in butyl cellosolve is sonicated for 30 minutes with a bath sonicator. Immediately following sonication, the sample is placed in a standard calibrated electrolytic conductivity cell (K=1). A Fisher Scientific AB 30 conductivity meter is introduced to the sample to measure the conductivity of the sample. The conductivity is plotted over the course of about 40 minutes.

[0038] In accordance with certain embodiments, percolation, defined as long range interconnectivity, occurs between the conductive graphene carbon particles. Such percolation may reduce the resistivity of the coating compositions. The
Conductive grapheme particles may occupy a minimum volume within the coating such that the particles form a continuous, or nearly continuous, network. In such a case, the aspect ratios of the grapheme carbon particles may affect the minimum volume required for percolation.

[0039] In certain embodiments, at least a portion of the grapheme carbon particles to be dispersed in the compositions of the present invention are may be made by thermal processes. In accordance with embodiments of the invention, thermally produced grapheme carbon particles are made from carbon-containing precursor materials that are heated to high temperatures in a thermal zone such as a plasma. As more fully described below, the carbon-containing precursor materials are heated to a sufficiently high temperature, e.g., above 3,500°C, to produce grapheme carbon particles having characteristics as described above. The carbon-containing precursor, such as a hydrocarbon provided in gaseous or liquid form, is heated in the thermal zone to produce the grapheme carbon particles in the thermal zone or downstream therefrom. For example, thermally produced grapheme carbon particles may be made by the systems and methods disclosed in U.S. Patent Nos. 8,486,363 and 8,486,364.

[0040] In certain embodiments, the thermally produced grapheme carbon particles may be made by using the apparatus and method described in U.S. Patent No. 8,486,363 at [0022] to [0048] in which (i) one or more hydrocarbon precursor materials capable of forming a two-carbon fragment species (such as n-propanol, ethane, ethylene, acetylene, vinyl chloride, 1,2-dichloroethane, allyl alcohol, propionaldehyde, and/or vinyl bromide) is introduced into a thermal zone (such as a plasma), and (ii) the hydrocarbon is heated in the thermal zone to form the grapheme carbon particles. In other embodiments, the thermally produced grapheme carbon particles may be made by using the apparatus and method described in U.S. Patent No. 8,486,364 at [0015] to [0042] in which (i) a methane precursor material (such as a material comprising at least 50 percent methane, or, in some cases, gaseous or liquid methane of at least 95 or 99 percent purity or higher) is introduced into a thermal zone (such as a plasma), and (ii) the methane precursor is heated in the thermal zone to form the grapheme carbon particles. Such methods can produce grapheme carbon particles having at least some, in some cases all, of the characteristics described above.
[0041] During production of the graphene carbon particles by the thermal production methods described above, a carbon-containing precursor is provided as a feed material that may be contacted with an inert carrier gas. The carbon-containing precursor material may be heated in a thermal zone, for example, by a plasma system. In certain embodiments, the precursor material is heated to a temperature of at least 3,500°C, for example, from a temperature of greater than 3,500°C or 4,000°C up to 10,000°C or 20,000°C. Although the thermal zone may be generated by a plasma system, it is to be understood that any other suitable heating system may be used to create the thermal zone, such as various types of furnaces including electrically heated tube furnaces and the like.

[0042] The gaseous stream may be contacted with one or more quench streams that are injected into the plasma chamber through at least one quench stream injection port. The quench stream may cool the gaseous stream to facilitate the formation or control the particle size or morphology of the graphene carbon particles. In certain embodiments of the invention, after contacting the gaseous product stream with the quench streams, the ultrafine particles may be passed through a converging member. After the graphene carbon particles exit the plasma system, they may be collected. Any suitable means may be used to separate the graphene carbon particles from the gas flow, such as, for example, a bag filter, cyclone separator or deposition on a substrate.

[0043] In certain embodiments, at least a portion of the graphene carbon particles may be obtained from commercial sources, for example, from Angstron, XG Sciences and other commercial sources. In such embodiments, the commercially available graphene carbon particles may comprise exfoliated graphite and have different characteristics in comparison with the thermally produced graphene carbon particles, such as different size distributions, thicknesses, aspect ratios, structural morphologies, oxygen contents, and chemical functionalities at the basal planes/edges.

