BINDER COMPOSITION FOR SECONDARY BATTERY

FIELD OF THE INVENTION

[001] The present invention relates to the field of batteries. In particular, this invention relates to a binder composition for lithium-ion batteries.

BACKGROUND OF THE INVENTION

[002] Over the past decades, lithium-ion batteries (LIBs) have come to be widely utilized in various applications, especially consumer electronics, because of their outstanding energy density, long cycle life and high discharging capability. Due to rapid market development of electric vehicles (EV) and grid energy storage, high-performance, low-cost LIBs are currently offering one of the most promising options for large-scale energy storage devices.

[003] Generally, lithium-ion battery electrodes are manufactured by casting an organic-based slurry onto a metallic current collector. The slurry contains electrode active material, conductive carbon, and binder in an organic solvent. The binder provides a good electrochemical stability, holds together the electrode active materials and adheres them to the current collector in the fabrication of electrodes. Polyvinylidene fluoride (PVDF) is one of the most commonly used binders in the commercial lithium-ion battery industry. However, PVDF is insoluble in water and can only dissolve in some specific organic solvents such as N-methyl-2-pyrrolidone (NMP) which is flammable and toxic and hence requires specific handling.

[004] An NMP recovery system must be in place during the drying process to recover NMP vapors. This will generate significant costs in the manufacturing process since it requires a large capital investment. The use of less expensive and more environmentally-friendly solvents, such as aqueous solvents, most commonly water, is preferred in the present invention since it can reduce the large capital cost of the recovery system.

[005] In view of such a problem, attempts have been made to replace conventional PVDF with more environmentally-friendly water-soluble binder materials or to make use of the known advantages of PVDF as a binder for electrode slurries without using organic solvents that require specific recovery treatment during manufacture.

[006] Known aqueous binders such as carboxymethyl cellulose (CMC) and styrene-butadiene rubber (SBR) exhibit only marginal adhesion capability and poor cycle life. SBR, in particular, requires a thickening agent in regulating the binder viscosity. Further, SBR exhibits high expandability and undesirable agglomeration characteristics resulting in inhomogeneous dispersion, high electrode resistance and poor performance. In addition to that, within the battery, cathodes are at high voltage. Most rubbers including SBR are only stable at the low

voltage of the anode and will decompose at high voltage. Therefore, their applications, especially in cathodes, are somewhat limited.

[007] EP Patent Application Publication No. 2555293 B1 discloses a water-based electrode slurry for a lithium ion-containing electrochemical cell. The slurry comprises a combination of PVDF and SBR and at least one of polyacrylic acid (PAA) and CMC in an aqueous solution and an electrochemically active material. The proposed invention attempts to combine PVDF with a water-based slurry which allows easier handling and less environmental pollution and reduced costs while keeping the known chemical and electrochemical advantages of PVDF as a binder i.e. electrochemical stability, lifetime stability, reduced binder content enables higher C-rates, etc. Despite the fact that organic solvent free slurries could be prepared based on the proposed invention, the slurry consists of fluorine-containing binder material nonetheless. PVDF is highly fluorinated and toxic when exposed to thermal decomposition, which pose risks to the health of people and the environment.

[008] In view of the above, there is always a need for an aqueous binder composition for lithium-ion batteries that exhibits excellent adhesion capability and high electrochemical stability with such properties sustained in the making of a cathode slurry, and thus contribute to exceptional battery electrochemical performance.

SUMMARY OF THE INVENTION

[009] The aforementioned needs are met by various aspects and embodiments disclosed herein. Provided herein is a binder composition for a secondary battery electrode comprising a copolymer and a dispersion medium, wherein the copolymer comprises a structural unit (a) derived from a carboxylic acid group-containing monomer, a structural unit (b) derived from an amide group-containing monomer and a structural unit (c) derived from a nitrile group-containing monomer, with an improved binding capability. In addition, battery cells comprising the cathode prepared using the binder composition disclosed herein exhibits exceptional electrochemical performance.

BRIEF DESCRIPTION OF THE DRAWINGS

[0010] Figure 1 is a flow chart of an embodiment illustrating the steps for preparing a binder composition.

DETAILED DESCRIPTION OF THE INVENTION

[0011] Provided herein is a binder composition for a secondary battery electrode comprising a copolymer and a dispersion medium, wherein the copolymer comprises a structural unit (a) derived from a carboxylic acid group-containing monomer, a structural unit (b) derived from an amide group-containing monomer and a structural unit (c) derived from a nitrile group-containing monomer.

[0012] The term “electrode” refers to a “cathode” or an “anode.”

[0013] The term “positive electrode” is used interchangeably with cathode. Likewise, the term “negative electrode” is used interchangeably with anode.

[0014] The term “binder”, “binder material” or “binder composition” refers to a chemical compound, mixture of compounds, or polymer which form colloidal solutions or colloidal dispersions in a dispersion medium such as water, and is used to hold an electrode material and/or a conductive agent in place and adhere them onto a conductive metal part to form an electrode. In some embodiments, the electrode does not comprise any conductive agent.

[0015] The term “conductive agent” refers to a material which is chemically inactive and has good electrical conductivity. Therefore, the conductive agent is often mixed with an electrode active material at the time of forming an electrode to improve electrical conductivity of the electrode.

[0016] The term “polymer” refers to a polymeric compound prepared by polymerizing monomers, whether of the same or a different type. The generic term “polymer” embraces the terms “homopolymer” as well as “copolymer”.

[0017] The term “homopolymer” refers to a polymer prepared by the polymerization of the same type of monomer.

[0018] The term “copolymer” refers to a polymer prepared by the polymerization of two or more different types of monomers.

[0019] The term “unsaturated” as used herein, refers to a moiety having one or more units of unsaturation.

[0020] The term “alkyl” or “alkyl group” refers to a univalent group having the general formula C_nH_{2n+1} derived from removing a hydrogen atom from a saturated, unbranched or branched aliphatic hydrocarbon, where n is an integer, or an integer between 1 and 20, or between 1 and 8. Examples of alkyl groups include, but are not limited to, (C_1–C_8)alkyl groups, such as methyl, ethyl, propyl, isopropyl, 2-methyl-1-propyl, 2-methyl-2-propyl, 2-methyl-1-butyl, 3-methyl-1-butyl, 2-methyl-3-butyl, 2,2-dimethyl-1-propyl, 2-methyl-1-pentyl, 3-methyl-1-pentyl, 4-methyl-1-pentyl, 2-methyl-2-pentyl, 3-methyl-2-pentyl, 4-methyl-2-pentyl, 2,2-dimethyl-1-butyl, 3,3-dimethyl-1-butyl, 2-ethyl-1-butyl, butyl, isobutyl, *t*–butyl, pentyl, isopentyl, neopentyl, hexyl, heptyl and octyl. Longer alkyl groups include nonyl and decyl groups. An alkyl group can be unsubstituted or substituted with one or more suitable substituents. Furthermore, the alkyl group can be branched or unbranched. In some embodiments, the alkyl group contains at least 2, 3, 4, 5, 6, 7, or 8 carbon atoms.

[0021] The term “cycloalkyl” or “cycloalkyl group” refers to a saturated or unsaturated cyclic non-aromatic hydrocarbon radical having a single ring or multiple condensed rings.

Examples of cycloalkyl groups include, but are not limited to, (C_3-C_7)cycloalkyl groups, such as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, and cycloheptyl, and saturated cyclic and bicyclic terpenes and (C_3-C_7)cycloalkenyl groups, such as cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclohexenyl, and cycloheptenyl, and unsaturated cyclic and bicyclic terpenes. A cycloalkyl group can be unsubstituted or substituted by one or two suitable substituents. Furthermore, the cycloalkyl group can be monocyclic or polycyclic. In some embodiments, the cycloalkyl group contains at least 5, 6, 7, 8, 9, or 10 carbon atoms.

[0022] The term "alkoxy" refers to an alkyl group, as previously defined, attached to the principal carbon chain through an oxygen atom. Some non-limiting examples of the alkoxy group include methoxy, ethoxy, propoxy, butoxy, and the like. And the alkoxy defined above may be substituted or unsubstituted, wherein the substituent may be, but is not limited to, deuterium, hydroxy, amino, halo, cyano, alkoxy, alkyl, alkenyl, alkynyl, mercapto, nitro, and the like.

[0023] The term "alkenyl" refers to an unsaturated straight chain, branched chain, or cyclic hydrocarbon radical that contains one or more carbon-carbon double bonds. Examples of alkenyl groups include, but are not limited to, ethenyl, 1-propenyl, or 2-propenyl, which may optionally be substituted on one or more of the carbon atoms of the radical.

[0024] The term "aryl" or "aryl group" refers to an organic radical derived from a monocyclic or polycyclic aromatic hydrocarbon by removing a hydrogen atom. Non-limiting examples of the aryl group include phenyl, naphthyl, benzyl, or tolanyl group, sexiphenylene, phenanthrenyl, anthracenyl, coronenyl, and tolanylphenyl. An aryl group can be unsubstituted or substituted with one or more suitable substituents. Furthermore, the aryl group can be monocyclic or polycyclic. In some embodiments, the aryl group contains at least 6, 7, 8, 9, or 10 carbon atoms.

[0025] The term "aliphatic" refers to a C_1 to C_{30} alkyl group, a C_2 to C_{30} alkenyl group, a C_2 to C_{30} alkynyl group, a C_1 to C_{30} alkylene group, a C_2 to C_{30} alkenylene group, or a C_2 to C_{30} alkynylene group. In some embodiments, the alkyl group contains at least 2, 3, 4, 5, 6, 7, or 8 carbon atoms.

[0026] The term "aromatic" refers to groups comprising aromatic hydrocarbon rings, optionally including heteroatoms or substituents. Examples of such groups include, but are not limited to, phenyl, tolyl, biphenyl, o-terphenyl, m-terphenyl, p-terphenyl, naphthyl, anthryl, phenanthryl, pyrenyl, triphenylenyl, and derivatives thereof.

[0027] The term "substituted" as used to describe a compound or chemical moiety refers to that at least one hydrogen atom of that compound or chemical moiety is replaced with a second chemical moiety. Examples of substituents include, but are not limited to, halogen; alkyl; heteroalkyl; alkenyl; alkynyl; aryl, heteroaryl, hydroxyl; alkoxyl; amino; nitro; thiol; thioether;

imine; cyano; amido; phosphonato; phosphine; carboxyl; thiocarbonyl; sulfonyl; sulfonamide; acyl; formyl; acyloxy; alkoxycarbonyl; oxo; haloalkyl (*e.g.*, trifluoromethyl); carbocyclic cycloalkyl, which can be monocyclic or fused or non-fused polycyclic (*e.g.*, cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl) or a heterocycloalkyl, which can be monocyclic or fused or non-fused polycyclic (*e.g.*, pyrrolidinyl, piperidinyl, piperazinyl, morpholinyl or thiazinyl); carbocyclic or heterocyclic, monocyclic or fused or non-fused polycyclic aryl (*e.g.*, phenyl, naphthyl, pyrrolyl, indolyl, furanyl, thiophenyl, imidazolyl, oxazolyl, isoxazolyl, thiazolyl, triazolyl, tetrazolyl, pyrazolyl, pyridinyl, quinolinyl, isoquinolinyl, acridinyl, pyrazinyl, pyridazinyl, pyrimidinyl, benzimidazolyl, benzothiophenyl or benzofuranyl); amino (primary, secondary or tertiary); o-lower alkyl; o-aryl, aryl; aryl-lower alkyl; $-CO_2CH_3$; $-CONH_2$; $-OCH_2CONH_2$; $-NH_2$; $-SO_2NH_2$; $-OCHF_2$; $-CF_3$; $-OCF_3$; –NH(alkyl); $–N(alkyl)_2$; –NH(aryl); –N(alkyl)(aryl); $–N(aryl)_2$; –CHO; –CO(alkyl); -CO(aryl); $-CO_2$(alkyl); and $–CO_2$(aryl); and such moieties can also be optionally substituted by a fused-ring structure or bridge, for example $-OCH_2O-$. These substituents can optionally be further substituted with a substituent selected from such groups. All chemical groups disclosed herein can be substituted, unless it is specified otherwise.

[0028] The term “halogen” or “halo” refers to F, Cl, Br or I.

[0029] The term “monomeric unit” refers to the constitutional unit contributed by a single monomer to the structure of a polymer.

[0030] The term “structural unit” refers to the total monomeric units contributed by the same monomer type in a polymer.

[0031] The term “carboxylic salt group” refers to the carboxylate salt formed when a carboxylic acid reacts with a base. In some embodiments, the proton of the carboxylic acid is replaced with a metal cation. In some embodiments, the proton of the carboxylic acid is replaced with an ammonium ion.

[0032] The term “applying” refers to an act of laying or spreading a substance on a surface.

[0033] The term “current collector” refers to any conductive substrate, which is in contact with an electrode layer and is capable of conducting an electrical current flowing to electrodes during discharging or charging a secondary battery. Some non-limiting examples of the current collector include a single conductive metal layer or substrate and a single conductive metal layer or substrate with an overlying conductive coating layer, such as a carbon black-based coating layer. The conductive metal layer or substrate may be in the form of a foil or a porous body having a three-dimensional network structure, and may be a polymeric or metallic material or a metalized polymer. In some embodiments, the three-dimensional porous current collector is covered with a conformal carbon layer.

[0034] The term “electrode layer” refers to a layer, which is in contact with a current collector, that comprises an electrochemically active material. In some embodiments, the electrode layer is made by applying a coating on to the current collector. In some embodiments, the electrode layer is located on the surface of the current collector. In other embodiments, the three-dimensional porous current collector is coated conformally with an electrode layer.

[0035] The term “room temperature” refers to indoor temperatures from about 18 °C to about 30 °C, e.g., 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, or 30 °C. In some embodiments, room temperature refers to a temperature of about 20 °C +/- 1 °C or +/- 2 °C or +/- 3 °C. In other embodiments, room temperature refers to a temperature of about 22 °C or about 25 °C.

[0036] The term “particle size D50” refers to a volume-based accumulative 50% size (D50), which is a particle size at a point of 50% on an accumulative curve (i.e., a diameter of a particle in the 50th percentile (median) of the volumes of particles) when the accumulative curve is drawn so that a particle size distribution is obtained on the volume basis and the whole volume is 100%. Further, with respect to the cathode active material of the present invention, the particle size D50 means a volume-averaged particle size of secondary particles which can be formed by mutual agglomeration of primary particles, and in a case where the particles are composed of the primary particles only, it means a volume-averaged particle size of the primary particles.

[0037] The term “polydispersity index” or “PDI” refers to a ratio of the weight average molecular weight (M_w) relative to the number average molecular weight (M_n). It is a measure of the distribution of the molecular weight within a given binder composition sample.

[0038] The term “solid content” refers to the amount of non-volatile material remaining after evaporation.

[0039] The term “mean roughness depth” or “R_z” refers to the arithmetic mean value of the single roughness depths of consecutive sampling lengths of a current collector.

[0040] The term “peeling strength” refers to the amount of force required to separate a current collector and an electrode active material coating that are bonded to each other. It is a measure of the binding strength between such two materials and is usually expressed in N/cm.

[0041] The term “adhesive strength” refers to the amount of force required to separate a current collector and a binder composition coating that are bonded to each other. It is a measure of the adhesion strength between such two materials and is usually expressed in N/cm.

[0042] The term “swelling” refers to a volumetric evolution of binder composition after soaking in the electrolyte or the uptake of electrolyte due to electrolyte-binder interactions.

[0043] The term “C rate” refers to the charging or discharging rate of a cell or battery, expressed in terms of its total storage capacity in Ah or mAh. For example, a rate of 1 C means utilization of all of the stored energy in one hour; a 0.1 C means utilization of 10% of the energy

in one hour or full energy in 10 hours; and a 5 C means utilization of full energy in 12 minutes.

[0044] The term “ampere-hour (Ah)” refers to a unit used in specifying the storage capacity of a battery. For example, a battery with 1 Ah capacity can supply a current of one ampere for one hour or 0.5 A for two hours, etc. Therefore, 1 ampere-hour (Ah) is the equivalent of 3,600 coulombs of electrical charge. Similarly, the term “miniampere-hour (mAh)” also refers to a unit of the storage capacity of a battery and is 1/1,000 of an ampere-hour.

[0045] The term “battery cycle life” refers to the number of complete charge/discharge cycles a battery can perform before its nominal capacity falls below 80% of its initial rated capacity.

[0046] The term “capacity” is a characteristic of an electrochemical cell that refers to the total amount of electrical charge an electrochemical cell, such as a battery, is able to hold. Capacity is typically expressed in units of ampere-hours. The term “specific capacity” refers to the capacity output of an electrochemical cell, such as a battery, per unit weight, usually expressed in Ah/kg or mAh/g.

[0047] In the following description, all numbers disclosed herein are approximate values, regardless whether the word “about” or “approximate” is used in connection therewith. They may vary by 1 percent, 2 percent, 5 percent, or, sometimes, 10 to 20 percent. Whenever a numerical range with a lower limit, R^L, and an upper limit, R^U, is disclosed, any number falling within the range is specifically disclosed. In particular, the following numbers within the range are specifically disclosed: $R=R^L+k*(R^U-R^L)$, wherein k is a variable ranging from 0 percent to 100 percent. Moreover, any numerical range defined by two R numbers as defined in the above is also specifically disclosed.

[0048] Currently, cathodes are often prepared by dispersing a cathode active material, a binder material and a conductive agent in an organic solvent such as N-methyl-2-pyrrolidone (NMP) to form a cathode slurry, then coating the cathode slurry onto a current collector and drying it.

[0049] Very often the binders influence on cell performance is underestimated, as they are considered as electrochemically inactive materials. The aim of a binder is to adhere the active material particles and the conductive agent together to form a continuous electrical conduction path to the current collector. Along with the binding capability, a binder material should be capable of facilitating electron and ion transportation to reduce the impedance between the current collector and the electrode materials and have sufficient elasticity to prevent the electrode from swelling due to volume expansion and contraction during charging and discharging.

[0050] Polyvinylidene fluoride (PVDF) has been widely used as a binder material in the

production of lithium-ion batteries. However, PVDF can only dissolve in specific organic solvents such as NMP which is flammable and toxic and hence require specific handling. An NMP recovery system must be in place during the drying process to recover NMP vapors. This will generate significant energy consumption and production costs in the manufacturing process. Accordingly, exploring a novel environmentally-friendly binder material to replace PVDF has become imperative in the development of lithium-ion battery binder materials.

[0051] Carboxymethyl cellulose (CMC) and styrene butadiene rubber (SBR) are some of the typical aqueous binders that have already been used in large-scale commercial applications. However, these binders have limited binding strengths and capabilities in preventing electrode swelling. Furthermore, within the battery, cathodes are at high voltage. Most rubbers including SBR are only stable at the low voltage of the anode and will decompose at high voltage. Therefore, their applications, especially in cathodes, are somewhat limited.

[0052] Accordingly, the present invention provides a method of preparing an aqueous binder composition comprising a copolymer and a dispersion medium, wherein the copolymer comprises a structural unit (a) derived from a carboxylic acid group-containing monomer, a structural unit (b) derived from an amide group-containing monomer and a structural unit (c) derived from a nitrile group-containing monomer. Figure 1 is a flow chart of an embodiment illustrating the steps of method **100** for preparing a binder composition. It is found that the binder composition described herein exhibits an enhanced adhesive capability and simultaneously has the unexpected effect of improving the capacity and electrochemical performance of cathodes formed therefrom.

[0053] In some embodiments, the binder composition described herein is produced via polymerization that involves monomers, polymers or monomer-polymer complexes being dispersed in an aqueous phase, with generation of free radicals with a water-soluble free radical initiator.

[0054] In some embodiments, a neutralizing solution is prepared by dissolving the neutralizing agent in water. In some embodiments, the first suspension is formed by adding the neutralizing solution in a dispersion medium in step **101**. Addition of neutralizing solution aims to improve polymerization stability and provide a pH range in which initiator that is added at a later stage is capable of generating free radicals.

[0055] Establishing a desirable range of pH to work with is especially important in aqueous systems. Neutralizing agents are commonly used for pH adjustment. In some embodiments, the neutralizing agent include alkaline aqueous solutions. In some embodiments, the neutralizing agent may be selected from the group consisting of ammonia, sodium bicarbonate, sodium hydroxide, potassium hydroxide, lithium hydroxide, ammonium hydroxide, magnesium hydroxide, calcium hydroxide, triethylamine, dimethylethanolamine (DMEA),

sodium carbonate, lithium carbonate, lithium bicarbonate and combinations thereof.

[0056] A dispersion medium is used as a solvent for the free radical initiators, neutralizing agents and other constituents. In some embodiments, the binder composition disclosed herein is prepared by an aqueous processing method in which water is used as the dispersion medium.

[0057] In some embodiments, the dispersion medium can further comprise a hydrophilic solvent selected from the group consisting of ethanol, isopropanol, n-propanol, tert-butanol, n-butanol, dimethylacetamide (DMAc), dimethylformamide (DMF), N-methylpyrrolidone (NMP), methyl ethyl ketone (MEK), ethyl acetate (EA), butyl acetate (BA) and combinations thereof. In some embodiments, the dispersion medium is free of water, ethanol, isopropanol, n-propanol, tert-butanol, n-butanol, dimethylacetamide (DMAc), dimethylformamide (DMF), N-methylpyrrolidone (NMP), methyl ethyl ketone (MEK), ethyl acetate (EA) or butyl acetate (BA).

[0058] In some embodiments, the first suspension is stirred for a time period of from about 5 minutes to about 45 minutes, from about 5 minutes to about 40 minutes, from about 5 minutes to about 35 minutes, from about 5 minutes to about 30 minutes, from about 5 minutes to about 25 minutes, from about 5 minutes to about 20 minutes or from about 10 minutes to about 20 minutes. In some embodiments, the first suspension is stirred for a time period of less than 45 minutes, less than 40 minutes, less than 35 minutes, less than 30 minutes, less than 25 minutes, less than 20 minutes, less than 15 minutes or less than 10 minutes. In some embodiments, the first suspension is stirred for a time period of more than 5 minutes, more than 10 minutes, more than 15 minutes, more than 20 minutes, more than 25 minutes, more than 30 minutes, more than 35 minutes or more than 40 minutes.