[0044] In certain embodiments, different types of graphene carbon particles may be co-dispersed in the composition. For example, when thermally produced graphene carbon particles are combined with commercially available graphene carbon particles in accordance with embodiments of the invention, a bi-modal distribution, tri-modal distribution, etc. of graphene carbon particle characteristics
may be achieved. The grapheme carbon particles contained in the compositions may have multi-modal particle size distributions, aspect ratio distributions, structural morphologies, edge functionality differences, oxygen content, and the like.

[0045] In an embodiment of the present invention in which both thermally produced grapheme carbon particles and commercially available grapheme carbon particles, e.g., from exfoliated graphite, are co-dispersed and added to a coating composition to produce a bi-modal grapheme particle size distribution, the relative amounts of the different types of grapheme carbon particles are controlled to produce desired conductivity properties of the coatings. For example, the thermally produced grapheme particles may comprise from 1 to 50 weight percent, and the commercially available grapheme carbon particles may comprise from 50 to 99 weight percent, based on the total weight of the grapheme carbon particles. In certain embodiments, the thermally produced grapheme carbon particles may comprise from 2 or 4 to 40 weight percent, or from 6 or 8 to 35 weight percent, or from 10 to 30 weight percent. When co-dispersions of the present invention having such relative amounts of thermally produced grapheme carbon particles and commercially available grapheme carbon particles are incorporated in coatings, inks, or other materials, such materials may exhibit significantly increased electrical conductivities in comparison with similar materials containing mixtures of such types of grapheme carbon particles at similar ratios. For example, the co-dispersions may increase electrical conductivity by at least 10 or 20 percent compared with the mixtures. In certain embodiments, the electrical conductivity may be increased by at least 50, 70 or 90 percent, or more.

[0046] In certain embodiments, the grapheme carbon particles are functionalized. As used herein, "functionalized", when referring to grapheme carbon particles, means covalent bonding of any non-carbon atom or any organic group to the grapheme carbon particles. The grapheme carbon particles may be functionalized through the formation of covalent bonds between the carbon atoms of a particle and other chemical moieties such as carboxylic acid groups, sulfonic acid groups, hydroxyl groups, halogen atoms, nitro groups, amine groups, aliphatic hydrocarbon groups, phenyl groups and the like. For example, functionalization with carbonaceous materials may result in the formation of carboxylic acid groups on the grapheme carbon particles. The grapheme carbon particles may also be functionalized by other reactions such as Diels-Alder addition reactions, 1,3-dipolar cycloaddition reactions,
free radical addition reactions and diazonium addition reactions. In certain
embodiments, the hydrocarbon and phenyl groups may be further functionalized. If
the graphene carbon particles already have some hydroxyl functionality, the
functionality can be modified and extended by reacting these groups with, for
example, an organic isocyanate.

[0047] The following examples are intended to illustrate various aspects of the
invention, and are not intended to limit the scope of the invention.

Example 1

[0048] A coin cell configuration as shown in Fig. 1 was used to represent the
physical configuration, internal voltages and charge transfer that occurs in a packaged
supercapacitor in order to provide an indication of the performance of the electrode
material. Two identical electrodes with the same mass loading were used for testing. The
electrodes included thermally produced graphene carbon particles as the sole active
material (80 wt%), conductive carbon black (10 wt%) and a PVDF binder (10
wt%) applied on opposite sides of an aluminum foil layer. The electrolyte was KC1 in
water.

[0049] Fig. 3 shows the typical charge/discharge curve at 100 mA/g of
thermally produced graphene carbon particles with specific surface area (SSA) of 416
m²/g. The charge curve and discharge curves are nearly linear, indicating ideal
capacitive behaviors. The specific capacitance of thermally produced graphene
carbon particles is 25 F/g, which is lower than commercial active carbons with SSA
ranging from 1,000 to 3,500 m²/g. However, the theoretical specific capacitance of
the thermally produced graphene carbon particles is calculated to be 88 F/g
(416/2600*550), which is higher than the measured specific capacitance.

Example 2

[0050] To investigate the effects of thermally produced graphene carbon
particles in electrodes comprising activated carbon, six samples including those with
different ratios of thermally produced graphene carbon particles (TPGC) and Super C
conductive carbon black, and commercial graphene carbon particles as comparisons
(Table 1) were prepared and tested. The active charge supporting particles of each
electrode were commercially available YP-80F activated carbon particles. The specific capacitance is calculated based on the mass of activated carbon only.

[0051] Table 2 shows the performance of thin electrodes with mass loading of 1 mg/cm² per electrode and typical thickness of 15 µm. In general, introduction of graphene carbon particles increases specific capacitance and decreases internal resistance (IR) compared to pure conductive carbon at both low and high current densities. The difference is more striking at a high current density of 1 A/g. Sample 3 with 5 wt% thermally produced graphene carbon particles delivers a capacitance of 112 F/g, which doubles that of pure conductive carbon (55 F/g). Increasing the thermally produced graphene carbon particles to higher loading adversely affects the performance as demonstrated by sample 4 due to decreased conductivity. The preferred mass content of thermally produced graphene carbon particles should be in the range of about 2 to 5 weight percent. The electrode including commercially available M5 graphene produced acceptable results, but the electrode including commercially available C300 graphene produced less favorable results.

Table 1
Formulas of Electrode Coatings

<table>
<thead>
<tr>
<th>Sample</th>
<th>YP-80F (g)</th>
<th>Graphene (g)</th>
<th>Super C (g)</th>
<th>2 wt% PVDF/NMP (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.8</td>
<td>0</td>
<td>0.1</td>
<td>5</td>
</tr>
<tr>
<td>2</td>
<td>0.8</td>
<td>TPGC 0.02</td>
<td>0.08</td>
<td>5</td>
</tr>
<tr>
<td>3</td>
<td>0.8</td>
<td>TPGC 0.05</td>
<td>0.05</td>
<td>5</td>
</tr>
<tr>
<td>4</td>
<td>0.8</td>
<td>TPGC 0.08</td>
<td>0.02</td>
<td>5</td>
</tr>
<tr>
<td>5</td>
<td>0.8</td>
<td>M5 0.05</td>
<td>0.05</td>
<td>5</td>
</tr>
<tr>
<td>6</td>
<td>0.8</td>
<td>C300 0.05</td>
<td>0.05</td>
<td>5</td>
</tr>
</tbody>
</table>

Table 2
Performance of Thin Electrodes

<table>
<thead>
<tr>
<th>Sample</th>
<th>Capacitance (F/g)</th>
<th>IR drop (V)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0.1 A/g</td>
<td>1 A/g</td>
</tr>
<tr>
<td>1</td>
<td>102</td>
<td>55</td>
</tr>
<tr>
<td>2</td>
<td>145</td>
<td>108</td>
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<tr>
<td>3</td>
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<td>108</td>
</tr>
<tr>
<td>6</td>
<td>146</td>
<td>67</td>
</tr>
</tbody>
</table>
Example 3

[0052] Relatively thick electrodes were fabricated and evaluated. Carbon-coated Al foil (from Exopack) was used to enhance the adhesion. The electrodes have mass loadings of about 6.0 mg/cm$^2$ and thicknesses of about 115 $\mu$m. Overall, each electrode has lower capacitance at the same current density compared to its thin counterpart. As shown in Table 3 and Fig. 4, pure conductive carbon exhibits a low capacitance of 63.5 F/g at 1$^\text{st}$ cycle and has severe fading. In addition, the IR drop is large and increases upon cycling. In contrast, conductive carbon with thermally produced grapheme carbon particles shows higher capacitance with stable cyclability and much lower IR drop. Electrodes with thermally produced grapheme carbon particles also show higher capacitance at a high current density of 1A/g. Similar results were obtained with the electrodes including commercially available M5 and C300 graphene. The electrode with 8 weight percent thermally produced grapheme carbon particles has the lowest capacitance and highest IR drop among electrodes with thermally produced grapheme carbon particles, which confirms that the optimal thermally produced grapheme carbon particles content should range from about 2 to 5 weight percent.