[0059] In some embodiments, the first suspension is stirred at a speed of from about 10 rpm to about 600 rpm, from about 50 rpm to about 600 rpm, from about 100 rpm to about 600 rpm, from about 150 rpm to about 600 rpm, from about 200 rpm to about 600 rpm, from about 250 rpm to about 600 rpm, from about 300 rpm to about 600 rpm, from about 300 rpm to about 550 rpm, from about 300 rpm to about 500 rpm, from about 320 rpm to about 480 rpm, from about 340 rpm to about 460 rpm or from about 360 rpm to about 440 rpm. In some embodiments, the first suspension is stirred at a speed of less than 600 rpm, less than 550 rpm, less than 500 rpm, less than 450 rpm, less than 400 rpm, less than 350 rpm, less than 300 rpm, less than 250 rpm, less than 200 rpm, less than 150 rpm, less than 100 rpm or less than 50 rpm. In some embodiments, the first suspension is stirred at a speed of more than 10 rpm, more than 50 rpm, more than 100 rpm, more than 150 rpm, more than 200 rpm, more than 250 rpm, more than 300 rpm, more than 350 rpm, more than 400 rpm, more than 450 rpm, more than 500 rpm or more than 550 rpm.

[0060] In some embodiments, the second suspension is formed by adding a carboxylic

acid group-containing monomer into the first suspension in step **102**.

[0061] Structural unit (a) is derived from a carboxylic acid group-containing monomer. Any monomer that has at least one carboxylic acid group may be used as carboxylic acid group-containing monomer without any specific limitations. In some embodiments, the carboxylic acid group-containing monomer is acrylic acid, methacrylic acid, crotonic acid, 2-butyl crotonic acid, cinnamic acid, maleic acid, maleic anhydride, fumaric acid, itaconic acid, itaconic anhydride, tetraconic acid or a combination thereof. In certain embodiments, the carboxylic acid group-containing monomer is 2-ethylacrylic acid, isocrotonic acid, cis-2-pentenoic acid, trans-2-pentenoic acid, angelic acid, tiglic acid, 3,3-dimethyl acrylic acid, 3-propyl acrylic acid, trans-2-methyl-3-ethyl acrylic acid, cis-2-methyl-3-ethyl acrylic acid, 3-isopropyl acrylic acid, trans-3-methyl-3-ethyl acrylic acid, cis-3-methyl-3-ethyl acrylic acid, 2-isopropyl acrylic acid, trimethyl acrylic acid, 2-methyl-3,3-diethyl acrylic acid, 3-butyl acrylic acid, 2-butyl acrylic acid, 2-pentyl acrylic acid, 2-methyl-2-hexenoic acid, trans-3-methyl-2-hexenoic acid, 3-methyl-3-propyl acrylic acid, 2-ethyl-3-propyl acrylic acid, 2,3-diethyl acrylic acid, 3,3-diethyl acrylic acid, 3-methyl-3-hexyl acrylic acid, 3-methyl-3-tert-butyl acrylic acid, 2-methyl-3-pentyl acrylic acid, 3-methyl-3-pentyl acrylic acid, 4-methyl-2-hexenoic acid, 4-ethyl-2-hexenoic acid, 3-methyl-2-ethyl-2-hexenoic acid, 3-tert-butyl acrylic acid, 2,3-dimethyl-3-ethyl acrylic acid, 3,3-dimethyl-2-ethyl acrylic acid, 3-methyl-3-isopropyl acrylic acid, 2-methyl-3-isopropyl acrylic acid, trans-2-octenoic acid, cis-2-octenoic acid, trans-2-decenoic acid, α-acetoxyacrylic acid, β-trans-aryloxyacrylic acid, α-chloro-β-E-methoxyacrylic acid or a combination thereof. In some embodiments, the carboxylic acid group-containing monomer is methyl maleic acid, dimethyl maleic acid, phenyl maleic acid, bromo maleic acid, chloromaleic acid, dichloromaleic acid, fluoromaleic acid, difluoro maleic acid, nonyl hydrogen maleate, decyl hydrogen maleate, dodecyl hydrogen maleate, octadecyl hydrogen maleate, fluoroalkyl hydrogen maleate or a combination thereof. In some embodiments, the carboxylic acid group-containing monomer is maleic anhydride, methyl maleic anhydride, dimethyl maleic anhydride, acrylic anhydride, methacrylic anhydride, methacrolein, methacryloyl chloride, methacryloyl fluoride, methacryloyl bromide, or a combination thereof.

[0062] In some embodiments, the proportion of the carboxylic acid group-containing monomer is from about 10% to about 30%, from about 10% to about 25%, from about 10% to about 20%, from about 10% to about 15%, from about 11% to about 30%, from about 12% to about 30%, from about 13% to about 30%, from about 14% to about 30%, from about 15% to about 30%, from about 15% to about 25%, from about 15% to about 20%, from about 15% to about 29%, from about 15% to about 28%, from about 15% to about 27%, from about 15% to about 26%, from about 15% to about 25%, from about 16% to about 25%, from about 17% to about 25%, from about 18% to about 25%, from about 19% to about 25%, from about 20% to about 30%, from about 20% to about 25%, from about 17% to about 23%, from about 15% to

about 20% or from about 17% to about 26% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[0063] In some embodiments, the proportion of the carboxylic acid group-containing monomer is less than 30%, less than 29%, less than 28%, less than 27%, less than 26%, less than 25%, less than 24%, less than 23%, less than 22%, less than 21%, less than 20%, less than 19%, less than 18%, less than 17%, less than 16%, less than 15%, less than 14%, less than 13% or less than 12% by weight, based on the total weight of monomers added in the preparation of the binder composition. In some embodiments, the proportion of the carboxylic acid group-containing monomer is more than 10%, more than 11%, more than 12%, more than 13%, more than 14%, more than 15%, more than 16%, more than 17%, more than 18%, more than 19%, more than 20%, more than 21%, more than 22%, more than 23%, more than 24%, more than 25%, more than 26%, more than 27%, more than 28% or more than 29% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[0064] In some embodiments, more than one carboxylic acid group-containing monomers may be added into the first suspension in step **102** to form a second suspension. This may be advantageous as it allows better dispersion and prevents the sedimentation of materials in the processing of the binder composition disclosed herein. It is found that addition of more than one carboxylic acid group-containing monomers to the binder composition enhances a higher monomer conversion rate, thus maximizes the full use of starting materials and simultaneously could bring about significant costing savings and environmental benefits. In addition, a slightly better battery electrochemical performance is subsequently observed with the application of more than one carboxylic acid group-containing monomers in the preparation of the binder composition.

[0065] In some embodiments, the copolymer comprises a structural unit (a1) derived from a first carboxylic acid group-containing monomer, a structural unit (a2) derived from a second carboxylic acid group-containing monomer, a structural unit (b) derived from an amide group-containing monomer and a structural unit (c) derived from a nitrile group-containing monomer. In some embodiments, a first carboxylic acid group-containing monomer and a second carboxylic acid group-containing monomer may be added into the first suspension in step **102** to form a second suspension.

[0066] In some embodiments, the copolymer consists of a structural unit (a1) derived from a first carboxylic acid group-containing monomer, a structural unit (a2) derived from a second carboxylic acid group-containing monomer, a structural unit (b) derived from an amide group-containing monomer and a structural unit (c) derived from a nitrile group-containing monomer.

[0067] In some embodiments, structural unit (a1) is derived from a first carboxylic acid

group-containing monomer. In some embodiments, the first carboxylic acid group-containing monomer is acrylic acid.

[0068] In some embodiments, structural unit (a2) is derived from a second carboxylic acid group-containing monomer. In some embodiments, the second carboxylic acid group-containing monomer is an alkyl group substituted acrylic acid. In some embodiments, the second carboxylic acid group-containing monomer is methacrylic acid, crotonic acid, 2-butyl crotonic acid, 2-ethylacrylic acid, isocrotonic acid, cis-2-pentenoic acid, trans-2-pentenoic acid, angelic acid, tiglic acid, 3,3-dimethyl acrylic acid, 3-propyl acrylic acid, trans-2-methyl-3-ethyl acrylic acid, cis-2-methyl-3-ethyl acrylic acid, 3-isopropyl acrylic acid, trans-3-methyl-3-ethyl acrylic acid, cis-3-methyl-3-ethyl acrylic acid, 2-isopropyl acrylic acid, trimethyl acrylic acid, 2-methyl-3,3-diethyl acrylic acid, 3-butyl acrylic acid, 2-butyl acrylic acid, 2-pentyl acrylic acid, 2-methyl-2-hexenoic acid, trans-3-methyl-2-hexenoic acid, 3-methyl-3-propyl acrylic acid, 2-ethyl-3-propyl acrylic acid, 2,3-diethyl acrylic acid, 3,3-diethyl acrylic acid, 3-methyl-3-hexyl acrylic acid, 3-methyl-3-tert-butyl acrylic acid, 2-methyl-3-pentyl acrylic acid, 3-methyl-3-pentyl acrylic acid, 4-methyl-2-hexenoic acid, 4-ethyl-2-hexenoic acid, 3-methyl-2-ethyl-2-hexenoic acid, 3-tert-butyl acrylic acid, 2,3-dimethyl-3-ethyl acrylic acid, 3,3-dimethyl-2-ethyl acrylic acid, 3-methyl-3-isopropyl acrylic acid, 2-methyl-3-isopropyl acrylic acid, trans-2-octenoic acid, cis-2-octenoic acid, trans-2-decenoic acid, or a combination thereof.

[0069] In some embodiments, the proportion of the first carboxylic acid group-containing monomer is from about 5% to about 30%, from about 5.5% to about 30%, from about 6% to about 30%, from about 6.5% to about 30%, from about 7% to about 30%, from about 7.5% to about 30%, from about 8% to about 30%, from about 8.5% to about 30%, from about 9% to about 30%, from about 9.5% to about 30%, from about 10% to about 30%, from about 10% to about 29.5%, from about 10% to about 29%, from about 10% to about 28.5%, from about 10% to about 28%, from about 10% to about 27.5%, from about 10% to about 27%, from about 10% to about 26.5%, from about 10% to about 26%, from about 10% to about 25.5%, from about 10% to about 25%, from about 10% to about 24.5%, from about 10% to about 24%, from about 10% to about 23.5%, from about 10% to about 23%, from about 10% to about 22.5%, from about 10% to about 22%, from about 10% to about 21.5%, from about 10% to about 21%, from about 10% to about 20.5% or from about 10% to about 20% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[0070] In some embodiments, the proportion of the first carboxylic acid group-containing monomer is less than 30%, less than 29%, less than 28%, less than 27%, less than 26%, less than 25%, less than 24%, less than 23%, less than 22%, less than 21%, less than 20%, less than 19%, less than 18%, less than 17%, less than 16%, less than 15%, less than 14%, less than 13%, less than 12%, less than 11%, less than 10%, less than 9%, less than 8%, less than 7% or less than 6% by weight, based on the total weight of monomers added in the preparation of

the binder composition. In some embodiments, the proportion of the first carboxylic acid group-containing monomer is more than 5%, more than 6%, more than 7%, more than 8%, more than 9%, more than 10%, more than 11%, more than 12%, more than 13%, more than 14%, more than 15%, more than 16%, more than 17%, more than 18%, more than 19%, more than 20%, more than 21%, more than 22%, more than 23%, more than 24%, more than 25%, more than 26%, more than 27%, more than 28% or more than 29% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[0071] In some embodiments, the proportion of the second carboxylic acid group-containing monomer is from about 1% to about 7%, from about 1.2% to about 7%, from about 1.4% to about 7%, from about 1.6% to about 7%, from about 1.8% to about 7%, from about 2% to about 7%, from about 2.2% to about 7%, from about 2.4% to about 7%, from about 2.6% to about 7%, from about 2.8% to about 7%, from about 3% to about 7%, from about 3% to about 6.8%, from about 3% to about 6.6%, from about 3% to about 6.4%, from about 3.2% to about 6.4%, from about 3.4% to about 6.4%, from about 3.6% to about 6.4%, from about 3.8% to about 6.4%, from about 4% to about 6.4%, from about 4% to about 6.2%, from about 4% to about 6%, from about 3.5% to about 6%, from about 3% to about 6% or from about 3% to about 6.5% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[0072] In some embodiments, the proportion of the second carboxylic acid group-containing monomer is less than 7%, less than 6.8%, less than 6.6%, less than 6.4%, less than 6.2%, less than 6%, less than 5.8%, less than 5.6%, less than 5.4%, less than 5.2%, less than 5%, less than 4.8%, less than 4.6%, less than 4.4%, less than 4.2%, less than 4%, less than 3.8%, less than 3.6%, less than 3.4%, less than 3.2%, less than 3%, less than 2.8%, less than 2.6%, less than 2.4%, less than 2.2%, less than 2%, less than 1.8%, less than 1.6% or less than 1.4% by weight, based on the total weight of monomers added in the preparation of the binder composition. In some embodiments, the proportion of the second carboxylic acid group-containing monomer is more than 1%, more than 1.2%, more than 1.4%, more than 1.6%, more than 1.8%, more than 2%, more than 2.2%, more than 2.4%, more than 2.6%, more than 2.8%, more than 3%, more than 3.2%, more than 3.4%, more than 3.6%, more than 3.8%, more than 4%, more than 4.2%, more than 4.4%, more than 4.6%, more than 4.8%, more than 5%, more than 5.2%, more than 5.4%, more than 5.6%, more than 5.8%, more than 6%, more than 6.2%, more than 6.4% or more than 6.6% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[0073] In some embodiments, the weight ratio of the first carboxylic acid group-containing monomer to the second carboxylic acid group-containing monomer added in the preparation of the binder composition is from about 1 to about 15, from about 1 to about 14.5, from about 1 to about 14, from about 1 to about 13.5, from about 1 to about 13, from about 1 to

about 12.5, from about 1 to about 12, from about 1 to about 11.5, from about 1 to about 11, from about 1 to about 10.5, from about 1 to about 10, from about 1 to about 9.5, from about 1 to about 9, from about 1 to about 8.5, from about 1 to about 8, from about 1.5 to about 10, from about 2 to about 10, from about 2.5 to about 10, from about 3 to about 10, from about 3.5 to about 10, from about 4 to about 10, from about 4.5 to about 10, from about 5 to about 10 or from about 2 to about 8.

[0074] In some embodiments, the weight ratio of the first carboxylic acid group-containing monomer to the second carboxylic acid group-containing monomer added in the preparation of the binder composition is less than 15, less than 14, less than 13, less than 12, less than 11, less than 10, less than 9, less than 8, less than 7, less than 6, less than 5, less than 4, less than 3 or less than 2. In some embodiments, the weight ratio of the first carboxylic acid group-containing monomer to the second carboxylic acid group-containing monomer added in the preparation of the binder composition is more than 1, more than 2, more than 3, more than 4, more than 5, more than 6, more than 7, more than 8, more than 9, more than 10, more than 11, more than 12, more than 13 or more than 14.

[0075] In some embodiments, a third suspension is formed by adding an amide group-containing monomer into the second suspension in step **103**.

[0076] In some embodiments, an amide group-containing monomer solution is prepared by dissolving the amide group-containing monomer in water. In some embodiments, the third suspension is formed by adding the amide group-containing monomer solution into the second suspension in step **103**.

[0077] Structural unit (b) is derived from an amide group-containing monomer. Any monomer that has at least one amide group may be used as amide group-containing monomer without any specific limitations. In some embodiments, the amide group-containing monomer is acrylamide, methacrylamide, N-methyl methacrylamide, N-ethyl methacrylamide, N-n-propyl methacrylamide, N-isopropyl methacrylamide, isopropyl acrylamide, N-n-butyl methacrylamide, N-isobutyl methacrylamide, N,N-dimethyl acrylamide, N,N-dimethyl methacrylamide, N,N-diethyl acrylamide, N,N-diethyl methacrylamide, N-methylol methacrylamide, N-(methoxymethyl)methacrylamide, N-(ethoxymethyl)methacrylamide, N-(propoxymethyl)methacrylamide, N-(butoxymethyl)methacrylamide, N,N-dimethylaminopropyl methacrylamide, N,N-dimethylaminoethyl methacrylamide, N,N-dimethylol methacrylamide, diacetone methacrylamide, diacetone acrylamide, methacryloyl morpholine, N-hydroxyl methacrylamide, N-methoxymethyl acrylamide, N-methoxymethyl methacrylamide, N,N'-methylene-bis-acrylamide (MBA), N-hydroxymethyl acrylamide or a combination thereof.

[0078] In some embodiments, the proportion of the amide group-containing monomer is from about 5% to about 20%, from about 5% to about 15%, from about 5% to about 10%, from

about 6% to about 20%, from about 7% to about 20%, from about 8% to about 20%, from about 9% to about 20%, from about 10% to about 20%, from about 10% to about 19%, from about 10% to about 18%, from about 10% to about 17%, from about 10% to about 16%, from about 10% to about 15%, from about 8% to about 17%, from about 7% to about 13%, from about 12% to about 18% or from about 15% to about 20% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[0079] In some embodiments, the proportion of the amide group-containing monomer is less than 20%, less than 19%, less than 18%, less than 17%, less than 16%, less than 15%, less than 14%, less than 13%, less than 12%, less than 11%, less than 10%, less than 9%, less than 8%, less than 7% or less than 6% by weight, based on the total weight of monomers added in the preparation of the binder composition. In some embodiments, the proportion of the amide group-containing monomer is more than 5%, more than 6%, more than 7%, more than 8%, more than 9%, more than 10%, more than 11%, more than 12%, more than 13%, more than 14%, more than 15%, more than 16%, more than 17%, more than 18% or more than 19% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[0080] In some embodiments, the fourth suspension is formed by adding a nitrile group-containing monomer into the third suspension in step **104**.

[0081] Structural unit (c) is derived from a nitrile group-containing monomer. Any monomer that has at least one nitrile group may be used as nitrile group-containing monomer without any specific limitations. In some embodiments, the nitrile group-containing monomer include α,β-ethylenically unsaturated nitrile monomers. In some embodiments, the nitrile group-containing monomer is acrylonitrile, α-halogenoacrylonitrile, α-alkylacrylonitrile or a combination thereof. In some embodiments, the nitrile group-containing monomer is α-chloroacrylonitrile, α-bromoacrylonitrile, α-fluoroacrylonitrile, methacrylonitrile, α-ethylacrylonitrile, α-isopropylacrylonitrile, α-n-hexylacrylonitrile, α-methoxyacrylonitrile, 3-methoxyacrylonitrile, 3-ethoxyacrylonitrile, α-acetoxyacrylonitrile, α-phenylacrylonitrile, α-tolylacrylonitrile, α-(methoxyphenyl)acrylonitrile, α-(chlorophenyl)acrylonitrile, α-(cyanophenyl)acrylonitrile, vinylidene cyanide, or a combination thereof.

[0082] In some embodiments, the proportion of the nitrile group-containing monomer is from about 60% to about 75%, from about 60% to about 74.5%, from about 60% to about 74%, from about 60% to about 73.5%, from about 60% to about 73%, from about 60% to about 72.5%, from about 60% to about 72%, from about 60% to about 71.5%, from about 60% to about 71%, from about 60% to about 70.5%, from about 60% to about 70%, from about 60% to about 69.5%, from about 60% to about 69%, from about 60% to about 68.5%, from about 60% to about 68%, from about 60% to about 67.5%, from about 60% to about 67%, from about 60% to about 66.5%, from about 60% to about 66%, from about 60% to about 65.5%, from about

60% to about 65%, from about 65% to about 75%, from about 65% to about 70%, from about 63% to about 75% or from about 70% to about 75% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[0083] In some embodiments, the proportion of the nitrile group-containing monomer is more than 60%, more than 61%, more than 62%, more than 63%, more than 64%, more than 65%, more than 66%, more than 67%, more than 68%, more than 69%, more than 70%, more than 71%, more than 72%, more than 73% or more than 74% by weight, based on the total weight of monomers added in the preparation of the binder composition. In some embodiments, the proportion of the nitrile group-containing monomer is less than 75%, less than 74%, less than 73%, less than 72%, less than 71%, less than 70%, less than 69%, less than 68%, less than 67%, less than 66%, less than 65%, less than 64%, less than 63%, less than 62% or less than 61% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[0084] In certain embodiments, combinations of a carboxylic acid group-containing monomer, a nitrile group-containing monomer and an amide group-containing monomer may be added into the first suspension to form a second suspension without forming the third suspension and the fourth suspension. In other embodiments, a carboxylic acid group-containing monomer, a nitrile group-containing monomer, an amide group-containing monomer or combinations thereof are added sequentially into the first suspension to form a second suspension, a third suspension or the fourth suspension. Stirring or dispersion may be employed between the additions. This is advantageous as it allows better dispersion of materials. In the case where combinations of monomers are added sequentially, formation of the third suspension or the fourth suspension may be omitted.

[0085] In some embodiments, the copolymer is obtained via polymerization of a composition. In some embodiments, the composition comprises a carboxylic salt group-containing monomer, a carboxylic acid group-containing monomer, a nitrile group-containing monomer and an amide group-containing monomer. In some embodiments, formation of a carboxylic salt group-containing monomer is resulted from the neutralization of a carboxylic acid group-containing monomer by the neutralizing agent added in step **101**.