Table 3

<table>
<thead>
<tr>
<th>Sample</th>
<th>Mass loading (mg/cm$^2$)</th>
<th>Capacitance (F/g)</th>
<th>IR drop (V)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>6.3</td>
<td>63.5 (1$^\text{st}$) 40 (30$^\text{th}$)</td>
<td>0.23 (1$^\text{st}$) 0.42 (30$^\text{th}$)</td>
</tr>
<tr>
<td>2</td>
<td>6.4</td>
<td>87</td>
<td>0.06</td>
</tr>
<tr>
<td>3</td>
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<td>87</td>
<td>0.08</td>
</tr>
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<td>0.07</td>
</tr>
<tr>
<td>6</td>
<td>6.0</td>
<td>84</td>
<td>0.1</td>
</tr>
</tbody>
</table>

Example 4

[0053] Thick electrodes of the same composition were also made and tested using a different carbon-coated Al foil (from MTI). As shown in Table 4, the difference between grapheme carbon particle-free and thermally produced grapheme carbon particle-containing electrodes is less pronounced in terms of capacitance.
However, the electrode with 2 weight percent thermally produced grapheme carbon particles (Sample 2) shows lower IR drop at both low and high current densities than grapheme carbon particle-free electrode (sample 1), indicating an improvement in conductivity of the electrode.

Table 4
Performance of Thick Electrodes

<table>
<thead>
<tr>
<th>Sample</th>
<th>Mass loading (mg/cm²)</th>
<th>Capacitance (F/g)</th>
<th>IR drop (V)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>0.1 A/g</td>
<td>1 A/g</td>
</tr>
<tr>
<td>1</td>
<td>5.8</td>
<td>90</td>
<td>80</td>
</tr>
<tr>
<td>2</td>
<td>5.8</td>
<td>90</td>
<td>80</td>
</tr>
<tr>
<td>3</td>
<td>6</td>
<td>93</td>
<td>74</td>
</tr>
<tr>
<td>4</td>
<td>5.9</td>
<td>83</td>
<td>45</td>
</tr>
<tr>
<td>5</td>
<td>6.3</td>
<td>84</td>
<td>71</td>
</tr>
<tr>
<td>6</td>
<td>6.1</td>
<td>88</td>
<td>53</td>
</tr>
</tbody>
</table>

[0054] Figs. 5 and 6 show charge/discharge curves of electrodes made of Samples 2 and 6 (Table 4). As shown in Fig. 5, at low current density (100 mA/g) both of the samples show symmetric charge and discharge cycles, but with Sample 2 showing a comparatively lesser IR drop than Sample 6. As shown in Fig. 6, at a higher current density (1 A/g), Sample 2 sustains the symmetric charge and discharge cycle profile (more linear), but the profile of Sample 6 turns asymmetric (and curved), which shows cyclic instability. Also, there is a significant drop in specific capacitance for Sample 6 which also confirms this behavior (see Table 4).

Example 5

[0055] Impedance spectroscopy was performed on electrodes containing activated carbon particles (Sample 1) and activated carbon particles with thermally produced grapheme carbon particles (Sample 2). The results are shown in the Nyquist plot of Fig. 7. With the addition of thermally produced grapheme carbon particles as an additive there is a decrease in impedance which indicates less resistance and leads to higher specific capacitance.

[0056] In accordance with embodiments of the invention, introducing thermally produced grapheme carbon particles effectively maintains the capacitance of activated carbon at high current densities and increases capacitance for thick
electrodes by increasing overall electrical conductivity of the whole electrodes. Such an effect becomes less pronounced when highly conductive substrate is used. Also, such improvements are not limited to thermally produced graphene carbon particles as commercial graphene carbon particles such as M5 can also achieve similar results.