[0086] In some embodiments, the carboxylic salt group-containing monomer is acrylic acid salt, methacrylic acid salt, crotonic acid salt, 2-butyl crotonic acid salt, cinnamic acid salt, maleic acid salt, maleic anhydride salt, fumaric acid salt, itaconic acid salt, itaconic anhydride salt, tetraconic acid salt or a combination thereof. In certain embodiments, the carboxylic salt group-containing monomer is 2-ethylacrylic acid salt, isocrotonic acid salt, cis-2-pentenoic acid salt, trans-2-pentenoic acid salt, angelic acid salt, tiglic acid salt, 3,3-dimethyl acrylic acid salt, 3-propyl acrylic acid salt, trans-2-methyl-3-ethyl acrylic acid salt, cis-2-methyl-3-ethyl acrylic

acid salt, 3-isopropyl acrylic acid salt, trans-3-methyl-3-ethyl acrylic acid salt, cis-3-methyl-3-ethyl acrylic acid salt, 2-isopropyl acrylic acid salt, trimethyl acrylic acid salt, 2-methyl-3,3-diethyl acrylic acid salt, 3-butyl acrylic acid salt, 2-butyl acrylic acid salt, 2-pentyl acrylic acid salt, 2-methyl-2-hexenoic acid salt, trans-3-methyl-2-hexenoic acid salt, 3-methyl-3-propyl acrylic acid salt, 2-ethyl-3-propyl acrylic acid salt, 2,3-diethyl acrylic acid salt, 3,3-diethyl acrylic acid salt, 3-methyl-3-hexyl acrylic acid salt, 3-methyl-3-tert-butyl acrylic acid salt, 2-methyl-3-pentyl acrylic acid salt, 3-methyl-3-pentyl acrylic acid salt, 4-methyl-2-hexenoic acid salt, 4-ethyl-2-hexenoic acid salt, 3-methyl-2-ethyl-2-hexenoic acid salt, 3-tert-butyl acrylic acid salt, 2,3-dimethyl-3-ethyl acrylic acid salt, 3,3-dimethyl-2-ethyl acrylic acid salt, 3-methyl-3-isopropyl acrylic acid salt, 2-methyl-3-isopropyl acrylic acid salt, trans-2-octenoic acid salt, cis-2-octenoic acid salt, trans-2-decenoic acid salt, α-acetoxyacrylic acid salt, β-trans-aryloxyacrylic acid salt, α-chloro-β-E-methoxyacrylic acid salt or a combination thereof. In some embodiments, the carboxylic salt group-containing monomer is methyl maleic acid salt, dimethyl maleic acid salt, phenyl maleic acid salt, bromo maleic acid salt, chloromaleic acid salt, dichloromaleic acid salt, fluoromaleic acid salt, difluoro maleic acid salt or a combination thereof.

[0087] In some embodiments, the carboxylic salt group-containing monomer is an alkali metal carboxylic salt group-containing monomer. Examples of an alkali metal forming the alkali metal carboxylic salt include lithium, sodium and potassium. In some embodiments, the carboxylic salt group-containing monomer is an ammonium carboxylic salt group-containing monomer.

[0088] In some embodiments, the molar ratio of the carboxylic acid group-containing monomer to the carboxylic salt group-containing monomer in the composition is from about 0 to about 1.5, from about 0 to about 1.45, from about 0 to about 1.4, from about 0 to about 1.35, from about 0 to about 1.3, from about 0 to about 1.25, from about 0 to about 1.2, from about 0 to about 1.15, from about 0 to about 1.1, from about 0 to about 1.05, from about 0 to about 1, from about 0 to about 0.95, from about 0 to about 0.9, from about 0 to about 0.85, from about 0 to about 0.8, from about 0 to about 0.75, from about 0 to about 0.7, from about 0 to about 0.65, from about 0 to about 0.6, from about 0 to about 0.55, from about 0 to about 0.5, from about 0 to about 0.45, from about 0 to about 0.4, from about 0.05 to about 0.5, from about 0.1 to about 0.7 or from about 0.1 to about 1.

[0089] In some embodiments, the molar ratio of the carboxylic acid group-containing monomer to the carboxylic salt group-containing monomer in the composition is less than 1.5, less than 1.4, less than 1.3, less than 1.2, less than 1.1, less than 1, less than 0.9, less than 0.8, less than 0.7, less than 0.6, less than 0.5, less than 0.4, less than 0.3 or less than 0.2. In some embodiments, the molar ratio of the carboxylic acid group-containing monomer to the carboxylic salt group-containing monomer in the composition is more than 0, more than 0.1, more than 0.2, more than 0.3, more than 0.4, more than 0.5, more than 0.6, more than 0.7, more

than 0.8, more than 0.9, more than 1, more than 1.1, more than 1.2 or more than 1.3.

[0090] In some embodiments, the proportion of the carboxylic acid group-containing monomer is from about 0% to about 15%, from about 0% to about 14.5%, from about 0% to about 14%, from about 0% to about 13.5%, from about 0% to about 13%, from about 0% to about 12.5%, from about 0% to about 12%, from about 0% to about 11.5%, from about 0% to about 11%, from about 0% to about 10.5%, from about 0% to about 10%, from about 0% to about 9.5%, from about 0% to about 9%, from about 0% to about 8.5%, from about 0% to about 8%, from about 0% to about 7.5%, from about 0% to about 7%, from about 0% to about 6.5%, from about 0% to about 6%, from about 0% to about 5.5%, from about 0% to about 5%, from about 0.5% to about 10%, from about 1% to about 10% or from about 1% to about 8% by mole, based on the total number of moles of monomers in the composition.

[0091] In some embodiments, the proportion of the carboxylic acid group-containing monomer is less than 15%, less than 14%, less than 13%, less than 12%, less than 11%, less than 10%, less than 9%, less than 8%, less than 7%, less than 6%, less than 5%, less than 4%, less than 3% or less than 2% by mole, based on the total number of moles of monomers in the composition. In some embodiments, the proportion of the carboxylic acid group-containing monomer is more than 0%, more than 1%, more than 2%, more than 3%, more than 4%, more than 5%, more than 6%, more than 7%, more than 8%, more than 9%, more than 10%, more than 11%, more than 12%, more than 13% or more than 14% by mole, based on the total number of moles of monomers in the composition.

[0092] In some embodiments, proportion of the carboxylic salt group-containing monomer is from about 5% to about 16%, from about 5.5% to about 16%, from about 6% to about 16%, from about 6.5% to about 16%, from about 7% to about 16%, from about 7.5% to about 16%, from about 8% to about 16%, from about 8.5% to about 16%, from about 9% to about 16%, from about 9.5% to about 16%, from about 10% to about 16%, from about 10% to about 15.5%, from about 10% to about 15%, from about 10.5% to about 15%, from about 11% to about 15% or from about 8% to about 15% by mole, based on the total number of moles of monomers in the composition.

[0093] In some embodiments, proportion of the carboxylic salt group-containing monomer is less than 16%, less than 15%, less than 14%, less than 13%, less than 12%, less than 11%, less than 10%, less than 9%, less than 8%, less than 7% or less than 6% by mole, based on the total number of moles of monomers in the composition. In some embodiments, proportion of the carboxylic salt group-containing monomer is more than 5%, more than 6%, more than 7%, more than 8%, more than 9%, more than 10%, more than 11%, more than 12%, more than 13%, more than 14% or more than 15% by mole, based on the total number of moles of monomers in the composition.

[0094] In some embodiments, the proportion of the nitrile group-containing monomer is from about 65% to about 80%, from about 65.5% to about 80%, from about 66% to about 80%, from about 66.5% to about 80%, from about 67% to about 80%, from about 67.5% to about 80%, from about 68% to about 80%, from about 68.5% to about 80%, from about 69% to about 80%, from about 69.5% to about 80%, from about 70% to about 80%, from about 70.5% to about 80%, from about 71% to about 80%, from about 71.5% to about 80%, from about 72% to about 80%, from about 65% to about 78%, from about 65% to about 75%, from about 68% to about 76%, from about 70% to about 78% or from about 70% to about 75% by mole, based on the total number of moles of monomers in the composition.

[0095] In some embodiments, the proportion of the nitrile group-containing monomer is less than 80%, less than 79%, less than 78%, less than 77%, less than 76%, less than 75%, less than 74%, less than 73%, less than 72%, less than 71%, less than 70%, less than 69%, less than 68% or less than 67% by mole, based on the total number of moles of monomers in the composition. In some embodiments, the proportion of the nitrile group-containing monomer is more than 65%, more than 66%, more than 67%, more than 68%, more than 69%, more than 70%, more than 71%, more than 72%, more than 73%, more than 74%, more than 75%, more than 76%, more than 77% or more than 78% by mole, based on the total number of moles of monomers in the composition.

[0096] In some embodiments, the proportion of the amide group-containing monomer is from about 5% to about 20%, from about 5% to about 15%, from about 5% to about 10%, from about 6% to about 20%, from about 7% to about 20%, from about 8% to about 20%, from about 9% to about 20%, from about 10% to about 20%, from about 10% to about 19%, from about 10% to about 18%, from about 10% to about 17%, from about 10% to about 16%, from about 10% to about 15%, from about 8% to about 17%, from about 7% to about 13%, from about 12% to about 18% or from about 15% to about 20% by mole, based on the total number of moles of monomers in the composition.

[0097] In some embodiments, the proportion of the amide group-containing monomer is less than 20%, less than 19%, less than 18%, less than 17%, less than 16%, less than 15%, less than 14%, less than 13%, less than 12%, less than 11%, less than 10%, less than 9%, less than 8%, less than 7% or less than 6% by mole, based on the total number of moles of monomers in the composition. In some embodiments, the proportion of the amide group-containing monomer is more than 5%, more than 6%, more than 7%, more than 8%, more than 9%, more than 10%, more than 11%, more than 12%, more than 13%, more than 14%, more than 15%, more than 16%, more than 17%, more than 18% or more than 19% by mole, based on the total number of moles of monomers in the composition.

[0098] In some embodiments, each of the second suspension, the third suspension and

the fourth suspension is independently stirred at a speed of from about 20 rpm to about 300 rpm, from about 20 rpm to about 280 rpm, from about 20 rpm to about 260 rpm, from about 20 rpm to about 240 rpm, from about 20 rpm to about 220 rpm, from about 20 rpm to about 200 rpm, from about 20 rpm to about 180 rpm, from about 20 rpm to about 160 rpm, from about 40 rpm to about 160 rpm, from about 60 rpm to about 160 rpm, from about 60 rpm to about 140 rpm, from about 80 rpm to about 140 rpm, from about 80 rpm to about 120 rpm, from about 50 rpm to about 150 rpm or from about 50 rpm to about 200 rpm.

[0099] In some embodiments, each of the second suspension, the third suspension and the fourth suspension is independently stirred at a speed of less than 300 rpm, less than 280 rpm, less than 260 rpm, less than 240 rpm, less than 220 rpm, less than 200 rpm, less than 180 rpm, less than 160 rpm, less than 140 rpm, less than 120 rpm, less than 100 rpm, less than 80 rpm, less than 60 rpm or less than 40 rpm. In some embodiments, each of the second suspension, the third suspension and the fourth suspension is independently stirred at a speed of more than 20 rpm, more than 40 rpm, more than 60 rpm, more than 80 rpm, more than 100 rpm, more than 120 rpm, more than 140 rpm, more than 160 rpm, more than 180 rpm, more than 200 rpm, more than 220 rpm, more than 240 rpm, more than 260 rpm or more than 280 rpm.

[00100] In some embodiments, each of the second suspension, the third suspension and the fourth suspension is independently stirred for a time period of from about 30 minutes to about 120 minutes, from about 30 minutes to about 105 minutes, from about 30 minutes to about 90 minutes, from about 45 minutes to about 90 minutes, from about 45 minutes to about 75 minutes, from about 50 minutes to about 70 minutes or from about 40 minutes to about 80 minutes. In some embodiments, each of the second suspension, the third suspension and the fourth suspension is independently stirred for a time period of less than 120 minutes, less than 110 minutes, less than 100 minutes, less than 90 minutes, less than 80 minutes, less than 70 minutes, less than 60 minutes, less than 50 minutes or less than 40 minutes. In some embodiments, each of the second suspension, the third suspension and the fourth suspension is independently stirred for a time period of more than 30 minutes, more than 40 minutes, more than 50 minutes, more than 60 minutes, more than 70 minutes, more than 80 minutes, more than 90 minutes, more than 100 minutes or more than 110 minutes.

[00101] In some embodiments, an initiator solution is prepared by dissolving the initiator in water. In some embodiments, a fifth suspension is formed by adding the initiator solution into the fourth suspension dropwise in step **105**.

[00102] In some embodiments, the temperature of the fourth suspension is elevated to from about 30 ºC to about 70 ºC, from about 32 ºC to about 70 ºC, from about 34 ºC to about 70 ºC, from about 36 ºC to about 70 ºC, from about 38 ºC to about 70 ºC, from about 40 ºC to about 70 ºC, from about 42 ºC to about 70 ºC, from about 44 ºC to about 70 ºC, from about 46 ºC to

about 70 ºC, from about 48 ºC to about 70 ºC or from about 50 ºC to about 70 ºC, before the addition of the initiator solution into the fourth suspension to form the fifth suspension.

[00103] In some embodiments, the temperature of the fourth suspension is elevated to less than 70 ºC, less than 68 ºC, less than 66 ºC, less than 64 ºC, less than 62 ºC, less than 60 ºC, less than 58 ºC, less than 56 ºC, less than 54 ºC, less than 52 ºC, less than 50 ºC, less than 48 ºC, less than 46 ºC, less than 44 ºC, less than 42 ºC, less than 40 ºC, less than 38 ºC, less than 36 ºC or less than 34 ºC, before the addition of the initiator solution into the fourth suspension to form the fifth suspension. In some embodiments, the temperature of the fourth suspension is elevated to more than 30 ºC, more than 32 ºC, more than 34 ºC, more than 36 ºC, more than 38 ºC, more than 40 ºC, more than 42 ºC, more than 44 ºC, more than 46 ºC, more than 48 ºC, more than 50 ºC, more than 52 ºC, more than 54 ºC, more than 56 ºC, more than 58 ºC, more than 60 ºC, more than 62 ºC, more than 64 ºC or more than 66 ºC, before the addition of the initiator solution into the fourth suspension to form the fifth suspension.

[00104] In some embodiments, the fourth suspension is stirred at a speed of from about 50 rpm to about 500 rpm, from about 50 rpm to about 450 rpm, from about 50 rpm to about 400 rpm, from about 50 rpm to about 350 rpm, from about 50 rpm to about 300 rpm, from about 50 rpm to about 280 rpm, from about 50 rpm to about 260 rpm, from about 50 rpm to about 240 rpm, from about 50 rpm to about 220 rpm, from about 50 rpm to about 200 rpm, from about 50 rpm to about 180 rpm, from about 50 rpm to about 160 rpm, from about 50 rpm to about 140 rpm, from about 50 rpm to about 120 rpm or from about 50 rpm to about 100 rpm, before the addition of the initiator solution into the fourth suspension to form the fifth suspension.

[00105] In some embodiments, the fourth suspension is stirred at speed of less than 500 rpm, less than 450 rpm, less than 400 rpm, less than 350 rpm, less than 300 rpm, less than 250 rpm, less than 200 rpm, less than 150 rpm or less than 100 rpm, before the addition of the initiator solution into the fourth suspension to form the fifth suspension. In some embodiments, the fourth suspension is stirred at a speed of more than 50 rpm, more than 100 rpm, more than 150 rpm, more than 200 rpm, more than 250 rpm, more than 300 rpm, more than 350 rpm, more than 400 rpm or more than 450 rpm, before the addition of the initiator solution into the fourth suspension to form the fifth suspension.

[00106] In some embodiments, the fourth suspension is stirred for a time period of from about 30 minutes to about 120 minutes, from about 30 minutes to about 105 minutes, from about 30 minutes to about 90 minutes, from about 45 minutes to about 90 minutes, from about 45 minutes to about 75 minutes, from about 50 minutes to about 70 minutes or from about 40 minutes to about 80 minutes, before the addition of the initiator solution into the fourth suspension to form the fifth suspension. In some embodiments, the fourth suspension is stirred for a time period of less than 120 minutes, less than 110 minutes, less than 100 minutes, less

than 90 minutes, less than 80 minutes, less than 70 minutes, less than 60 minutes, less than 50 minutes or less than 40 minutes, before the addition of the initiator solution into the fourth suspension to form the fifth suspension. In some embodiments, the fourth suspension is stirred for a time period of more than 30 minutes, more than 40 minutes, more than 50 minutes, more than 60 minutes, more than 70 minutes, more than 80 minutes, more than 90 minutes, more than 100 minutes or more than 110 minutes, before the addition of the initiator solution into the fourth suspension to form the fifth suspension.

[00107] Polymerization occurred in the present invention follows the radical mechanism, wherein an initiator acts to generate free radicals, which in turn lead to polymer chains propagation. The free radicals used herein can be produced using thermal decomposition or redox reactions. The free radical initiator(s) disclosed herein is/are water-soluble.

[00108] The water-soluble free radical initiator decomposes thermally in the aqueous phase to give radicals which can initiate the polymerization. In some embodiments, the water-soluble initiator may be selected from the group consisting of persulfate-based initiators such as ammonium persulfate, sodium persulfate, potassium persulfate and the like; azo-based initiators such as azobis (isobutyl-amidine hydrochloride) (AIBA), 2,2'-azobis(2-methylpropionamidine) dihydrochloride, 2,2-azobis(2-amidinopropane) dihydrochloride (AAPH), 2,2'-azobis[2-(2-imidazolin-2-yl)propane]dihydrochloride, bis[2-(4’-sulfophenyl)alkyl]-2,2’-azodiisobutyrate ammonium salts, 2,2′-azobis(N-2′-methylpropanoyl-2-amino-alkyl-1-sulfonate and the like; peroxide-based initiators such as hydrogen peroxide, t-butyl hydroperoxide, succinic acid peroxide and the like and combinations thereof.

[00109] In some embodiments, the water-soluble free radical initiator can be used together with a reducing agent to establish a redox initiator system. This allows generation of free radicals through an oxidation-reduction reaction at relatively low temperatures and promotes an enhancement in polymerization rate.

[00110] In some embodiments, the reducing agent can be selected from the group consisting of sodium bisulfite, sodium metabisulfite, sodium sulfite, sodium thiosulfate, thiourea dioxide, ferrous sulfate, ferrous chloride, ascorbic acid, citric acid, tartaric acid, erythorbic acid, glucose, metal salt of formaldehyde sulfoxylate, Bruggolite FF6M, and combinations thereof.

[00111] In some embodiments, the proportion of water-soluble free radical initiator is from about 0.05% to about 0.4%, from about 0.07% to about 0.4%, from about 0.1% to about 0.4%, from about 0.1% to about 0.39%, from about 0.1% to about 0.38%, from about 0.1% to about 0.37%, from about 0.1% to about 0.36%, from about 0.1% to about 0.35%, from about 0.1% to about 0.34%, from about 0.1% to about 0.33%, from about 0.1% to about 0.32%, from about 0.1% to about 0.31%, from about 0.1% to about 0.3%, from about 0.1% to about 0.29%, from about 0.1% to about 0.28%, from about 0.1% to about 0.27% or from about 0.1% to about

0.26% by weight, based on the total weight of monomers added in the preparation of the binder composition. When the proportion of the water-soluble initiator out of the total weight of monomers added in the preparation of the binder composition is within the above range, a higher monomer conversion rate could be achieved and the binder composition could exhibit a better overall binding performance.

[00112] In some embodiments, the proportion of water-soluble initiator is less than 0.4%, less than 0.38%, less than 0.36%, less than 0.34%, less than 0.32%, less than 0.3%, less than 0.28%, less than 0.26%, less than 0.24%, less than 0.22%, less than 0.2%, less than 0.18%, less than 0.16%, less than 0.14%, less than 0.12%, less than 0.1% or less than 0.08% by weight, based on the total weight of monomers added in the preparation of the binder composition. In some embodiments, the proportion of water-soluble initiator is more than 0.05%, more than 0.07%, more than 0.1%, more than 0.12%, more than 0.14%, more than 0.16%, more than 0.18%, more than 0.2%, more than 0.22%, more than 0.24%, more than 0.26%, more than 0.28%, more than 0.3%, more than 0.32%, more than 0.34%, more than 0.36% or more than 0.38% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[00113] In some embodiments, the proportion of reducing agent is from about 0.01% to about 0.2%, from about 0.02% to about 0.2%, from about 0.03% to about 0.2%, from about 0.04% to about 0.2%, from about 0.05% to about 0.2%, from about 0.06% to about 0.2%, from about 0.07% to about 0.2%, from about 0.08% to about 0.2%, from about 0.09% to about 0.2% or from about 0.1% to about 0.2% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[00114] In some embodiments, the proportion of reducing agent is less than 0.2%, less than 0.19%, less than 0.18%, less than 0.17%, less than 0.16%, less than 0.15%, less than 0.14%, less than 0.13%, less than 0.12%, less than 0.11%, less than 0.1%, less than 0.09%, less than 0.08%, less than 0.07%, less than 0.06%, less than 0.05% or less than 0.04% by weight, based on the total weight of monomers added in the preparation of the binder composition. In some embodiments, the proportion of reducing agent is more than 0.01%, more than 0.02%, more than 0.03%, more than 0.04%, more than 0.05%, more than 0.06%, more than 0.07%, more than 0.08%, more than 0.09%, more than 0.1%, more than 0.11%, more than 0.12%, more than 0.13%, more than 0.14%, more than 0.15% or more than 0.16% by weight, based on the total weight of monomers added in the preparation of the binder composition.

[00115] In some embodiments, when a redox initiator system is selected as the initiator, the molar ratio of the water-soluble free radical initiator to the reducing agent is from about 0.2 to about 10, from about 0.2 to about 9, from about 0.2 to about 8, from about 0.2 to about 7, from about 0.2 to about 6, from about 0.2 to about 5, from about 0.3 to about 5, from about 0.4

to about 5, from about 0.5 to about 5, from about 0.6 to about 5, from about 0.7 to about 5, from about 0.8 to about 5, from about 0.9 to about 5, from about 1 to about 5, from about 0.5 to about 4.5, from about 0.5 to about 4, from about 0.6 to about 3.5, from about 0.6 to about 3, from about 0.8 to about 3 or from about 0.2 to about 1.

[00116] In some embodiments, when a redox initiator system is selected as the initiator, the molar ratio of the water-soluble free radical initiator to the reducing agent is less than 10, less than 9, less than 8, less than 7, less than 6, less than 5, less than 4.8, less than 4.6, less than 4.4, less than 4.2, less than 4, less than 3.8, less than 3.6, less than 3.4, less than 3.2, less than 3, less than 2.8, less than 2.6, less than 2.4, less than 2.2, less than 2, less than 1.8, less than 1.6, less than 1.4, less than 1.2, less than 1, less than 0.8, less than 0.6 or less than 0.4. In some embodiments, when a redox initiator system is selected as the initiator, the molar ratio of the water-soluble free radical initiator to the reducing agent is more than 0.2, more than 0.4, more than 0.6, more than 0.8, more than 1, more than 1.2, more than 1.4, more than 1.6, more than 1.8, more than 2, more than 2.2, more than 2.4, more than 2.6, more than 2.8, more than 3, more than 3.2, more than 3.4, more than 3.6, more than 3.8, more than 4, more than 4.2, more than 4.4, more than 4.6, more than 4.8, more than 5, more than 6, more than 7, more than 8 or more than 9.