For purposes of this detailed description, it is to be understood that the invention may assume various alternative variations and step sequences, except where expressly specified to the contrary. Moreover, other than in any operating examples, or where otherwise indicated, all numbers expressing, for example, quantities of ingredients used in the specification and claims are to be understood as being modified in all instances by the term "about". Accordingly, unless indicated to the contrary, the numerical parameters set forth in the following specification and attached claims are approximations that may vary depending upon the desired properties to be obtained by the present invention. At the very least, and not as an attempt to limit the application of the doctrine of equivalents to the scope of the claims, each numerical parameter should at least be construed in light of the number of reported significant digits and by applying ordinary rounding techniques.

Notwithstanding that the numerical ranges and parameters setting forth the broad scope of the invention are approximations, the numerical values set forth in the specific examples are reported as precisely as possible. Any numerical value, however, inherently contains certain errors necessarily resulting from the standard variation found in their respective testing measurements.

Also, it should be understood that any numerical range recited herein is intended to include all sub-ranges subsumed therein. For example, a range of "1 to 10" is intended to include all sub-ranges between (and including) the recited minimum value of 1 and the recited maximum value of 10, that is, having a minimum value equal to or greater than 1 and a maximum value of equal to or less than 10.

In this application, the use of the singular includes the plural and plural encompasses singular, unless specifically stated otherwise. In addition, in this application, the use of "or" means "and/or" unless specifically stated otherwise, even though "and/or" may be explicitly used in certain instances.

It will be readily appreciated by those skilled in the art that modifications may be made to the invention without departing from the concepts
disclosed in the foregoing description. Such modifications are to be considered as included within the following claims unless the claims, by their language, expressly state otherwise. Accordingly, the particular embodiments described in detail herein are illustrative only and are not limiting to the scope of the invention which is to be given the full breadth of the appended claims and any and all equivalents thereof.
WE CLAIM:

1. A supercapacitor electrode comprising:
   a conductive foil substrate layer; and
   electrode coating layers on opposite sides of the conductive foil
substrate layer, wherein each electrode coating layer comprises:
   active charge supporting particles;
   grapheme carbon particles; and
   a binder.

2. The supercapacitor electrode of Claim 1, wherein the active
   charge supporting particles comprise activated carbon or a transition metal oxide.

3. The supercapacitor electrode of Claim 1, wherein the active
   charge supporting particles comprise activated carbon in an amount of from 60 to 90
   weight percent of each of the electrode coating layers.

4. The supercapacitor electrode of Claim 3, wherein the grapheme
   carbon particles comprise thermally produced grapheme carbon particles.

5. The supercapacitor electrode of Claim 4, wherein the thermally
   produced grapheme carbon particles comprise from 1 to 10 weight percent of each of
   the electrode coating layers.

6. The supercapacitor electrode of Claim 4, wherein the thermally
   produced grapheme carbon particles comprise from 2 to 5 weight percent of each of
   the electrode coating layers.

7. The supercapacitor electrode of Claim 4, wherein the binder
   comprises from 1 to 15 weight percent of each of the electrode coating layers.

8. The supercapacitor electrode of Claim 4, wherein each of the
   electrode coating layers further comprises conductive carbon black in an amount of up
   to 20 weight percent of the electrode coating layer.

9. The supercapacitor electrode of Claim 8, wherein the weight
   ratio of the thermally produced grapheme carbon particles to the conductive carbon
   black is 50 percent or less.

10. The supercapacitor electrode of Claim 1, wherein the
    supercapacitor electrode has a specific capacitance of at least 100 F/g at a current
    density of 1 A/g.
11. The supercapacitor electrode of Claim 1, wherein the supercapacitor electrode has a specific capacitance of at least 110 F/g at a current density of 1 A/g.

12. The supercapacitor electrode of Claim 1, wherein the supercapacitor electrode has an energy density of at least 2 Wh/kg.


14. An electrode coating comprising:
   from 50 to 95 weight percent active carbon particles;
   from 1 to 10 weight percent thermally produced grapheme carbon particles; and
   from 1 to 15 weight percent binder.
FIG. 4