[00117] In some embodiments, a reducing agent solution is prepared by dissolving the reducing agent in water. In some embodiments, when a redox initiator system is selected as the initiator, the reducing agent is added to the fourth suspension before the addition of the initiator solution to form the fifth suspension.

[00118] In some embodiments, the initiator solution is added dropwise into the fourth suspension for a time period of from about 2 hours to about 5 hours, from about 2 hours to about 4.75 hours, from about 2 hours to about 4.5 hours, from about 2 hours to about 4.25 hours, from about 2 hours to about 4 hours, from about 2 hours to about 3.75 hours, from about 2 hours to about 3.5 hours, from about 2.25 hours to about 3.5 hours or from about 2.5 hours to about 3.5 hours. In some embodiments, the initiator solution is added dropwise into the fourth suspension for a time period of less than 5 hours, less than 4.75 hours, less than 4.5 hours, less than 4.25 hours, less than 4 hours, less than 3.75 hours, less than 3.5 hours, less than 3.25 hours, less than 3 hours, less than 2.75 hours or less than 2.5 hours. In some embodiments, the initiator solution is added dropwise into the fourth suspension for a time period of more than 2 hours, more than 2.25 hours, more than 2.5 hours, more than 2.75 hours, more than 3 hours, more than 3.25 hours, more than 3.5 hours, more than 3.75 hours, more than 4 hours, more than 4.25 hours or more than 4.5 hours.

[00119] The polymerization temperature is dependent on the type of initiator applied. In some embodiments, the reaction temperature of the polymerization is from about 50 °C to about

90 °C, from about 50 °C to about 85 °C, from about 50 °C to about 80 °C, from about 50 °C to about 75 °C, from about 50 °C to about 70 °C, from about 55 °C to about 75 °C, from about 55 °C to about 80 °C, from about 55 °C to about 85 °C, from about 60 °C to about 80 °C, from about 60 °C to about 75 °C, from about 60 °C to about 70 °C or from about 55 °C to about 70 °C. When the reaction temperature of the polymerization is within the above range, a higher reaction stability could be achieved and the binder composition could exhibit a better overall binding performance.

[00120] In some embodiments, the reaction temperature of the polymerization is less than 90 °C, less than 88 °C, less than 86 °C, less than 84 °C, less than 82 °C, less than 80 °C, less than 78 °C, less than 76 °C, less than 74 °C, less than 72 °C, less than 70 °C, less than 68 °C, less than 66 °C, less than 64 °C, less than 62 °C, less than 60 °C, less than 58 °C, less than 56 °C or less than 54 °C. In some embodiments, the reaction temperature of the polymerization is more than 50 °C, more than 52 °C, more than 54 °C, more than 56 °C, more than 58 °C, more than 60 °C, more than 62 °C, more than 64 °C, more than 66 °C, more than 68 °C, more than 70 °C, more than 72 °C, more than 74 °C, more than 76 °C, more than 78 °C, more than 80 °C, more than 82 °C, more than 84 °C or more than 86 °C.

[00121] In some embodiments, the stirring speed of the mixer during polymerization is from about 100 rpm to about 1000 rpm, from about 100 rpm to about 950 rpm, from about 100 rpm to about 900 rpm, from about 100 rpm to about 850 rpm, from about 100 rpm to about 800 rpm, from about 100 rpm to about 750 rpm, from about 100 rpm to about 700 rpm, from about 100 rpm to about 650 rpm, from about 100 rpm to about 600 rpm, from about 100 rpm to about 550 rpm, from about 100 rpm to about 500 rpm, from about 150 rpm to about 500 rpm, from about 200 rpm to about 500 rpm, from about 250 rpm to about 500 rpm, from about 250 rpm to about 450 rpm, from about 300 rpm to about 450 rpm or from about 300 rpm to about 400 rpm. In some embodiments, the stirring speed of the mixer during polymerization is less than 1000 rpm, less than 950 rpm, less than 900 rpm, less than 850 rpm, less than 800 rpm, less than 750 rpm, less than 700 rpm, less than 650 rpm, less than 600 rpm, less than 550 rpm, less than 500 rpm, less than 450 rpm, less than 400 rpm, less than 350 rpm, less than 300 rpm, less than 250 rpm, less than 200 rpm or less than 150 rpm. In some embodiments, the stirring speed of the mixer during polymerization is more than 100 rpm, more than 150 rpm, more than 200 rpm, more than 250 rpm, more than 300 rpm, more than 350 rpm, more than 400 rpm, more than 450 rpm, more than 500 rpm, more than 550 rpm, more than 600 rpm, more than 650 rpm, more than 700 rpm, more than 750 rpm, more than 800 rpm, more than 850 rpm, more than 900 rpm or more than 950 rpm.

[00122] In some embodiments, the reaction time of the polymerization is from about 20 hours to about 24 hours, from about 20.25 hours to about 24 hours, from about 20.5 hours to about 24 hours, from about 20.75 hours to about 24 hours, from about 21 hours to about 24

hours, from about 21.25 hours to about 24 hours, from about 21.5 hours to about 24 hours, from about 21.75 hours to about 24 hours, from about 22 hours to about 24 hours, from about 20 hours to about 23.75 hours, from about 20 hours to about 23.5 hours, from about 20 hours to about 23.25 hours, from about 20 hours to about 23 hours, from about 20 hours to about 22.75 hours, from about 20 hours to about 22.5 hours, from about 20 hours to about 22.25 hours, from about 20 hours to about 22 hours or from about 22 hours to about 23 hours.

[00123] In some embodiments, the reaction time of the polymerization is less than 24 hours, less than 23.75 hours, less than 23.5 hours, less than 23.25 hours, less than 23 hours, less than 22.75 hours, less than 22.5 hours, less than 22.25 hours, less than 22 hours, less than 21.75 hours, less than 21.5 hours, less than 21.25 hours, less than 21 hours, less than 20.75 hours, less than 20.5 hours or less than 20.25 hours. In some embodiments, the reaction time of the polymerization is more than 20 hours, more than 20.25 hours, more than 20.5 hours, more than 20.75 hours, more than 21 hours, more than 21.25 hours, more than 21.5 hours, more than 21.75 hours, more than 22 hours, more than 22.25 hours, more than 22.5 hours, more than 22.75 hours, more than 23 hours, more than 23.25 hours, more than 23.5 hours or more than 23.75 hours.

[00124] In some embodiments, the fifth suspension is stirred for a time period of from about 22 hours to about 30 hours, from about 22 hours to about 29.5 hours, from about 22 hours to about 29 hours, from about 22 hours to about 28.5 hours, from about 22 hours to about 28 hours, from about 22.5 hours to about 28 hours, from about 23 hours to about 28 hours, from about 23.5 hours to about 28 hours, from about 24 hours to about 28 hours, from about 24 hours to about 27.5 hours, from about 24 hours to about 27 hours, from about 24.5 hours to about 27 hours, from about 25 hours to about 27 hours, from about 24 hours to about 26 hours or from about 26 hours to about 28 hours during the addition of the initiator solution and as polymerization takes place.

[00125] In some embodiments, the fifth suspension is stirred for a time period of less than 30 hours, less than 29.5 hours, less than 29 hours, less than 28.5 hours, less than 28 hours, less than 27.5 hours, less than 27 hours, less than 26.5 hours, less than 26 hours, less than 25.5 hours, less than 25 hours, less than 24.5 hours, less than 24 hours, less than 23.5 hours, less than 23 hours or less than 22.5 hours during the addition of the initiator solution and as polymerization takes place. In some embodiments, the fifth suspension is stirred for a time period of more than 22 hours, more than 22.5 hours, more than 23 hours, more than 23.5 hours, more than 24 hours, more than 24.5 hours, more than 25 hours, more than 25.5 hours, more than 26 hours, more than 26.5 hours, more than 27 hours, more than 27.5 hours, more than 28 hours, more than 28.5 hours, more than 29 hours or more than 29.5 hours during the addition of the initiator solution and as polymerization takes place.

[00126] In some embodiments, a neutralizing solution is prepared by dissolving the

neutralizing agent in water. In some embodiments, a sixth suspension is formed by adding the neutralizing solution into the fifth suspension in step **106**. The above-mentioned neutralizing agent in step **101** may be selected. In some embodiments, the neutralizing agent applied in step **101** may correspond to the neutralizing agent used in step **106**. In some embodiments, the neutralizing agents applied in steps **101** and **106** may not be the same.

[00127] In some embodiments, the temperature of the fifth suspension is lowered to from about 40 ºC to about 50 ºC, from about 40 ºC to about 49 ºC, from about 40 ºC to about 48 ºC, from about 40 ºC to about 47 ºC, from about 40 ºC to about 46 ºC, from about 40 ºC to about 45 ºC, from about 41 ºC to about 50 ºC, from about 42 ºC to about 50 ºC, from about 43 ºC to about 50 ºC, from about 44 ºC to about 50 ºC, from about 45 ºC to about 50 ºC or from about 42 ºC to about 48 ºC, prior to adding a neutralizing solution to form a sixth suspension. In some embodiments, the temperature of the fifth suspension is lowered to less than 50 ºC, less than 49 ºC, less than 48 ºC, less than 47 ºC, less than 46 ºC, less than 45 ºC, less than 44 ºC, less than 43 ºC, less than 42 ºC or less than 41 ºC, prior to adding a neutralizing solution to form a sixth suspension. In some embodiments, the temperature of the fifth suspension is lowered to more than 40 ºC, more than 41 ºC, more than 42 ºC, more than 43 ºC, more than 44 ºC, more than 45 ºC, more than 46 ºC, more than 47 ºC, more than 48 ºC or more than 49 ºC, prior to adding a neutralizing solution to form a sixth suspension.

[00128] In some embodiments, the total proportion of the neutralizing agent is from about 10% to about 40%, from about 10% to about 38%, from about 10% to about 36%, from about 10% to about 34%, from about 10% to about 32%, from about 10% to about 30%, from about 10% to about 28%, from about 10% to about 26%, from about 10% to about 25.5%, from about 10% to about 25%, from about 10% to about 24.5%, from about 10% to about 24%, from about 10% to about 23.5%, from about 10% to about 23%, from about 10% to about 22.5%, from about 10% to about 22%, from about 10% to about 21.5%, from about 10% to about 21%, from about 10% to about 20.5%, from about 10% to about 20%, from about 10% to about 19.5%, from about 10% to about 19%, from about 10% to about 18.5%, from about 10% to about 18%, from about 10% to about 17.5%, from about 10% to about 17%, from about 10% to about 16.5%, from about 10% to about 16%, from about 10% to about 15.5%, from about 10% to about 15%, from about 10.5% to about 19%, from about 11% to about 19% or from about 11% to about 15% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00129] In some embodiments, the total proportion of the neutralizing agent is less than 40%, less than 38%, less than 36%, less than 34%, less than 32%, less than 30%, less than 29%, less than 28%, less than 27%, less than 26%, less than 25%, less than 24%, less than 23%, less than 22%, less than 21%, less than 20%, less than 19%, less than 18%, less than 17%, less than 16%, less than 15%, less than 14%, less than 13%, less than 12% or less than 11% by mole,

based on the total number of moles of monomeric units in the copolymer in the binder composition. In some embodiments, the total proportion of the neutralizing agent is more than 10%, more than 11%, more than 12%, more than 13%, more than 14%, more than 15%, more than 16%, more than 17%, more than 18%, more than 19%, more than 20%, more than 21%, more than 22%, more than 23%, more than 24%, more than 25%, more than 26%, more than 28%, more than 30%, more than 32%, more than 34%, more than 36% or more than 38% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00130] In some embodiments, the sixth suspension is stirred for a time period of from about 1 hour to about 4 hours, from about 1.25 hours to about 4 hours, from about 1.5 hours to about 4 hours, from about 1.75 hours to about 4 hours, from about 2 hours to about 4 hours, from about 2.25 hours to about 4 hours, from about 2.5 hours to about 4 hours, from about 2.5 hours to about 3.75 hours, from about 2.5 hours to about 3.5 hours, from about 2.75 hours to about 3.5 hours, from about 2.75 hours to about 3.25 hours, from about 2.5 hours to about 3 hours, from about 2 hours to about 3 hours, from about 3 hours to about 3.5 hours or from about 3 hours to about 4 hours. In some embodiments, the sixth suspension is stirred for a time period of less than 4 hours, less than 3.75 hours, less than 3.5 hours, less than 3.25 hours, less than 3 hours, less than 2.75 hours, less than 2.5 hours, less than 2.25 hours, less than 2 hours, less than 1.75 hours, less than 1.5 hours or less than 1.25 hours. In some embodiments, the sixth suspension is stirred for a time period of more than 1 hour, more than 1.25 hours, more than 1.5 hours, more than 1.75 hours, more than 2 hours, more than 2.25 hours, more than 2.5 hours, more than 2.75 hours, more than 3 hours, more than 3.25 hours, more than 3.5 hours or more than 3.75 hours.

[00131] In some embodiments, the temperature of the sixth suspension is lowered to from 20 ºC to about 35 ºC, from about 21 ºC to about 35 ºC, from about 22 ºC to about 35 ºC, from about 23 ºC to about 35 ºC, from about 24 ºC to about 35 ºC, from about 25 ºC to about 35 ºC, from about 26 ºC to about 35 ºC, from about 27 ºC to about 35 ºC, from about 28 ºC to about 35 ºC, from about 29 ºC to about 35 ºC, from about 30 ºC to about 35 ºC, from about 20 ºC to about 34 ºC, from about 20 ºC to about 33 ºC, from about 20 ºC to about 32 ºC, from about 20 ºC to about 31 ºC, from about 20 ºC to about 30 ºC, from about 20 ºC to about 29 ºC, from about 20 ºC to about 28 ºC, from about 20 ºC to about 27 ºC or from about 25 ºC to about 30 ºC. In some embodiments, the temperature of the sixth suspension is lowered to less than 35 ºC, less than 34 ºC, less than 33 ºC, less than 32 ºC, less than 31 ºC, less than 30 ºC, less than 29 ºC, less than 28 ºC, less than 27 ºC, less than 26 ºC, less than 25 ºC, less than 24 ºC, less than 23 ºC, less than 22 ºC or less than 21 ºC. In some embodiments, the temperature of the sixth suspension is lowered to more than 20 ºC, more than 21 ºC, more than 22 ºC, more than 23 ºC, more than 24 ºC, more than 25 ºC, more than 26 ºC, more than 27 ºC, more than 28 ºC, more than 29 ºC, more than 30 ºC, more than 31 ºC, more than 32 ºC, more than 33 ºC or more than 34 ºC.

[00132] In some embodiments, a binder composition is formed by filtering the sixth suspension in step **107**.

[00133] Filtration aims to remove sediments or unconverted monomers that are present in the suspension in order to obtain a well-dispersed binder composition. The polymerization step in the making of binder composition comprising both structural units (a1) and (a2) derived from a first carboxylic acid group-containing monomer and a second carboxylic acid group-containing monomer respectively is observed to be capable of achieving a yield of ≥95% (Examples 4 to 23), with a negligible amount of retentate being retained during filtration. On the other hand, in the case where only one carboxylic acid group-containing monomer (Examples 1 to 3, 24 and 25) is employed in the polymerization step, a monomer conversion rate of 60-80% is resulted. This signifies a significant quantity of monomers that are not being unconverted and is likely to incur substantial expenditure, regardless of whether measures of recovery and reuse of unconverted monomers are in place. However, in spite of yielding a lower monomer conversion rate with the application of only one carboxylic acid group-containing monomer, the proportions of constituent structural units remain within the range disclosed herein. In addition, precipitation of unwanted side products is observed in the making of binder composition comprising only structural unit (a) derived from a carboxylic acid group-containing monomer, possibly emerging from formation of a relatively unstable copolymer system.

[00134] Additions of neutralizing agents in steps **101** and **106** aims to neutralize the carboxylic acid group-containing monomer added in step **102** in order to produce a binder composition that is slightly alkaline in nature. Exposing binder composition to acidic conditions is undesirable as this may disrupt the dispersion of the binder composition.

[00135] In some embodiments, structural unit (a) derived from a carboxylic acid group-containing monomer comprises a carboxylic salt group. In some embodiments, a carboxylic salt group is a salt of a carboxylic acid group. In some embodiments, structural unit (a) derived from a carboxylic acid group-containing monomer comprises a combination of a carboxylic salt group and a carboxylic acid group. In some embodiments, structural unit (a) comprises an alkali metal carboxylic salt group. Examples of an alkali metal forming the alkali metal carboxylic salt include lithium, sodium and potassium. In some embodiments, structural unit (a) comprises an ammonium carboxylic salt group.

[00136] In some embodiments, structural unit (a1) derived from a first carboxylic acid group-containing monomer comprises a carboxylic salt group. In some embodiments, a carboxylic salt group is a salt of a carboxylic acid group. In some embodiments, structural unit (a1) derived from a first carboxylic acid group-containing monomer comprises a combination of a carboxylic salt group and a carboxylic acid group. In some embodiments, structural unit (a1) comprises an alkali metal carboxylic salt group. Examples of an alkali metal forming the alkali

metal carboxylic salt include lithium, sodium and potassium. In some embodiments, structural unit (a1) comprises an ammonium carboxylic salt group.

[00137] In some embodiments, structural unit (a2) derived from a second carboxylic acid group-containing monomer comprises a carboxylic salt group. In some embodiments, a carboxylic salt group is a salt of a carboxylic acid group. In some embodiments, structural unit (a2) derived from a second carboxylic acid group-containing monomer comprises a combination of a carboxylic salt group and a carboxylic acid group. In some embodiments, structural unit (a2) comprises an alkali metal carboxylic salt group. Examples of an alkali metal forming the alkali metal carboxylic salt include lithium, sodium and potassium. In some embodiments, structural unit (a2) comprises an ammonium carboxylic salt group.

[00138] In some embodiments, when the copolymer comprises structural unit (a), the molar ratio of the carboxylic acid group to the carboxylic salt group in the copolymer is from about 0 to about 0.25, from about 0 to about 0.24, from about 0 to about 0.23, from about 0 to about 0.22, from about 0 to about 0.21, from about 0 to about 0.2, from about 0 to about 0.19, from about 0 to about 0.18, from about 0 to about 0.17, from about 0 to about 0.16, from about 0 to about 0.15, from about 0 to about 0.14, from about 0 to about 0.13, from about 0 to about 0.12, from about 0 to about 0.11, from about 0 to about 0.1, from about 0 to about 0.09, from about 0 to about 0.08, from about 0 to about 0.07, from about 0 to about 0.06 or from about 0 to about 0.05.

[00139] In some embodiments, when the copolymer comprises structural unit (a), the molar ratio of the carboxylic acid group to the carboxylic salt group in the copolymer is less than 0.25, less than 0.24, less than 0.23, less than 0.22, less than 0.21, less than 0.2, less than 0.18, less than 0.16, less than 0.14, less than 0.12, less than 0.1, less than 0.08, less than 0.06, less than 0.04 or less than 0.02. In some embodiments, the molar ratio of the carboxylic acid group to the carboxylic salt group in the copolymer is more than 0, more than 0.02, more than 0.04, more than 0.06, more than 0.08, more than 0.1, more than 0.12, more than 0.14, more than 0.16, more than 0.18, more than 0.2, more than 0.22 or more than 0.24.

[00140] In some embodiment, when the copolymer comprises structural unit (a1) and structural unit (a2), the molar ratio of the carboxylic acid group to the carboxylic salt group in the copolymer is from about 0 to about 0.25, from about 0 to about 0.24, from about 0 to about 0.23, from about 0 to about 0.22, from about 0 to about 0.21, from about 0 to about 0.2, from about 0 to about 0.19, from about 0 to about 0.18, from about 0 to about 0.17, from about 0 to about 0.16, from about 0 to about 0.15, from about 0 to about 0.14, from about 0 to about 0.13, from about 0 to about 0.12, from about 0 to about 0.11, from about 0 to about 0.1, from about 0 to about 0.09, from about 0 to about 0.08, from about 0 to about 0.07, from about 0 to about 0.06 or from about 0 to about 0.05.

[00141] In some embodiments, when the copolymer comprises structural unit (a1) and structural unit (a2), the molar ratio of the carboxylic acid group to the carboxylic salt group in the copolymer is less than 0.25, less than 0.24, less than 0.23, less than 0.22, less than 0.21, less than 0.2, less than 0.18, less than 0.16, less than 0.14, less than 0.12, less than 0.1, less than 0.08, less than 0.06, less than 0.04 or less than 0.02. In some embodiments, when the copolymer comprises structural unit (a1) and structural unit (a2), the molar ratio of the carboxylic acid group to the carboxylic salt group in the copolymer is more than 0, more than 0.02, more than 0.04, more than 0.06, more than 0.08, more than 0.1, more than 0.12, more than 0.14, more than 0.16, more than 0.18, more than 0.2, more than 0.22 or more than 0.24.

[00142] In some embodiments, the proportion of structural unit (a) in the copolymer is from about 7% to about 25%, from about 8% to about 25%, from about 9% to about 25%, from about 10% to about 25%, from about 10% to about 24%, from about 10% to about 23%, from about 10% to about 22%, from about 10% to about 21%, from about 10% to about 20%, from about 10% to about 19%, from about 10% to about 18%, from about 10% to about 17%, from about 10% to about 16%, from about 10% to about 15%, from about 12% to about 25%, from about 12% to about 20% or from about 12% to about 18% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00143] In some embodiments, the proportion of structural unit (a) in the copolymer is less than 25%, less than 24%, less than 23%, less than 22%, less than 21%, less than 20%, less than 19%, less than 18%, less than 17%, less than 16%, less than 15%, less than 14%, less than 13%, less than 12%, less than 11%, less than 10%, less than 9% or less than 8% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition. In some embodiments, the proportion of structural unit (a) in the copolymer is more than 7%, more than 8%, more than 9%, more than 10%, more than 11%, more than 12%, more than 13%, more than 14%, more than 15%, more than 16%, more than 17%, more than 18%, more than 19%, more than 20%, more than 21%, more than 22%, more than 23% or more than 24% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00144] In some embodiments, when the copolymer comprises structural unit (a1) and structural unit (a2), the proportion of structural unit (a1) in the copolymer is from about 4% to about 25%, from about 5% to about 25%, from about 6% to about 25%, from about 7% to about 25%, from about 8% to about 25%, from about 9% to about 25%, from about 10% to about 25%, from about 10% to about 24%, from about 10% to about 23%, from about 10% to about 22%, from about 10% to about 21%, from about 10% to about 20%, from about 10% to about 19%, from about 10% to about 18%, from about 10% to about 17%, from about 10% to about 16%, from about 10% to about 15%, from about 8% to about 25%, from about 8% to about 20%, from about 8% to about 18% or from about 8% to about 15% by mole, based on the total number of

moles of monomeric units in the copolymer in the binder composition.

[00145] In some embodiments, when the copolymer comprises structural unit (a1) and structural unit (a2), the proportion of structural unit (a1) in the copolymer is less than 25%, less than 24%, less than 23%, less than 22%, less than 21%, less than 20%, less than 19%, less than 18%, less than 17%, less than 16%, less than 15%, less than 14%, less than 13%, less than 12%, less than 11%, less than 10%, less than 9%, less than 8%, less than 7%, less than 6% or less than 5% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition. In some embodiments, when the copolymer comprises structural unit (a1) and structural unit (a2), the proportion of structural unit (a1) in the copolymer is more than 4%, more than 5%, more than 6%, more than 7%, more than 8%, more than 9%, more than 10%, more than 11%, more than 12%, more than 13%, more than 14%, more than 15%, more than 16%, more than 17%, more than 18%, more than 19%, more than 20%, more than 21%, more than 22%, more than 23% or more than 24% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00146] In some embodiments, when the copolymer comprises structural unit (a1) and structural unit (a2), the proportion of structural unit (a2) in the copolymer is from about 1% to about 4.5%, from about 1% to about 4.4%, from about 1% to about 4.3%, from about 1% to about 4.2%, from about 1% to about 4.1%, from about 1% to about 4%, from about 1.1% to about 4%, from about 1.2% to about 4%, from about 1.3% to about 4%, from about 1.4% to about 4%, from about 1.5% to about 4%, from about 1.6% to about 4%, from about 1.7% to about 4%, from about 1.8% to about 4%, from about 1.9% to about 4%, from about 2% to about 4%, from about 1.5% to about 4.5%, from about 1.5% to about 4% or from about 2% to about 4.5% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00147] In some embodiments, when the copolymer comprises structural unit (a1) and structural unit (a2), the proportion of structural unit (a2) in the copolymer is less than 4.5%, less than 4.4%, less than 4.2%, less than 4%, less than 3.8%, less than 3.6%, less than 3.4%, less than 3.2%, less than 3%, less than 2.8%, less than 2.6%, less than 2.4%, less than 2.2%, less than 2%, less than 1.8%, less than 1.6% or less than 1.4% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition. In some embodiments, when the copolymer comprises structural unit (a1) and structural unit (a2), the proportion of structural unit (a2) in the copolymer is more than 1%, more than 1.2%, more than 1.4%, more than 1.6%, more than 1.8%, more than 2%, more than 2.2%, more than 2.4%, more than 2.6%, more than 2.8%, more than 3%, more than 3.2%, more than 3.4%, more than 3.6%, more than 3.8%, more than 4%, more than 4.2% or more than 4.4% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00148] In some embodiments, when the copolymer comprises structural unit (a1) and structural unit (a2), the molar ratio of the structural unit (a1) to the structural unit (a2) in the copolymer is from about 1 to about 12, from about 1 to about 11.5, from about 1 to about 11, from about 1 to about 10.5, from about 1 to about 10, from about 1 to about 9.5, from about 1 to about 9, from about 1 to about 8.5, from about 1 to about 8, from about 1 to about 7.5, from about 1 to about 7, from about 1 to about 6.5, from about 1 to about 6, from about 1.5 to about 6, from about 2 to about 6, from about 2 to about 10, from about 2 to about 8, from about 2.5 to about 10, from about 2.5 to about 8, from about 3 to about 10 or from about 3 to about 8.

[00149] In some embodiments, when the copolymer comprises structural unit (a1) and structural unit (a2), the molar ratio of the structural unit (a1) to the structural unit (a2) in the copolymer is less than 12, less than 11.5, less than 11, less than 10.5, less than 10, less than 9.5, less than 9, less than 8.5, less than 8, less than 7.5, less than 7, less than 6.5, less than 6, less than 5.5, less than 5, less than 4.5, less than 4, less than 3.5, less than 3, less than 2.5 or less than 2. In some embodiments, when the copolymer comprises structural unit (a1) and structural unit (a2), the molar ratio of the structural unit (a1) to the structural unit (a2) in the copolymer is more than 1, more than 1.5, more than 2, more than 2.5, more than 3, more than 3.5, more than 4, more than 4.5, more than 5, more than 5.5, more than 6, more than 6.5, more than 7, more than 7.5, more than 8, more than 8.5, more than 9, more than 9.5, more than 10, more than 10.5 or more than 11.

[00150] In some embodiments, the proportion of structural unit (b) in the copolymer is from about 4% to about 17%, from about 4% to about 15%, from about 4% to about 10%, from about 5% to about 17%, from about 6% to about 17%, from about 7% to about 17%, from about 8% to about 17%, from about 9% to about 17%, from about 10% to about 17%, from about 10% to about 16%, from about 10% to about 15%, from about 8% to about 13%, from about 7% to about 13%, from about 12% to about 17% or from about 13% to about 17% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00151] In some embodiments, the proportion of structural unit (b) in the copolymer is less than 17%, less than 16%, less than 15%, less than 14%, less than 13%, less than 12%, less than 11%, less than 10%, less than 9%, less than 8%, less than 7%, less than 6% or less than 5% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition. In some embodiments, the proportion of structural unit (b) in the copolymer is more than 4%, more than 5%, more than 6%, more than 7%, more than 8%, more than 9%, more than 10%, more than 11%, more than 12%, more than 13%, more than 14%, more than 15% or more than 16% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00152] In some embodiments, the proportion of structural unit (c) in the copolymer is

from about 65% to about 80%, from about 65.5% to about 80%, from about 66% to about 80%, from about 66.5% to about 80%, from about 67% to about 80%, from about 67.5% to about 80%, from about 68% to about 80%, from about 68.5% to about 80%, from about 69% to about 80%, from about 69.5% to about 80%, from about 70% to about 80%, from about 70% to about 79.5%, from about 70% to about 79%, from about 70% to about 78.5%, from about 70% to about 78%, from about 70% to about 77.5%, from about 70% to about 77%, from about 70% to about 76.5%, from about 70% to about 76%, from about 70.5% to about 76%, from about 71% to about 76%, from about 71.5% to about 76%, from about 72% to about 76%, from about 67% to about 77% or from about 68% to about 75% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00153] In some embodiments, the proportion of structural unit (c) in the copolymer is more than 65%, more than 66%, more than 67%, more than 68%, more than 69%, more than 70%, more than 71%, more than 72%, more than 73%, more than 74%, more than 75%, more than 76%, more than 77%, more than 78% or more than 79% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition. In some embodiments, the proportion of structural unit (c) in the copolymer is less than 80%, less than 79%, less than 78%, less than 77%, less than 76%, less than 75%, less than 74%, less than 73%, less than 72%, less than 71%, less than 70%, less than 69%, less than 68%, less than 67% or less than 66% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00154] In some embodiments, structural unit (a) and structural unit (b) constitute as the hydrophilic portion of the copolymer. In some embodiments, structural unit (a1), structural unit (a2) and structural unit (b) constitute as the hydrophilic portion of the copolymer. In some embodiments, structural unit (c) constitutes as the hydrophobic portion of the copolymer.

[00155] In some embodiments, the proportion of the sum of structural unit (a) and structural unit (b) in the copolymer is from about 18% to about 35%, from about 18.5% to about 35%, from about 19% to about 35%, from about 19.5% to about 35%, from about 20% to about 35%, from about 20% to about 34.5%, from about 20% to about 34%, from about 20% to about 33.5%, from about 20% to about 33%, from about 20% to about 32.5%, from about 20% to about 32%, from about 20% to about 31.5%, from about 20% to about 31%, from about 20% to about 30.5%, from about 20% to about 30%, from about 20.5% to about 30%, from about 21% to about 30%, from about 21.5% to about 30%, from about 22% to about 30%, from about 22% to about 32%, from about 25% to about 35% or from about 25% to about 30% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00156] In some embodiments, the proportion of the sum of structural unit (a) and structural unit (b) in the copolymer is less than 35%, less than 34%, less than 33%, less than

32%, less than 31%, less than 30%, less than 29%, less than 28%, less than 27%, less than 26%, less than 25%, less than 24%, less than 23%, less than 22%, less than 21%, less than 20% or less than 19% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition. In some embodiments, the proportion of the sum of structural unit (a) and structural unit (b) in the copolymer is more than 18%, more than 19%, more than 20%, more than 21%, more than 22%, more than 23%, more than 24%, more than 25%, more than 26%, more than 27%, more than 28%, more than 29%, more than 30%, more than 31%, more than 32%, more than 33% or more than 34% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00157] In some embodiments, the proportion of the sum of structural unit (a1), structural unit (a2) and structural unit (b) in the copolymer is from about 18% to about 35%, from about 18.5% to about 35%, from about 19% to about 35%, from about 19.5% to about 35%, from about 20% to about 35%, from about 20% to about 34.5%, from about 20% to about 34%, from about 20% to about 33.5%, from about 20% to about 33%, from about 20% to about 32.5%, from about 20% to about 32%, from about 20% to about 31.5%, from about 20% to about 31%, from about 20% to about 30.5%, from about 20% to about 30%, from about 20.5% to about 30%, from about 21% to about 30%, from about 21.5% to about 30%, from about 22% to about 30%, from about 22% to about 32%, from about 25% to about 35% or from about 25% to about 30% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00158] In some embodiments, the proportion of the sum of structural unit (a1), structural unit (a2) and structural unit (b) in the copolymer is less than 35%, less than 34%, less than 33%, less than 32%, less than 31%, less than 30%, less than 29%, less than 28%, less than 27%, less than 26%, less than 25%, less than 24%, less than 23%, less than 22%, less than 21%, less than 20% or less than 19% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition. In some embodiments, the proportion of the sum of structural unit (a1), structural unit (a2) and structural unit (b) in the copolymer is more than 18%, more than 19%, more than 20%, more than 21%, more than 22%, more than 23%, more than 24%, more than 25%, more than 26%, more than 27%, more than 28%, more than 29%, more than 30%, more than 31%, more than 32%, more than 33% or more than 34% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

[00159] In some embodiments, the molar ratio of the structural unit (c) to the sum of the structural unit (a) and structural unit (b) in the copolymer is from about 1.5 to about 4, from about 1.6 to about 4, from about 1.7 to about 4, from about 1.8 to about 4, from about 1.9 to about 4, from about 2 to about 4, from about 2 to about 3.9, from about 2 to about 3.8, from about 2 to about 3.7, from about 2 to about 3.6, from about 2 to about 3.5, from about 2 to about

3.4, from about 2 to about 3.3, from about 2 to about 3.2, from about 2 to about 3.1, from about 2 to about 3, from about 2.2 to about 3.5 or from about 2.4 to about 3.8.

[00160] In some embodiments, the molar ratio of the structural unit (c) to the sum of the structural unit (a) and structural unit (b) in the copolymer is less than 4, less than 3.9, less than 3.8, less than 3.7, less than 3.6, less than 3.5, less than 3.4, less than 3.3, less than 3.2, less than 3.1, less than 3, less than 2.9, less than 2.8, less than 2.7, less than 2.6, less than 2.5, less than 2.4, less than 2.3, less than 2.2, less than 2.1, less than 2, less than 1.9, less than 1.8, less than 1.7 or less than 1.6. In some embodiments, the molar ratio of the structural unit (c) to the sum of the structural unit (a) and structural unit (b) in the copolymer is more than 1.5, more than 1.6, more than 1.7, more than 1.8, more than 1.9, more than 2, more than 2.1, more than 2.2, more than 2.3, more than 2.4, more than 2.5, more than 2.6, more than 2.7, more than 2.8, more than 2.9, more than 3, more than 3.1, more than 3.2, more than 3.3, more than 3.4, more than 3.5, more than 3.6, more than 3.7, more than 3.8 or more than 3.9.

[00161] In some embodiments, the molar ratio of the structural unit (c) to the sum of the structural unit (a1), structural unit (a2) and structural unit (b) in the copolymer is from about 1.5 to about 4, from about 1.6 to about 4, from about 1.7 to about 4, from about 1.8 to about 4, from about 1.9 to about 4, from about 2 to about 4, from about 2 to about 3.9, from about 2 to about 3.8, from about 2 to about 3.7, from about 2 to about 3.6, from about 2 to about 3.5, from about 2 to about 3.4, from about 2 to about 3.3, from about 2 to about 3.2, from about 2 to about 3.1, from about 2 to about 3, from about 2.2 to about 3.5 or from about 2.4 to about 3.8.

[00162] In some embodiments, the molar ratio of the structural unit (c) to the sum of the structural unit (a1), structural unit (a2) and structural unit (b) in the copolymer is less than 4, less than 3.9, less than 3.8, less than 3.7, less than 3.6, less than 3.5, less than 3.4, less than 3.3, less than 3.2, less than 3.1, less than 3, less than 2.9, less than 2.8, less than 2.7, less than 2.6, less than 2.5, less than 2.4, less than 2.3, less than 2.2, less than 2.1, less than 2, less than 1.9, less than 1.8, less than 1.7 or less than 1.6. In some embodiments, the molar ratio of the structural unit (c) to the sum of the structural unit (a1), structural unit (a2) and structural unit (b) in the copolymer is more than 1.5, more than 1.6, more than 1.7, more than 1.8, more than 1.9, more than 2, more than 2.1, more than 2.2, more than 2.3, more than 2.4, more than 2.5, more than 2.6, more than 2.7, more than 2.8, more than 2.9, more than 3, more than 3.1, more than 3.2, more than 3.3, more than 3.4, more than 3.5, more than 3.6, more than 3.7, more than 3.8 or more than 3.9.

[00163] In some embodiments, the molar ratio of the sum of the structural unit (c) and structural unit (a) to the structural unit (b) in the copolymer is from about 5 to about 15, from about 5 to about 14.75, from about 5 to about 14.5, from about 5 to about 14.25, from about 5 to about 14, from about 5 to about 13.75, from about 5 to about 13.5, from about 5 to about 13,

from about 5 to about 12.75, from about 5 to about 12.5, from about 5 to about 12.25, from about 5 to about 12, from about 5 to about 11.75, from about 5 to about 11.5, from about 5 to about 11.25, from about 5 to about 11, from about 5 to about 10.75, from about 5 to about 10.5, from about 5 to about 10.25, from about 5 to about 10, from about 5.5 to about 15, from about 6 to about 15, from about 6.5 to about 15 or from about 7 to about 15.

[00164] In some embodiments, the molar ratio of the sum of the structural unit (c) and structural unit (a) to the structural unit (b) in the copolymer is less than 15, less than 14.5, less than 14, less than 13.5, less than 13, less than 12.5, less than 12, less than 11.5, less than 11, less than 10.5, less than 10, less than 9.5, less than 9, less than 8.5, less than 8, less than 7.5, less than 7, less than 6.5, less than 6 or less than 5.5. In some embodiments, the molar ratio of the sum of the structural unit (c) and structural unit (a) to the structural unit (b) in the copolymer is more than 5, more than 5.5, more than 6, more than 6.5, more than 7, more than 7.5, more than 8, more than 8.5, more than 9, more than 9.5, more than 10, more than 10.5, more than 11, more than 11.5, more than 12, more than 12.5, more than 13, more than 13.5, more than 14 or more than 14.5.

[00165] In some embodiments, the molar ratio of the sum of the structural unit (c), structural unit (a1) and structural unit (a2) to the structural unit (b) in the copolymer is from about 5 to about 15, from about 5 to about 14.75, from about 5 to about 14.5, from about 5 to about 14.25, from about 5 to about 14, from about 5 to about 13.75, from about 5 to about 13.5, from about 5 to about 13, from about 5 to about 12.75, from about 5 to about 12.5, from about 5 to about 12.25, from about 5 to about 12, from about 5 to about 11.75, from about 5 to about 11.5, from about 5 to about 11.25, from about 5 to about 11, from about 5 to about 10.75, from about 5 to about 10.5, from about 5 to about 10.25, from about 5 to about 10, from about 5.5 to about 15, from about 6 to about 15, from about 6.5 to about 15 or from about 7 to about 15.

[00166] In some embodiments, the molar ratio of the sum of the structural unit (c), structural unit (a1) and structural unit (a2) to the structural unit (b) in the copolymer is less than 15, less than 14.5, less than 14, less than 13.5, less than 13, less than 12.5, less than 12, less than 11.5, less than 11, less than 10.5, less than 10, less than 9.5, less than 9, less than 8.5, less than 8, less than 7.5, less than 7, less than 6.5, less than 6 or less than 5.5. In some embodiments, the molar ratio of the sum of the structural unit (c), structural unit (a1) and structural unit (a2) to the structural unit (b) in the copolymer is more than 5, more than 5.5, more than 6, more than 6.5, more than 7, more than 7.5, more than 8, more than 8.5, more than 9, more than 9.5, more than 10, more than 10.5, more than 11, more than 11.5, more than 12, more than 12.5, more than 13, more than 13.5, more than 14 or more than 14.5.

[00167] Addition of ester group-containing monomer in the preparation of the binder composition disclosed herein is found to result in a deterioration in electrochemical

performance. In some embodiments, the binder composition is free of structural unit derived from an ester group-containing monomer. In some embodiments, the ester group-containing monomer is C_1 to C_{20} alkyl acrylate, C_1 to C_{20} alkyl (meth)acrylate, cycloalkyl acrylate or a combination thereof. In some embodiments, the ester group-containing monomer is methyl acrylate, ethyl acrylate, n-propyl acrylate, isopropyl acrylate, n-butyl acrylate, sec-butyl acrylate, tert-butyl acrylate, pentyl acrylate, hexyl acrylate, heptyl acrylate, octyl acrylate, 3,3,5-trimethylhexyl acrylate, 2-ethylhexyl acrylate, nonyl acrylate, decyl acrylate, lauryl acrylate, n-tetradecyl acrylate, oxtadecyl acrylate, cyclohexyl acrylate, phenyl acrylate, methoxymethyl acrylate, methoxyethyl acrylate, ethoxymethyl acrylate, ethoxyethyl acrylate, perfluorooctyl acrylate, stearyl acrylate or a combination thereof. In some embodiments, the ester group-containing monomer is cyclohexyl acrylate, cyclohexyl methacrylate, isobornyl acrylate, isobornyl methacrylate, 3,3,5-trimethylcyclohexylacrylate, or a combination thereof. In some embodiments, the ester group-containing monomer is methyl methacrylate, ethyl methacrylate, n-propyl methacrylate, isopropyl methacrylate, n-butyl methacrylate, sec-butyl methacrylate, tert-butyl methacrylate, isobutyl methacrylate, n-pentyl methacrylate, isopentyl methacrylate, hexyl methacrylate, heptyl methacrylate, octyl methacrylate, 2-ethylhexyl methacrylate, nonyl methacrylate, decyl methacrylate, lauryl methacrylate, n-tetradecyl methacrylate, stearyl methacrylate, 2,2,2-trifluoroethyl methacrylate, phenyl methacrylate, benzyl methacrylate, or a combination thereof.

[00168] In some embodiments, the binder composition is free of structural unit derived from a conjugated diene group-containing monomer. Examples of conjugated diene group-containing monomer include aliphatic conjugated diene monomers such as 1,3-butadiene, 2-methyl-1,3-butadiene, 2,3-dimethyl-1,3-butadiene, 2-chloro-1,3-butadiene, substituted linear conjugated pentadienes, and substituted side chain conjugated hexadienes.

[00169] In some embodiments, the binder composition is free of structural unit derived from an aromatic vinyl group-containing monomer. Examples of aromatic vinyl group-containing monomer include styrene, α-methylstyrene, vinyltoluene, and divinylbenzene.

[00170] In some embodiments, the pH of the binder composition is from about 7 to about 9, from about 7 to about 8.9, from about 7 to about 8.8, from about 7 to about 8.7, from about 7 to about 8.6, from about 7 to about 8.5, from about 7 to about 8.4, from about 7 to about 8.3, from about 7 to about 8.2, from about 7 to about 8.1, from about 7 to about 8, from about 7.1 to about 9, from about 7.2 to about 9, from about 7.3 to about 9, from about 7.4 to about 9, from about 7.5 to about 9, from about 7.6 to about 9, from about 7.7 to about 9, from about 7.8 to about 9, from about 7.9 to about 9 or from about 8 to about 9.

[00171] In certain embodiments, the pH of the binder composition is less than 9, less than 8.9, less than 8.8, less than 8.7, less than 8.6, less than 8.5, less than 8.4, less than 8.3, less than

8.2, less than 8.1, less than 8, less than 7.9, less than 7.8, less than 7.7, less than 7.6, less than 7.5, less than 7.4, less than 7.3 or less than 7.2. In certain embodiments, the pH of the binder composition is more than 7, more than 7.1, more than 7.2, more than 7.3, more than 7.4, more than 7.5, more than 7.6, more than 7.7, more than 7.8, more than 7.9, more than 8, more than 8.1, more than 8.2, more than 8.3, more than 8.4, more than 8.5, more than 8.6, more than 8.7 or more than 8.8.

[00172] In some embodiments, the viscosity of the binder composition is from about 10,000 mPa·s to about 50,000 mPa·s, from about 10,000 mPa·s to about 47,500 mPa·s, from about 10,000 mPa·s to about 45,000 mPa·s, from about 10,000 mPa·s to about 42,500 mPa·s, from about 10,000 mPa·s to about 40,000 mPa·s, from about 10,000 mPa·s to about 37,500 mPa·s, from about 10,000 mPa·s to about 35,000 mPa·s, from about 10,000 mPa·s to about 32,500 mPa·s, from about 10,000 mPa·s to about 30,000 mPa·s, from about 10,000 mPa·s to about 29,000 mPa·s, from about 10,000 mPa·s to about 28,000 mPa·s, from about 10,000 mPa·s to about 27,000 mPa·s, from about 10,000 mPa·s to about 26,000 mPa·s, from about 10,000 mPa·s to about 25,000 mPa·s, from about 10,000 mPa·s to about 24,000 mPa·s, from about 10,000 mPa·s to about 23,000 mPa·s, from about 10,000 mPa·s to about 22,000 mPa·s, from about 10,000 mPa·s to about 21,000 mPa·s, from about 10,000 mPa·s to about 20,000 mPa·s, from about 15,000 mPa·s to about 30,000 mPa·s, from about 15,000 mPa·s to about 25,000 mPa·s, from about 15,000 mPa·s to about 35,000 mPa·s.

[00173] In some embodiments, the viscosity of the binder composition is less than 50,000 mPa·s, less than 47,500 mPa·s, less than 45,000 mPa·s, less than 42,500 mPa·s, less than 40,000 mPa·s, less than 37,500 mPa·s, less than 35,000 mPa·s, less than 32,500 mPa·s, less than 30,000 mPa·s, less than 27,500 mPa·s, less than 25,000 mPa·s, less than 22,500 mPa·s, less than 20,000 mPa·s, less than 17,500 mPa·s, less than 15,000 mPa·s or less than 12,500 mPa·s. In some embodiments, the viscosity of the binder composition is more than 10,000 mPa·s, more than 12,500 mPa·s, more than 15,000 mPa·s, more than 17,500 mPa·s, more than 20,000 mPa·s, more than 22,500 mPa·s, more than 25,000 mPa·s, more than 27,500 mPa·s, more than 30,000 mPa·s, more than 32,500 mPa·s, more than 35,000 mPa·s, more than 37,500 mPa·s, more than 40,000 mPa·s, more than 42,500 mPa·s, more than 45,000 mPa·s, more than 47,500 mPa·s.

[00174] In some embodiments, the solid content of the binder composition is from about 12% to about 18%, from about 12.2% to about 18%, from about 12.4% to about 18%, from about 12.6% to about 18%, from about 12.8% to about 18%, from about 13% to about 18%, from about 13% to about 17.8%, from about 13% to about 17.6%, from about 13% to about 17.4%, from about 13% to about 17.2%, from about 13% to about 17%, from about 13.1% to about 17%, from about 13.2% to about 17%, from about 13.3% to about 17%, from about 13.4% to about 17%, from about 13.5% to about 17%, from about 13.6% to about 17%, from about 13.7% to about 17%, from about 13.8% to about 17%, from about 13.9% to about 17%, from

about 14% to about 16.9%, from about 14% to about 16.8%, from about 14% to about 16.7%, from about 14% to about 16.6%, from about 14% to about 16.5%, from about 14% to about 16.4%, from about 14% to about 16.3%, from about 14% to about 16.2%, from about 14% to about 16.1%, from about 14% to about 16% by weight, based on the total weight of the binder composition.

[00175] In some embodiments, the solid content of the binder composition is less than 18%, less than 17.8%, less than 17.6%, less than 17.4%, less than 17.2%, less than 17%, less than 16.8%, less than 16.6%, less than 16.4%, less than 16.2%, less than 16%, less than 15.8%, less than 15.6%, less than 15.4%, less than 15.2%, less than 15%, less than 14.8%, less than 14.6%, less than 14.4%, less than 14.2%, less than 14%, less than 13.8%, less than 13.6%, less than 13.4%, less than 13.2%, less than 13%, less than 12.8%, less than 12.6%, less than 12.4% or less than 12.2% by weight, based on the total weight of the binder composition. In some embodiments, the solid content of the binder composition is more than 12%, more than 12.2%, more than 12.4%, more than 12.6%, more than 12.8%, more than 13%, more than 13.2%, more than 13.4%, more than 13.6%, more than 13.8%, more than 14%, more than 14.2%, more than 14.4%, more than 14.6%, more than 14.8%, more than 15%, more than 15.2%, more than 15.4%, more than 15.6%, more than 15.8%, more than 16%, more than 16.2%, more than 16.4%, more than 16.6%, more than 16.8%, more than 17%, more than 17.2%, more than 17.4%, more than 17.6% or more than 17.8% by weight, based on the total weight of the binder composition.

[00176] In some embodiments, the weight average molecular weight of the binder composition is from about 100,000 g/mol to about 200,000 g/mol, from about 105,000 g/mol to about 200,000 g/mol, from about 110,000 g/mol to about 200,000 g/mol, from about 115,000 g/mol to about 200,000 g/mol, from about 120,000 g/mol to about 200,000 g/mol, from about 125,000 g/mol to about 200,000 g/mol, from about 130,000 g/mol to about 200,000 g/mol, from about 130,000 g/mol to about 195,000 g/mol, from about 130,000 g/mol to about 190,000 g/mol, from about 130,000 g/mol to about 185,000 g/mol, from about 130,000 g/mol to about 180,000 g/mol, from about 130,000 g/mol to about 175,000 g/mol, from about 130,000 g/mol to about 170,000 g/mol, from about 135,000 g/mol to about 170,000 g/mol, from about 140,000 g/mol to about 170,000 g/mol, from about 145,000 g/mol to about 170,000 g/mol, from about 150,000 g/mol to about 170,000 g/mol, from about 150,000 g/mol to about 165,000 g/mol or from about 155,000 g/mol to about 165,000 g/mol. When the weight average molecular weight of the binder composition is not more than the upper limit set forth above, a smooth binder composition layer can be obtained because coatability of the binder composition is ensured, and adhesive strength of the binder composition can be improved. On the other hand, when the weight average molecular weight of the binder composition is not less than the lower limit set forth above, binding capability of the binder composition can be ensured, and adhesive strength of the binder

composition and secondary battery cycle characteristics can be improved.

[00177] In some embodiments, the weight average molecular weight of the binder composition is less than 200,000 g/mol, less than 195,000 g/mol, less than 190,000 g/mol, less than 185,000 g/mol, less than 180,000 g/mol, less than 175,000 g/mol, less than 170,000 g/mol, less than 165,000 g/mol, less than 160,000 g/mol, less than 155,000 g/mol, less than 150,000 g/mol, less than 145,000 g/mol, less than 140,000 g/mol, less than 135,000 g/mol, less than 130,000 g/mol, less than 125,000 g/mol, less than 120,000 g/mol, less than 115,000 g/mol, less than 110,000 g/mol or less than 105,000 g/mol. In some embodiments, the weight average molecular weight of the binder composition is more than 100,000 g/mol, more than 105,000 g/mol, more than 110,000 g/mol, more than 115,000 g/mol, more than 120,000 g/mol, more than 125,000 g/mol, more than 130,000 g/mol, more than 135,000 g/mol, more than 140,000 g/mol, more than 145,000 g/mol, more than 150,000 g/mol, more than 155,000 g/mol, more than 160,000 g/mol, more than 165,000 g/mol, more than 170,000 g/mol, more than 175,000 g/mol, more than 180,000 g/mol, more than 185,000 g/mol, more than 190,000 g/mol or more than 195,000 g/mol.

[00178] In some embodiments, the number average molecular weight of the binder composition is from about 10,000 g/mol to about 100,000 g/mol, from about 15,000 g/mol to about 100,000 g/mol, from about 20,000 g/mol to about 100,000 g/mol, from about 25,000 g/mol to about 100,000 g/mol, from about 30,000 g/mol to about 100,000 g/mol, from about 35,000 g/mol to about 100,000 g/mol, from about 40,000 g/mol to about 100,000 g/mol, from about 45,000 g/mol to about 100,000 g/mol, from about 50,000 g/mol to about 100,000 g/mol, from about 50,000 g/mol to about 95,000 g/mol, from about 50,000 g/mol to about 90,000 g/mol, from about 50,000 g/mol to about 85,000 g/mol, from about 50,000 g/mol to about 80,000 g/mol, from about 55,000 g/mol to about 80,000 g/mol, from about 60,000 g/mol to about 80,000 g/mol, from about 65,000 g/mol to about 75,000 g/mol or from about 60,000 g/mol to about 90,000 g/mol.

[00179] In some embodiments, the number average molecular weight of the binder composition is less than 100,000 g/mol, less than 95,000 g/mol, less than 90,000 g/mol, less than 85,000 g/mol, less than 80,000 g/mol, less than 75,000 g/mol, less than 70,000 g/mol, less than 65,000 g/mol, less than 60,000 g/mol, less than 55,000 g/mol, less than 50,000 g/mol, less than 45,000 g/mol, less than 40,000 g/mol, less than 35,000 g/mol, less than 30,000 g/mol, less than 25,000 g/mol, less than 20,000 g/mol or less than 15,000 g/mol. In some embodiments, the number average molecular weight of the binder composition is more than 10,000 g/mol, more than 15,000 g/mol, more than 20,000 g/mol, more than 25,000 g/mol, more than 30,000 g/mol, more than 35,000 g/mol, more than 40,000 g/mol, more than 45,000 g/mol, more than 50,000 g/mol, more than 55,000 g/mol, more than 60,000 g/mol, more than 65,000 g/mol, more than 70,000 g/mol, more than 75,000 g/mol, more than 80,000 g/mol, more than 85,000 g/mol, more

than 90,000 g/mol or more than 95,000 g/mol.

[00180] In some embodiments, the polydispersity index (PDI) of the binder composition is from about 1 to about 5, from about 1 to about 4.8, from about 1 to about 4.6, from about 1 to about 4.4, from about 1 to about 4.2, from about 1 to about 4, from about 1 to about 3.8, from about 1 to about 3.6, from about 1 to about 3.4, from about 1 to about 3.2, from about 1 to about 3, from about 1.1 to about 3, from about 1.2 to about 3, from about 1.3 to about 3, from about 1.4 to about 3, from about 1.5 to about 3, from about 1.6 to about 3, from about 1.6 to about 2.8, from about 1.6 to about 2.6, from about 1.8 to about 2.6 or from about 1.8 to about 2.8. Stability of the binder composition can be further improved when the polydispersity index of the binder composition is within the range set forth above.

[00181] In some embodiments, the polydispersity index of the binder composition is less than 5, less than 4.8, less than 4.6, less than 4.4, less than 4.2, less than 4, less than 3.8, less than 3.6, less than 3.4, less than 3.2, less than 3, less than 2.8, less than 2.6, less than 2.4, less than 2.2, less than 2, less than 1.8, less than 1.6, less than 1.4 or less than 1.2. In some embodiments, the polydispersity index of the binder composition is more than 1, more than 1.2, more than 1.4, more than 1.6, more than 1.8, more than 2, more than 2.2, more than 2.4, more than 2.6, more than 2.8, more than 3, more than 3.2, more than 3.4, more than 3.6, more than 3.8, more than 4, more than 4.2, more than 4.4, more than 4.6 or more than 4.8.

[00182] In some embodiments, the average particle diameter of the binder composition is from about 10 μm to about 50 μm, from about 12 μm to about 50 μm, from about 14 μm to about 50 μm, from about 16 μm to about 50 μm, from about 18 μm to about 50 μm, from about 20 μm to about 50 μm, from about 20 μm to about 48 μm, from about 20 μm to about 46 μm, from about 20 μm to about 44 μm, from about 20 μm to about 42 μm, from about 20 μm to about 40 μm, from about 22 μm to about 40 μm, from about 22 μm to about 38 μm, from about 24 μm to about 38 μm, from about 24 μm to about 36 μm, from about 26 μm to about 34 μm, from about 28 μm to about 34 μm or from about 28 μm to about 32 μm.

[00183] In some embodiments, the average particle diameter of the binder composition is less than 50 μm, less than 48 μm, less than 46 μm, less than 44 μm, less than 42 μm, less than 40 μm, less than 38 μm, less than 36 μm, less than 34 μm, less than 32 μm, less than 30 μm, less than 28 μm, less than 26 μm, less than 24 μm, less than 22 μm, less than 20 μm, less than 18 μm, less than 16 μm, less than 14 μm or less than 12 μm. In some embodiments, the average particle diameter of the binder composition is more than 10 μm, more than 12 μm, more than 14 μm, more than 16 μm, more than 18 μm, more than 20 μm, more than 22 μm, more than 24 μm, more than 26 μm, more than 28 μm, more than 30 μm, more than 32 μm, more than 34 μm, more than 36 μm, more than 38 μm, more than 40 μm, more than 42 μm, more than 44 μm, more than 46 μm or more than 48 μm.

[00184] In some embodiments, the D50 of the binder composition is from about 1 μm to about 100 μm, from about 1 μm to about 98 μm, from about 1 μm to about 96 μm, from about 1 μm to about 94 μm, from about 1 μm to about 92 μm, from about 1 μm to about 90 μm, from about 1 μm to about 88 μm, from about 1 μm to about 86 μm, from about 1 μm to about 84 μm, from about 1 μm to about 82 μm, from about 1 μm to about 80 μm, from about 1 μm to about 75 μm, from about 1 μm to about 70 μm, from about 1 μm to about 65 μm, from about 1 μm to about 60 μm, from about 1 μm to about 55 μm, from about 1 μm to about 50 μm, from about 1 μm to about 45 μm, from about 1 μm to about 40 μm, from about 1 μm to about 35 μm, from about 1 μm to about 30 μm, from about 1 μm to about 25 μm, from about 2 μm to about 25 μm, from about 3 μm to about 25 μm, from about 4 μm to about 25 μm or from about 5 μm to about 25 μm.

[00185] In some embodiments, the D50 of the binder composition is less than 100 μm, less than 95 μm, less than 90 μm, less than 85 μm, less than 80 μm, less than 75 μm, less than 70 μm, less than 65 μm, less than 60 μm, less than 55 μm, less than 50 μm, less than 45 μm, less than 40 μm, less than 35 μm, less than 30 μm, less than 25 μm, less than 20 μm, less than 15 μm, less than 10 μm or less than 5 μm. In some embodiments, the D50 of the binder composition is more than 1 μm, more than 5 μm, more than 10 μm, more than 15 μm, more than 20 μm, more than 25 μm, more than 30 μm, more than 35 μm, more than 40 μm, more than 45 μm, more than 50 μm, more than 55 μm, more than 60 μm, more than 65 μm, more than 70 μm, more than 75 μm, more than 80 μm, more than 85 μm, more than 90 μm or more than 95 μm.

[00186] In some embodiments, the D10 of the binder composition is from about 0.1 μm to about 20 μm, from about 0.1 μm to about 19.5 μm, from about 0.1 μm to about 19 μm, from about 0.1 μm to about 18.5 μm, from about 0.1 μm to about 18 μm, from about 0.1 μm to about 17.5 μm, from about 0.1 μm to about 17 μm, from about 0.1 μm to about 16.5 μm, from about 0.1 μm to about 16 μm, from about 0.1 μm to about 15.5 μm, from about 0.1 μm to about 15 μm, from about 0.1 μm to about 14.5 μm, from about 0.1 μm to about 14 μm, from about 0.1 μm to about 13.5 μm, from about 0.1 μm to about 13 μm, from about 0.1 μm to about 12.5 μm, from about 0.1 μm to about 12 μm, from about 0.1 μm to about 11.5 μm, from about 0.1 μm to about 11 μm, from about 0.1 μm to about 10.5 μm or from about 0.1 μm to about 10 μm.

[00187] In some embodiments, the D10 of the binder composition is less than 20 μm, less than 19 μm, less than 18 μm, less than 17 μm, less than 16 μm, less than 15 μm, less than 14 μm, less than 13 μm, less than 12 μm, less than 11 μm, less than 10 μm, less than 9 μm, less than 8 μm, less than 7 μm, less than 6 μm, less than 5 μm, less than 4 μm, less than 3 μm, less than 2 μm, less than 1 μm or less than 0.5 μm. In some embodiments, the D10 of the binder composition is more than 0.1 μm, more than 0.5 μm, more than 1 μm, more than 2 μm, more than 3 μm, more than 4 μm, more than 5 μm, more than 6 μm, more than 7 μm, more than 8 μm, more than 9 μm, more than 10 μm, more than 11 μm, more than 12 μm, more than 13 μm, more

than 14 μm, more than 15 μm, more than 16 μm, more than 17 μm, more than 18 μm or more than 19 μm.

[00188] In some embodiments, the D90 of the binder composition is from about 10 μm to about 300 μm, from about 15 μm to about 300 μm, from about 20 μm to about 300 μm, from about 25 μm to about 300 μm, from about 30 μm to about 300 μm, from about 35 μm to about 300 μm, from about 40 μm to about 300 μm, from about 45 μm to about 300 μm, from about 50 μm to about 300 μm, from about 60 μm to about 300 μm, from about 70 μm to about 300 μm, from about 80 μm to about 300 μm, from about 90 μm to about 300 μm, from about 100 μm to about 300 μm, from about 120 μm to about 300 μm, from about 140 μm to about 300 μm, from about 160 μm to about 300 μm, from about 180 μm to about 300 μm, from about 200 μm to about 300 μm, from about 220 μm to about 300 μm or from about 240 μm to about 300 μm.

[00189] In some embodiments, the D90 of the binder composition is less than 300 μm, less than 295 μm, less than 290 μm, less than 285 μm, less than 280 μm, less than 275 μm, less than 270 μm, less than 265 μm, less than 260 μm, less than 255 μm, less than 250 μm, less than 225 μm, less than 200 μm, less than 175 μm, less than 150 μm, less than 125 μm, less than 100 μm, less than 75 μm, less than 50 μm, less than 25 μm or less than 15 μm. In some embodiments, the D90 of the binder composition is more than 10 μm, more than 15 μm, more than 20 μm, more than 25 μm, more than 30 μm, more than 35 μm, more than 40 μm, more than 45 μm, more than 50 μm, more than 75 μm, more than 100 μm, more than 125 μm, more than 150 μm, more than 175 μm, more than 200 μm, more than 225 μm, more than 250 μm or more than 275 μm.

[00190] The binder composition of the present invention exhibits strong adhesion to the current collector. It is important for the binder composition to have good adhesive strength to the current collector as it promotes the binding force of the electrode layer to the current collector in the making of battery electrode, prevents separation and enhances the mechanical stability of the electrode. In some embodiments, the adhesive strength between the binder composition and the current collector is from about 2 N/cm to about 4 N/cm, from about 2.1 N/cm to about 4 N/cm, from about 2.2 N/cm to about 4 N/cm, from about 2.3 N/cm to about 4 N/cm, from about 2.4 N/cm to about 4 N/cm, from about 2.5 N/cm to about 4 N/cm, from about 2.6 N/cm to about 4 N/cm, from about 2.7 N/cm to about 4 N/cm, from about 2.8 N/cm to about 4 N/cm, from about 2.9 N/cm to about 4 N/cm, from about 3 N/cm to about 4 N/cm, from about 2 N/cm to about 3.9 N/cm, from about 2 N/cm to about 3.8 N/cm, from about 2 N/cm to about 3.7 N/cm, from about 2 N/cm to about 3.6 N/cm, from about 2 N/cm to about 3.5 N/cm, from about 2 N/cm to about 3.4 N/cm, from about 2 N/cm to about 3.3 N/cm, from about 2 N/cm to about 3.2 N/cm, from about 2 N/cm to about 3.1 N/cm, from about 2 N/cm to about 3 N/cm, from about 2.5 N/cm to about 3.5 N/cm, from about 2.3 N/cm to about 3.7 N/cm, from about 2.5 N/cm to about 3 N/cm or from about 3 N/cm to about 3.5 N/cm.

[00191] In some embodiments, the adhesive strength between the binder composition and the current collector is less than 4 N/cm, less than 3.9 N/cm, less than 3.8 N/cm, less than 3.7 N/cm, less than 3.6 N/cm, less than 3.5 N/cm, less than 3.4 N/cm, less than 3.3 N/cm, less than 3.2 N/cm, less than 3.1 N/cm, less than 3 N/cm, less than 2.9 N/cm, less than 2.8 N/cm, less than 2.7 N/cm, less than 2.6 N/cm, less than 2.5 N/cm, less than 2.4 N/cm, less than 2.3 N/cm or less than 2.2 N/cm. In some embodiments, the adhesive strength between the binder composition and the current collector is more than 2 N/cm, more than 2.1 N/cm, more than 2.2 N/cm, more than 2.3 N/cm, more than 2.4 N/cm, more than 2.5 N/cm, more than 2.6 N/cm, more than 2.7 N/cm, more than 2.8 N/cm, more than 2.9 N/cm, more than 3 N/cm, more than 3.1 N/cm, more than 3.2 N/cm, more than 3.3 N/cm, more than 3.4 N/cm, more than 3.5 N/cm, more than 3.6 N/cm, more than 3.7 N/cm or more than 3.8 N/cm.

[00192] In another aspect, provided herein is an electrode for a secondary battery, comprising an electrode active material, a current collector and the binder composition prepared by the method described above. In other embodiments, the electrode further comprises a conductive agent.

[00193] In some embodiments, the electrode active material is a cathode active material, wherein the cathode active material is selected from the group consisting of $LiCoO_2$, $LiNiO_2$, $LiNi_xMn_yO_2$, $Li_{1+z}Ni_xMn_yCo_{1-x-y}O_2$, $LiNi_xCo_yAl_zO_2$, LiV_2O_5, $LiTiS_2$, $LiMoS_2$, $LiMnO_2$, $LiCrO_2$, $LiMn_2O_4$, Li_2MnO_3, $LiFeO_2$, $LiFePO_4$, and combinations thereof, wherein each x is independently from 0.2 to 0.9; each y is independently from 0.1 to 0.45; and each z is independently from 0 to 0.2. In certain embodiments, the cathode active material is selected from the group consisting of $LiCoO_2$, $LiNiO_2$, $LiNi_xMn_yO_2$, $Li_{1+z}Ni_xMn_yCo_{1-x-y}O_2$ (NMC), $LiNi_xCo_yAl_zO_2$, LiV_2O_5, $LiTiS_2$, $LiMoS_2$, $LiMnO_2$, $LiCrO_2$, $LiMn_2O_4$, $LiFeO_2$, $LiFePO_4$, and combinations thereof, wherein each x is independently from 0.4 to 0.6; each y is independently from 0.2 to 0.4; and each z is independently from 0 to 0.1. In other embodiments, the cathode active material is not $LiCoO_2$, $LiNiO_2$, LiV_2O_5, $LiTiS_2$, $LiMoS_2$, $LiMnO_2$, $LiCrO_2$, $LiMn_2O_4$, $LiFeO_2$, or $LiFePO_4$. In further embodiments, the cathode active material is not $LiNi_xMn_yO_2$, $Li_{1+z}Ni_xMn_yCo_{1-x-y}O_2$, or $LiNi_xCo_yAl_zO_2$, wherein each x is independently from 0.2 to 0.9; each y is independently from 0.1 to 0.45; and each z is independently from 0 to 0.2. In certain embodiments, the cathode active material is $Li_{1+x}Ni_aMn_bCo_cAl_{(1-a-b-c)}O_2$; wherein $-0.2 \leq x \leq 0.2$, $0 \leq a < 1$, $0 \leq b < 1$, $0 \leq c < 1$, and $a+b+c \leq 1$. In some embodiments, the cathode active material has the general formula $Li_{1+x}Ni_aMn_bCo_cAl_{(1-a-b-c)}O_2$, with $0.33 \leq a \leq 0.92$, $0.33 \leq a \leq 0.9$, $0.33 \leq a \leq 0.8$, $0.5 \leq a \leq 0.92$, $0.5 \leq a \leq 0.9$, $0.5 \leq a \leq 0.8$, $0.6 \leq a \leq 0.92$, or $0.6 \leq a \leq 0.9$; $0 \leq b \leq 0.5$, $0 \leq b \leq 0.3$, $0.1 \leq b \leq 0.5$, $0.1 \leq b \leq 0.4$, $0.1 \leq b \leq 0.3$, $0.1 \leq b \leq 0.2$, or $0.2 \leq b \leq 0.5$; $0 \leq c \leq 0.5$, $0 \leq c \leq 0.3$, $0.1 \leq c \leq 0.5$, $0.1 \leq c \leq 0.4$, $0.1 \leq c \leq 0.3$, $0.1 \leq c \leq 0.2$, or $0.2 \leq c \leq 0.5$.

[00194] In certain embodiments, the cathode active material is doped with a dopant selected from the group consisting of Fe, Ni, Mn, Al, Mg, Zn, Ti, La, Ce, Sn, Zr, Ru, Si, Ge, and

combinations thereof. In some embodiments, the dopant is not Fe, Ni, Mn, Mg, Zn, Ti, La, Ce, Ru, Si, or Ge. In certain embodiments, the dopant is not Al, Sn, or Zr.

[00195] In some embodiments, the cathode active material is $LiNi_{0.33}Mn_{0.33}Co_{0.33}O_2$ (NMC333), $LiNi_{0.4}Mn_{0.4}Co_{0.2}O_2$, $LiNi_{0.5}Mn_{0.3}Co_{0.2}O_2$ (NMC532), $LiNi_{0.6}Mn_{0.2}Co_{0.2}O_2$ (NMC622), $LiNi_{0.7}Mn_{0.15}Co_{0.15}O_2$, $LiNi_{0.8}Mn_{0.1}Co_{0.1}O_2$ (NMC811), $LiNi_{0.92}Mn_{0.04}Co_{0.04}O_2$, $LiNi_{0.8}Co_{0.15}Al_{0.05}O_2$ (NCA), $LiNiO_2$ (LNO), and combinations thereof.

[00196] In other embodiments, the cathode active material is not $LiCoO_2$, $LiNiO_2$, $LiMnO_2$, $LiMn_2O_4$, or Li_2MnO_3. In further embodiments, the cathode active material is not $LiNi_{0.33}Mn_{0.33}Co_{0.33}O_2$, $LiNi_{0.4}Mn_{0.4}Co_{0.2}O_2$, $LiNi_{0.5}Mn_{0.3}Co_{0.2}O_2$, $LiNi_{0.6}Mn_{0.2}Co_{0.2}O_2$, $LiNi_{0.7}Mn_{0.15}Co_{0.15}O_2$, $LiNi_{0.8}Mn_{0.1}Co_{0.1}O_2$, $LiNi_{0.92}Mn_{0.04}Co_{0.04}O_2$, or $LiNi_{0.8}Co_{0.15}Al_{0.05}O_2$.

[00197] In certain embodiments, the cathode active material comprises or is a core-shell composite having a core and shell structure, wherein the core and the shell each independently comprise a lithium transition metal oxide selected from the group consisting of $Li_{1+x}Ni_aMn_bCo_cAl_{(1-a-b-c)}O_2$, $LiCoO_2$, $LiNiO_2$, $LiMnO_2$, $LiMn_2O_4$, Li_2MnO_3, $LiCrO_2$, $Li_4Ti_5O_{12}$, LiV_2O_5, $LiTiS_2$, $LiMoS_2$, and combinations thereof; wherein $-0.2 \leq x \leq 0.2$, $0 \leq a < 1$, $0 \leq b < 1$, $0 \leq c < 1$, and $a+b+c \leq 1$. In other embodiments, the core and the shell each independently comprise two or more lithium transition metal oxides. In some embodiments, one of the core or shell comprises only one lithium transition metal oxide, while the other comprises two or more lithium transition metal oxides. The lithium transition metal oxide or oxides in the core and the shell may be the same, or they may be different or partially different. In some embodiments, the two or more lithium transition metal oxides are uniformly distributed over the core. In certain embodiments, the two or more lithium transition metal oxides are not uniformly distributed over the core. In some embodiments, the cathode active material is not a core-shell composite.

[00198] In some embodiments, each of the lithium transition metal oxides in the core and the shell is independently doped with a dopant selected from the group consisting of Fe, Ni, Mn, Al, Mg, Zn, Ti, La, Ce, Sn, Zr, Ru, Si, Ge, and combinations thereof. In certain embodiments, the core and the shell each independently comprise two or more doped lithium transition metal oxides. In some embodiments, the two or more doped lithium transition metal oxides are uniformly distributed over the core and/or the shell. In certain embodiments, the two or more doped lithium transition metal oxides are not uniformly distributed over the core and/or the shell.

[00199] In some embodiments, the cathode active material comprises or is a core-shell composite comprising a core comprising a lithium transition metal oxide and a shell comprising a transition metal oxide. In certain embodiments, the lithium transition metal oxide is selected from the group consisting of $Li_{1+x}Ni_aMn_bCo_cAl_{(1-a-b-c)}O_2$, $LiCoO_2$, $LiNiO_2$, $LiMnO_2$, $LiMn_2O_4$, Li_2MnO_3, $LiCrO_2$, $Li_4Ti_5O_{12}$, LiV_2O_5, $LiTiS_2$, $LiMoS_2$, and combinations thereof; wherein $-0.2 \leq x \leq 0.2$, $0 \leq a < 1$, $0 \leq b < 1$, $0 \leq c < 1$, and $a+b+c \leq 1$. In some embodiments, the transition metal

oxide is selected from the group consisting of Fe_2O_3, MnO_2, Al_2O_3, MgO, ZnO, TiO_2, La_2O_3, CeO_2, SnO_2, ZrO_2, RuO_2, and combinations thereof. In certain embodiments, the shell comprises a lithium transition metal oxide and a transition metal oxide.

[00200] In some embodiments, the diameter of the core is from about 1 μm to about 15 μm, from about 3 μm to about 15 μm, from about 3 μm to about 10 μm, from about 5 μm to about 10 μm, from about 5 μm to about 45 μm, from about 5 μm to about 35 μm, from about 5 μm to about 25 μm, from about 10 μm to about 45 μm, from about 10 μm to about 40 μm, or from about 10 μm to about 35 μm, from about 10 μm to about 25 μm, from about 15 μm to about 45 μm, from about 15 μm to about 30 μm, from about 15 μm to about 25 μm, from about 20 μm to about 35 μm, or from about 20 μm to about 30 μm. In certain embodiments, the thickness of the shell is from about 1 μm to about 45 μm, from about 1 μm to about 35 μm, from about 1 μm to about 25 μm, from about 1 μm to about 15 μm, from about 1 μm to about 10 μm, from about 1 μm to about 5 μm, from about 3 μm to about 15 μm, from about 3 μm to about 10 μm, from about 5 μm to about 10 μm, from about 10 μm to about 35 μm, from about 10 μm to about 20 μm, from about 15 μm to about 30 μm, from about 15 μm to about 25 μm, or from about 20 μm to about 35 μm. In certain embodiments, the diameter or thickness ratio of the core and the shell are in the range of 15:85 to 85:15, 25:75 to 75:25, 30:70 to 70:30, or 40:60 to 60:40. In certain embodiments, the volume or weight ratio of the core and the shell is 95:5, 90:10, 80:20, 70:30, 60:40, 50:50, 40:60, or 30:70.

[00201] The current collector acts to collect electrons generated by electrochemical reactions of the cathode active material or to supply electrons required for the electrochemical reactions. In some embodiments, the current collector can be in the form of a foil, sheet or film. In certain embodiments, the current collector is stainless steel, titanium, nickel, aluminum, copper, or alloys thereof or electrically-conductive resin. In certain embodiments, the current collector has a two-layered structure comprising an outer layer and an inner layer, wherein the outer layer comprises a conductive material and the inner layer comprises an insulating material or another conductive material; for example, aluminum mounted with a conductive resin layer or a polymeric insulating material coated with an aluminum film. In some embodiments, the current collector has a three-layered structure comprising an outer layer, a middle layer and an inner layer, wherein the outer and inner layers comprise a conductive material and the middle layer comprises an insulating material or another conductive material; for example, a plastic substrate coated with a metal film on both sides. In certain embodiments, each of the outer layer, middle layer and inner layer is independently stainless steel, titanium, nickel, aluminum, copper, or alloys thereof or electrically-conductive resin. In some embodiments, the insulating material is a polymeric material selected from the group consisting of polycarbonate, polyacrylate, polyacrylonitrile, polyester, polyamide, polystyrene, polyurethane, polyepoxy, poly(acrylonitrile butadiene styrene), polyimide, polyolefin, polyethylene, polypropylene, polyphenylene sulfide,

poly(vinyl ester), polyvinyl chloride, polyether, polyphenylene oxide, cellulose polymer and combinations thereof. In certain embodiments, the current collector has more than three layers. In some embodiments, the current collector is coated with a protective coating. In certain embodiments, the protective coating comprises a carbon-containing material. In some embodiments, the current collector is not coated with a protective coating.

[00202] The thickness of the current collector affects the volume it occupies within the battery, the amount of the electrode active material needed, and hence the capacity in the battery. In some embodiments, the current collector has a thickness from about 5 μm to about 30 μm. In certain embodiments, the current collector has a thickness from about 5 μm to about 20 μm, from about 5 μm to about 15 μm, from about 10 μm to about 30 μm, from about 10 μm to about 25 μm, or from about 10 μm to about 20 μm.

[00203] In some embodiments, the current collector has a thickness of less than 30 μm, less than 28 μm, less than 26 μm, less than 24 μm, less than 22 μm, less than 20 μm, less than 18 μm, less than 16 μm, less than 14 μm, less than 12 μm, less than 10 μm, less than 8 μm or less than 6 μm. In some embodiments, the current collector has a thickness of more than 5 μm, more than 7 μm, more than 10 μm, more than 12 μm, more than 14 μm, more than 16 μm, more than 18 μm, more than 20 μm, more than 22 μm, more than 24 μm, more than 26 μm or more than 28 μm.

[00204] The conductive agent is for enhancing the electrically-conducting property of an electrode. Any suitable material can act as the conductive agent. In some embodiments, the conductive agent is a carbonaceous material. Some non-limiting examples include carbon, carbon black, graphite, expanded graphite, graphene, graphene nanoplatelets, carbon fibers, carbon nano-fibers, graphitized carbon flake, carbon tubes, activated carbon, Super P, 0-dimensional KS6, 1-dimensional vapor grown carbon fibers (VGCF), mesoporous carbon and combinations thereof.

[00205] In addition, the cathode prepared using the binder composition in the present invention exhibits strong adhesion of the electrode layer to the current collector. It is important for the electrode layer to have good peeling strength to the current collector as this prevents delamination or separation of the electrode, which would greatly influence the mechanical stability of the electrodes and the cyclability of the battery. Therefore, the electrodes should have sufficient peeling strength to withstand the rigors of battery manufacture.

[00206] In some embodiments, the peeling strength between the current collector and the electrode layer is in the range from about 1.0 N/cm to about 8.0 N/cm, from about 1.0 N/cm to about 6.0 N/cm, from about 1.0 N/cm to about 5.0 N/cm, from about 1.0 N/cm to about 4.0 N/cm, from about 1.0 N/cm to about 3.0 N/cm, from about 1.0 N/cm to about 2.5 N/cm, from about 1.0 N/cm to about 2.0 N/cm, from about 1.2 N/cm to about 3.0 N/cm, from about 1.2

N/cm to about 2.5 N/cm, from about 1.2 N/cm to about 2.0 N/cm, from about 1.5 N/cm to about 3.0 N/cm, from about 1.5 N/cm to about 2.5 N/cm, from about 1.5 N/cm to about 2.0 N/cm from about 1.8 N/cm to about 3.0 N/cm, from about 1.8 N/cm to about 2.5 N/cm, from about 2.0 N/cm to about 6.0 N/cm, from about 2.0 N/cm to about 5.0 N/cm, from about 2.0 N/cm to about 3.0 N/cm, from about 2.0 N/cm to about 2.5 N/cm, from about 2.2 N/cm to about 3.0 N/cm, from about 2.5 N/cm to about 3.0 N/cm, from about 3.0 N/cm to about 8.0 N/cm, from about 3.0 N/cm to about 6.0 N/cm, or from about 4.0 N/cm to about 6.0 N/cm.

[00207] In some embodiments, the peeling strength between the current collector and the electrode layer is 1.0 N/cm or more, 1.2 N/cm or more, 1.5 N/cm or more, 2.0 N/cm or more, 2.2 N/cm or more, 2.5 N/cm or more, 3.0 N/cm or more, 3.5 N/cm or more, 4.5 N/cm or more, 5.0 N/cm or more, 5.5 N/cm or more, 6.0 N/cm or more, 6.5 N/cm or more, 7.0 N/cm or more or 7.5 N/cm or more. In some embodiments, the peeling strength between the current collector and the electrode layer is less than 8.0 N/cm, less than 7.5 N/cm, less than 7.0 N/cm, less than 6.5 N/cm, less than 6.0 N/cm, less than 5.5 N/cm, less than 5.0 N/cm, less than 4.5 N/cm, less than 4.0 N/cm, less than 3.5 N/cm, less than 3.0 N/cm, less than 2.8 N/cm, less than 2.5 N/cm, less than 2.2 N/cm, less than 2.0 N/cm, less than 1.8 N/cm, or less than 1.5 N/cm.

[00208] The extent of swelling of the binder composition from an uptake of electrolyte in the secondary battery provides insight into the crystallinity of the binder composition and how the binder composition interacts with the electrolyte. On one hand, a binder composition of high crystallinity exhibits low swelling behavior that can act as barriers for solvent ingress, provides shorter ion transport pathways, thus reduces internal resistance and more importantly alter the mechanical properties of the swollen polymer, which are critical to stable battery performance. On the other hand, a binder composition of low crystallinity has a higher amount of amorphous regions where higher amounts of electrolyte can penetrate into the binder composition to ensure good ion transport. A semi-crystalline binder composition disclosed herein benefits from both influence factors and therefore demonstrates exceptional electrochemical performance.

[00209] In some embodiments, the electrolyte swelling of the binder composition is from about 2% to about 4%, from about 2.1% to about 4%, from about 2.2% to about 4%, from about 2.3% to about 4%, from about 2.4% to about 4%, from about 2.5% to about 4%, from about 2.6% to about 4%, from about 2.7% to about 4%, from about 2.8% to about 4%, from about 2.9% to about 4%, from about 3% to about 4%, from about 3.1% to about 4%, from about 3.2% to about 4%, from about 3.3% to about 4%, from about 3.4% to about 4%, from about 3.5% to about 4%, from about 3% to about 3.9%, from about 3% to about 3.8%, from about 3% to about 3.7%, from about 3% to about 3.6%, from about 3% to about 3.5%, from about 2.5% to about 3.5%, from about 2.5% to about 3.4%, from about 2.5% to about 3.3%, from about 2.5% to about 3.2%, from about 2.5% to about 3.1%, from about 2.5% to about 3%, from about 2% to about 3%, from about 2% to about 2.9%, from about 2% to about 2.8%, from about 2% to about

2.7%, from about 2% to about 2.6%, from about 2% to about 2.5%, from about 2.2% to about 3.7% or from about 2.7% to about 3.3%.

[00210] In some embodiments, the electrolyte swelling of the binder composition is less than 4%, less than 3.9%, less than 3.8%, less than 3.7%, less than 3.6%, less than 3.5%, less than 3.4%, less than 3.3%, less than 3.2%, less than 3.1%, less than 3%, less than 2.9%, less than 2.8%, less than 2.7%, less than 2.6%, less than 2.5%, less than 2.4%, less than 2.3%, less than 2.2% or less than 2.1%. In some embodiments, the electrolyte swelling of the binder composition is more than 2%, more than 2.1%, more than 2.2%, more than 2.3%, more than 2.4%, more than 2.5%, more than 2.6%, more than 2.7%, more than 2.8%, more than 2.9%, more than 3%, more than 3.1%, more than 3.2%, more than 3.3%, more than 3.4%, more than 3.5%, more than 3.6%, more than 3.7%, more than 3.8% or more than 3.9%.

[00211] The method disclosed herein has the advantage that aqueous solvents can be used in the manufacturing process, which can save on processing time and equipment, as well as improve safety by eliminating the need to handle or recycle hazardous organic solvents. In addition, costs are reduced by simplifying the overall process. Therefore, this method is especially suited for industrial processes because of its low cost and ease of handling.

[00212] The following examples are presented to exemplify embodiments of the invention but are not intended to limit the invention to the specific embodiments set forth. Unless indicated to the contrary, all parts and percentages are by weight. All numerical values are approximate. When numerical ranges are given, it should be understood that embodiments outside the stated ranges may still fall within the scope of the invention. Specific details described in each example should not be construed as necessary features of the invention.

EXAMPLES

[00213] The pH values of the binder composition were measured by an electrode-type pH meter (ION 2700, Eutech Instruments).

[00214] The viscosities of the binder composition were measured using a rotational viscosity meter (NDJ-5S, Shanghai JT Electronic Technology Co. Ltd., China) at 25 ºC.

[00215] The adhesive strengths of the dried binder composition layers were measured by a tensile testing machine (DZ-106A, obtained from Dongguan Zonhow Test Equipment Co. Ltd., China). This test measures the average force required to peel a binder composition layer from the current collector at 180° angle in Newtons. The mean roughness depth (R_z) of the current collector is 2 µm. The binder composition was coated on the current collector and dried to obtain a binder composition layer of thickness 10 µm to 12 µm. The coated current collector was then placed in an environment of constant temperature of 25 ºC and humidity of 50% to 60% for 30 minutes. A strip of adhesion tape (3M; US; model no. 810) with a width of 18 mm

and a length of 20 mm was attached onto the surface of the binder composition layer. The binder composition strip was clipped onto the testing machine and the tape was folded back on itself at 180 degrees, and placed in a moveable jaw and pulled at room temperature and a peel rate of 300 mm per minute. The maximum stripping force measured was taken as the adhesive strength. Measurements were repeated three times to find the average value.

[00216] The electrolyte swelling of the binder composition measures the extent of mass change of binder composition before and after electrolyte soaking. Test samples of dried binder composition strips with a length of 50 mm to 60 mm and a width of 1 mm were prepared. The dried binder composition strips were further dried at 80 ºC for 1 to 2 hours to completely remove the moisture in the strips. The weights of the dried binder composition strips were measured and the strips were placed in a sealed container with electrolyte after cooling. The binder composition strips were soaked in the electrolyte at 25 ºC for 3 days. After removing the binder composition strips from the electrolyte-containing container, the electrolyte on the surface of the strips were absorbed with oil-absorbing papers. The weights of the soaked binder composition strips were measured. The electrolyte swelling is the ratio of the weight change of the strip before and after electrolyte soaking to the weight of the strip before electrolyte soaking. Measurements were repeated three times to find the average value.

[00217] The solid content of the binder composition measures the extent of mass change of binder composition before and after drying. Approximately 1 g of a binder composition was weighed in a weighing bottle and dried at 110 ± 5℃ and -0.09 MPa for more than 5 hours by a vacuum dryer. The binder composition was cooled in a desiccator for about 15 minutes and then measured in terms of mass. The difference in mass of the binder composition before and after the drying was determined, and the solid content (%) of the binder composition was calculated according to the following formula:

[00218] $$Solid\ content\ (\%) = \frac{Mass\ of\ binder\ composition\ after\ drying}{Mass\ of\ binder\ composition\ before\ drying} \times 100\%$$

[00219] The weight average molecular weight and number average molecular weight of the binder composition were measured by gel permeation chromatography (GPC). The binder composition was firstly dissolved in dimethylformamide at room temperature. Once dissolution of the binder composition was completed, the solution was gently filtered through a 0.45 μm filter to prepare a measurement sample. A standard polystyrene was used to prepare a calibration curve such that the weight average molecular weight and the number average molecular weight were calculated as standard substance equivalent values. The distribution of molecular weights in the binder composition is described by the Polydispersity Index (PDI) that is the ratio of the weight average molecular weight to the number average molecular weight. The obtained measurement sample was analyzed under the following conditions:

[00220] Column: Agilent PLgel 5um MIXED-C column

[00221] Eluent: Dimethylformamide

[00222] Flow rate: 1ml/min

[00223] Weight of sample: 2mg

[00224] Detector: Waters 2414 Refractive Index (RI) Detector

[00225] Detection temperature: 35℃

[00226] Standard substance: Polystyrene

Example 1

A) Preparation of binder composition

[00227] 5.13 g of lithium hydroxide was dissolved in 3.85 g of DI water. Thereafter, 8.98 g of lithium hydroxide solution was added into a 500mL round-bottom flask containing 289.17 g of distilled water. The mixture was stirred at 200 rpm for 30 mins to obtain a first suspension.

[00228] Further, 19.15 g of acrylic acid (AA) was added into the first suspension. The mixture was further stirred at 200 rpm for 30 mins to obtain a second suspension.

[00229] 15.98 g of acrylamide (AM) was dissolved in 51.67 g of DI water. Thereafter, 67.65 g of AM solution was added into the second suspension. The mixture was further stirred at 200 rpm for 30 mins to obtain a third suspension.

[00230] 77.53 g of acrylonitrile (AN) was then added into the third suspension. The fourth suspension was obtained by stirring the mixture at 200 rpm for 40 mins.

[00231] The fourth suspension was heated up to 60 ℃ and stirred at 60 rpm for 45 mins. 0.23 g of water-soluble free radical initiator (ammonium persulfate, APS; obtained from Aladdin Industries Corporation, China) was dissolved in 82.68 g of DI water and 0.04 g of reducing agent (sodium bisulfite; obtained from Tianjin Damao Chemical Reagent Factory, China) was dissolved in 17.22 g of DI water. 17.26 g of sodium bisulfite solution was added into the fourth suspension and the mixture was stirred for 10 minutes. 82.91 g of APS solution was added into the mixture dropwise for 3 h to form a fifth suspension. The fifth suspension was further stirred at 200 rpm for 20 h at 65 ℃.

[00232] After the complete reaction, the temperature of the fifth suspension was lowered to 40 ºC and 0.69 g of lithium hydroxide (dissolved in 116.64 g of DI water) was added into the fifth suspension to adjust pH to 7.42 to form the sixth suspension. The temperature of the sixth suspension was lowered to 30 ºC and the binder composition was furnished by filtration using 200 mesh filter paper. The solid content of the binder composition was 14.85 wt.%. The weight average molecular weight, the number average molecular weight and the polydispersity index of

the binder composition were 163,282 g/mol, 71,877 g/mol and 2.27 respectively. The components of the binder composition of Example 1 and their respective proportions are shown in Table 1 below. The pH, solid content, viscosity, adhesive strength and electrolyte swelling of the binder composition of Example 1 were measured and are shown in Table 2 below.

B) Preparation of positive electrode

[00233] A first mixture was prepared by dispersing 0.9 g of conductive agent (SuperP; obtained from Timcal Ltd, Bodio, Switzerland) and 6 g of binder composition (14.85 wt.% solid content) in 7.4 g of deionized water while stirring with an overhead stirrer (R20, IKA). After the addition, the first mixture was further stirred for about 30 minutes at 25 °C at a speed of 1,200 rpm.

[00234] Thereafter, a second mixture was prepared by adding 28.2 g of NMC622 (obtained from Shandong Tianjiao New Energy Co.,Ltd, China) in the first mixture at 25 °C while stirring with an overhead stirrer. Then, the second mixture was degassed under a pressure of about 10 kPa for 1 hour. Then, the second mixture was further stirred for about 60 minutes at 25 °C at a speed of 1,200 rpm to form a homogenized slurry.

[00235] The homogenized slurry was coated onto one side of an aluminum foil having a thickness of 14 μm as a current collector using a doctor blade coater. The coated slurry film on the aluminum foil was dried at about 85 ºC for 120 minutes by a hot air dryer (DHG 10H, Huyue Equipment Co., Ltd., China) to form a cathode electrode layer. The electrode was then pressed to decrease the thickness of the cathode electrode layer to 27 μm and the surface density was 5.2 mg/cm^2.

C) Preparation of negative electrode

[00236] A negative electrode slurry was prepared by mixing 90 wt.% of hard carbon (BTR New Energy Materials Inc., Shenzhen, Guangdong, China) with 1.5 wt.% carboxymethyl cellulose (CMC, BSH-12, DKS Co. Ltd., Japan) and 3.5 wt.% SBR (AL-2001, NIPPON A&L INC., Japan) as a binder, and 5 wt.% carbon black as a conductive agent in deionized water. The solid content of the anode slurry was 50 wt.%. The slurry was coated onto one side of a copper foil having a thickness of 8 μm using a doctor blade coater. The coated film on the copper foil was dried at about 85 °C for 120 minutes by a hot air dryer to obtain a negative electrode. The electrode was then pressed to decrease the thickness of the coating to 18 μm.

D) Assembling of coin cell

[00237] CR2032 coin-type Li cells were assembled in an argon-filled glove box. The coated cathode and anode sheets were cut into disc-form positive and negative electrodes, which were then assembled into an electrode assembly by stacking the cathode and anode electrode plates alternatively and then packaged in a case made of stainless steel of the CR2032 type. The

cathode and anode electrode plates were kept apart by separators. The separator was a ceramic coated microporous membrane made of nonwoven fabric (MPM, Japan), which had a thickness of about 25 μm. The electrode assembly was then dried in a box-type resistance oven under vacuum (DZF-6020, obtained from Shenzhen Kejing Star Technology Co. Ltd., China) at 105 °C for about 16 hours.

[00238] An electrolyte was then injected into the case holding the packed electrodes under a high-purity argon atmosphere with a moisture and oxygen content of less than 3 ppm respectively. The electrolyte was a solution of $LiPF_6$ (1 M) in a mixture of ethylene carbonate (EC), ethyl methyl carbonate (EMC) and dimethyl carbonate (DMC) at a volume ratio of 1:1:1. After electrolyte filling, the coin cell was vacuum sealed and then mechanically pressed using a punch tooling with a standard circular shape.

E) Electrochemical measurements

[00239] The coin cells were analyzed in a constant current mode using a multi-channel battery tester (BTS-4008-5V10mA, obtained from Neware Electronics Co. Ltd, China). After 1 cycle at C/20 was completed, they were charged and discharged at a rate of C/2. The charging/discharging cycling tests of the cells were performed between 3.0 and 4.3 V at a current density of C/2 at 25 °C to obtain the discharge capacity. The electrochemical performance of the coin cell of Example 1 was measured and is shown in Table 2 below.

[00240] Example 2: A binder composition was prepared in the same manner as in Example 1, except that 11.27 g of AA was added in the preparation of the second suspension, 20.27 g of AM was added in the preparation of the third suspension and 81.12 g of AN was added in the preparation of the fourth suspension.

[00241] Example 3: A binder composition was prepared in the same manner as in Example 1, except that 31.54 g of AA was added in the preparation of the second suspension, 13.52 g of AM was added in the preparation of the third suspension, 67.60 g of AN was added in the preparation of the fourth suspension and 5.62 g of lithium hydroxide (dissolved in 116.64 g of DI water) was added in the preparation of the sixth suspension.

[00242] Example 4: A binder composition was prepared in the same manner as in Example 1, except that 14.74 g of AA and 4.41 g of methacrylic acid (MAA) were added in the preparation of the second suspension. The weight average molecular weight, the number average molecular weight and the polydispersity index of the binder composition were 159,836 g/mol, 70,980 g/mol and 2.25 respectively.

Preparation of binder composition of Example 5

[00243] 4.99 g of lithium hydroxide was dissolved in 3.75 g of DI water. Thereafter, 8.74 g of lithium hydroxide solution was added into a 500mL round-bottom flask containing 281.29 g

of distilled water. The mixture was stirred at 200 rpm for 30 mins to obtain a first suspension.

[00244] Further, 15.62 g of AA and 4.67 g of MAA were added into the first suspension. The mixture was further stirred at 200 rpm for 30 mins to obtain a second suspension.

[00245] 15.55 g of AM was dissolved in 50.27 g of DI water. Thereafter, 65.82 g of AM solution was added into the second suspension. The mixture was further stirred at 200 rpm for 30 mins to obtain a third suspension.

[00246] 73.77 g of AN was then added into the third suspension. The fourth suspension was obtained by stirring the mixture at 200 rpm for 40 mins.

[00247] The fourth suspension was heated up to 60 ºC and stirred at 60 rpm for 45 mins. 0.29 g of water-soluble free radical initiator (ammonium persulfate, APS; obtained from Aladdin Industries Corporation, China) was dissolved in 80.43 g of DI water and 0.05 g of reducing agent (sodium bisulfite; obtained from Tianjin Damao Chemical Reagent Factory, China) was dissolved in 16.76 g of DI water. 16.81 g of sodium bisulfite solution was added into the fourth suspension and the mixture was stirred for 10 minutes. 80.72 g of APS solution was added into the mixture dropwise for 3 h to form a fifth suspension. The fifth suspension was further stirred at 200 rpm for 20 h at 65 ºC.

[00248] After the complete reaction, the temperature of the fifth suspension was lowered to 40 ºC and 1.15 g of lithium hydroxide (dissolved in 101.39 g of DI water) was added into the fifth suspension to adjust pH to 7.90 to form the sixth suspension. The temperature of the sixth suspension was lowered to 30 ºC and the binder composition was furnished by filtration using 200 mesh filter paper. The solid content of the binder composition was 16.54 wt.%.

Preparation of binder composition of Example 6

[00249] 6.37 g of lithium hydroxide was dissolved in 4.79 g of DI water. Thereafter, 11.16 g of lithium hydroxide solution was added into a 500mL round-bottom flask containing 271.80 g of distilled water. The mixture was stirred at 200 rpm for 30 mins to obtain a first suspension.

[00250] Further, 20.45 g of AA and 6.10 g of MAA were added into the first suspension. The mixture was further stirred at 200 rpm for 30 mins to obtain a second suspension.

[00251] 10.93 g of AM was dissolved in 48.57 g of DI water. Thereafter, 59.50 g of AM solution was added into the second suspension. The mixture was further stirred at 200 rpm for 30 mins to obtain a third suspension.

[00252] 68.41 g of AN was then added into the third suspension. The fourth suspension was obtained by stirring the mixture at 200 rpm for 40 mins.

[00253] The fourth suspension was heated up to 60 ºC and stirred at 60 rpm for 45 mins.

0.28 g of water-soluble free radical initiator (ammonium persulfate, APS; obtained from Aladdin Industries Corporation, China) was dissolved in 77.71 g of DI water and 0.05 g of reducing agent (sodium bisulfite; obtained from Tianjin Damao Chemical Reagent Factory, China) was dissolved in 16.19 g of DI water. 16.24 g of sodium bisulfite solution was added into the fourth suspension and the mixture was stirred for 10 minutes. 77.99 g of APS solution was added into the mixture dropwise for 3 h to form a fifth suspension. The fifth suspension was further stirred at 200 rpm for 20 h at 65 ºC.

[00254] After the complete reaction, the temperature of the fifth suspension was lowered to 40 ºC and 1.57 g of lithium hydroxide (dissolved in 74.14 g of DI water) was added into the fifth suspension to adjust pH to 7.80 to form the sixth suspension. The sixth suspension was lowered to 30 ºC and the binder composition was furnished by filtration using 200 mesh filter paper. The solid content of the binder composition was 17.47 wt.%.

[00255] Example 7: A binder composition was prepared in the same manner as in Example 6, except that 15.80 g of AM was added in the preparation of the third suspension, 63.53 g of AN was added in the preparation of the fourth suspension and 0.46 g of lithium hydroxide (dissolved in 198 g of DI water) was added in the preparation of the sixth suspension.

[00256] Example 8: A binder composition was prepared in the same manner as in Example 6, except that 14.21 g of AM was added in the preparation of the third suspension, 65.12 g of AN was added in the preparation of the fourth suspension and 0.91 g of lithium hydroxide (dissolved in 197.60 g of DI water) was added in the preparation of the sixth suspension.

Preparation of binder composition of Example 9

[00257] 6.37 g of lithium hydroxide was dissolved in 4.79 g of DI water. Thereafter, 11.16 g of lithium hydroxide solution was added into a 500mL round-bottom flask containing 181.80 g of distilled water. The mixture was stirred at 200 rpm for 30 mins to obtain a first suspension.

[00258] Further, 20.45 g of AA and 6.10 g of MAA were added into the first suspension. The mixture was further stirred at 200 rpm for 30 mins to obtain a second suspension.

[00259] 12.62 g of AM was dissolved in 48.57 g of DI water. Thereafter, 61.19 g of AM solution was added into the second suspension. The mixture was further stirred at 200 rpm for 30 mins to obtain a third suspension.

[00260] 66.71 g of AN was then added into the third suspension. The fourth suspension was obtained by stirring the mixture at 200 rpm for 40 mins.

[00261] The fourth suspension was heated up to 60 ºC and stirred at 60 rpm for 45 mins. 0.11 g of water-soluble free radical initiator (ammonium persulfate, APS; obtained from Aladdin

Industries Corporation, China) was dissolved in 10.00 g of DI water and 0.02 g of reducing agent (sodium bisulfite; obtained from Tianjin Damao Chemical Reagent Factory, China) was dissolved in 16.19 g of DI water. 16.21 g of sodium bisulfite solution was added into the fourth suspension and the mixture was stirred for 10 minutes. 10.11 g of APS solution was added into the mixture dropwise for 3 h to form a fifth suspension. The fifth suspension was further stirred at 200 rpm for 20 h at 65 ºC.

[00262] After the complete reaction, the temperature of the fifth suspension was lowered to 40 ºC and 1.66 g of lithium hydroxide (dissolved in 314.66 g of DI water) was added into the fifth suspension to adjust pH to 7.24 to form the sixth suspension. The temperature of the sixth suspension was lowered to 30 ºC and the binder composition was furnished by filtration using 200 mesh filter paper. The solid content of the binder composition was 14.76 wt.%.

[00263] Example 10: A binder composition was prepared in the same manner as in Example 4, except that 7.89 g of AA and 5.63 g of MAA were added in the preparation of the second suspension, 16.90 g of AM was added in the preparation of the third suspension and 82.24 g of AN was added in the preparation of the fourth suspension.

[00264] Example 11: A binder composition was prepared in the same manner as in Example 4, except that 14.65 g of AA and 6.76 g of MAA were added in the preparation of the second suspension, 9.01 g of AM was added in the preparation of the third suspension and 82.24 g of AN was added in the preparation of the fourth suspension.

[00265] Example 12: A binder composition was prepared in the same manner as in Example 4, except that 30.42 g of AA and 3.38 g of MAA were added in the preparation of the second suspension, 11.27 g of AM was added in the preparation of the third suspension, 67.60 g of AN was added in the preparation of the fourth suspension and 6.12 g of lithium hydroxide (dissolved in 116.64 g of DI water) was added in the preparation of the sixth suspension.

[00266] Example 13: A binder composition was prepared in the same manner as in Example 4, except that 5.63 g of AA and 5.63 g of MAA were added in the preparation of the second suspension, 22.53 g of AM was added in the preparation of the third suspension and 78.87 g of AN was added in the preparation of the fourth suspension.

[00267] Example 14: A binder composition was prepared in the same manner as in Example 4, except that 27.04 g of AA and 5.63 g of MAA were added in the preparation of the second suspension, 9.01 g of AM was added in the preparation of the third suspension, 70.98 g of AN was added in the preparation of the fourth suspension and 6.24 g of lithium hydroxide (dissolved in 116.64 g of DI water) was added in the preparation of the sixth suspension.

[00268] Example 15: A binder composition was prepared in the same manner as in Example 4, except that 4.41 g of MAA was replaced with 2-ethylacrylic acid of the same weight

in the preparation of the second suspension.

[00269] Example 16: A binder composition was prepared in the same manner as in Example 4, except that 4.41 g of MAA was replaced with crotonic acid of the same weight in the preparation of the second suspension.

[00270] Example 17: A binder composition was prepared in the same manner as in Example 5, except that 7.93 g of lithium hydroxide (dissolved in 101.39 g of DI water) was added in the preparation of the sixth suspension.

[00271] Example 18: A binder composition was prepared in the same manner as in Example 6, except that 10.60 g of lithium hydroxide (dissolved in 74.14 g of DI water) was added in the preparation of the sixth suspension.

Preparation of binder composition of Examples 19-21

[00272] The binder compositions of Examples 19-21 were prepared in the same manner as in Example 4.

[00273] Example 22: A binder composition was prepared in the same manner as in Example 4, except that 0.29 g of APS was dissolved in 82.68 g of DI water and 0.05 g of sodium bisulfite was dissolved in 17.22 g of DI water so that 82.97 g of APS solution and 17.27 g of sodium bisulfite solution were added in the preparation of the fifth suspension. The weight average molecular weight, the number average molecular weight and the polydispersity index of the binder composition were 105,780 g/mol, 29,845 g/mol and 3.54 respectively.

[00274] Example 23: A binder composition was prepared in the same manner as in Example 4, except that 0.11 g of APS was dissolved in 82.68 g of DI water and 0.02 g of sodium bisulfite was dissolved in 17.22 g of DI water so that 82.79 g of APS solution and 17.24 g of sodium bisulfite solution were added in the preparation of the fifth suspension. The weight average molecular weight, the number average molecular weight and the polydispersity index of the binder composition were 193,226 g/mol, 89,641 g/mol and 2.16 respectively.

[00275] Example 24: A binder composition was prepared in the same manner as in Example 1, except that 0.29 g of APS was dissolved in 82.68 g of DI water and 0.05 g of sodium bisulfite was dissolved in 17.22 g of DI water so that 82.97 g of APS solution and 17.27 g of sodium bisulfite solution were added in the preparation of the fifth suspension. The weight average molecular weight, the number average molecular weight and the polydispersity index of the binder composition were 118,528 g/mol, 30,523 g/mol and 3.88 respectively.

[00276] Example 25: A binder composition was prepared in the same manner as in Example 1, except that 0.11 g of APS was dissolved in 82.68 g of DI water and 0.02 g of sodium bisulfite was dissolved in 17.22 g of DI water so that 82.79 g of APS solution and 17.24 g of sodium bisulfite solution were added in the preparation of the fifth suspension. The weight

average molecular weight, the number average molecular weight and the polydispersity index of the binder composition were 186,744 g/mol, 92,140 g/mol and 2.03 respectively.

Comparative Example 1

[00277] A binder composition was prepared in the same manner as in Example 1, except that 1.10 g of lithium hydroxide (dissolved in 3.85 g of DI water) was added in the preparation of the first suspension, 7.21 g of AA was added in the preparation of the second suspension, 22.75 g of AM was added in the preparation of the third suspension, 83.83 g of AN was added in the preparation of the fourth suspension and 1.5 g of lithium hydroxide (dissolved in 116.64 g of DI water) was added in the preparation of the sixth suspension.

Comparative Example 2

[00278] A binder composition was prepared in the same manner as in Example 1, except that 37.47 g of AA was added in the preparation of the second suspension, 7.11 g of AM was added in the preparation of the third suspension, 73.22 g of AN was added in the preparation of the fourth suspension and 7.34 g of lithium hydroxide (dissolved in 116.64 g of DI water) was added in the preparation of the sixth suspension.

Comparative Example 3

[00279] A binder composition was prepared in the same manner as in Example 4, except that 24.50 g of AA and 6.88 g of MAA were added in the preparation of the second suspension, 22.75 g of AM was added in the preparation of the third suspension, 66.86 g of AN was added in the preparation of the fourth suspension and 4.95 g of lithium hydroxide (dissolved in 116.64 g of DI water) was added in the preparation of the sixth suspension.

Comparative Example 4

[00280] A binder composition was prepared in the same manner as in Example 4, except that 10.09 g of AA and 5.16 g of MAA were added in the preparation of the second suspension, 7.11 g of AM was added in the preparation of the third suspension and 90.20 g of AN was added in the preparation of the fourth suspension.

Comparative Example 5

[00281] A binder composition was prepared in the same manner as in Example 4, except that 0.8 g of lithium hydroxide (dissolved in 3.85 g of DI water) was added in the preparation of the first suspension, 4.32 g of AA and 0.86 g of MAA were added in the preparation of the second suspension, 12.79 g of AM was added in the preparation of the third suspension, 92.86 g of AN was added in the preparation of the fourth suspension and 1.40 g of lithium hydroxide (dissolved in 116.64 g of DI water) was added in the preparation of the sixth suspension.

Comparative Example 6

[00282] A binder composition was prepared in the same manner as in Example 4, except that 28.10 g of AA and 10.33 g of MAA were added in the preparation of the second suspension, 7.11 g of AM was added in the preparation of the third suspension, 73.75 g of AN was added in the preparation of the fourth suspension and 7.38 g of lithium hydroxide (dissolved in 116.64 g of DI water) was added in the preparation of the sixth suspension.

Comparative Example 7

[00283] A binder composition was prepared in the same manner as in Example 4, except that 21.62 g of AA and 6.88 g of MAA were added in the preparation of the second suspension, 4.26 g of AM was added in the preparation of the third suspension, 82.77 g of AN was added in the preparation of the fourth suspension and 6.50 g of lithium hydroxide (dissolved in 116.64 g of DI water) was added in the preparation of the sixth suspension.

Comparative Example 8

[00284] A binder composition was prepared in the same manner as in Example 4, except that 5.95 g of AA and 5.96 g of MAA were added in the preparation of the second suspension, 33.99 g of AM was added in the preparation of the third suspension and 72.69 g of AN was added in the preparation of the fourth suspension.

Comparative Example 9

[00285] A binder composition was prepared in the same manner as in Example 4, except that 36.03 g of AA and 17.21 g of MAA were added in the preparation of the second suspension, 17.06 g of AM was added in the preparation of the third suspension, 56.24 g of AN was added in the preparation of the fourth suspension and 9.69 g of lithium hydroxide (dissolved in 116.64 g of DI water) was added in the preparation of the sixth suspension.

Comparative Example 10

[00286] A binder composition was prepared in the same manner as in Example 4, except that 17.29 g of AA, 5.16 g of MAA and 8.51 g of methyl acrylate (MA) were added in the preparation of the second suspension, 14.22 g of AM was added in the preparation of the third suspension and 74.28 g of AN was added in the preparation of the fourth suspension.

Preparation of positive electrode of Examples 2-18, 22-25 and Comparative Examples 1-10

[00287] The positive electrodes of Examples 2-18, 22-25 and Comparative Examples 1-10 were prepared in the same manner as in Example 1.

Preparation of positive electrode of Example 19

[00288] The positive electrode of Example 19 was prepared in the same manner as in Example 1, except that the 28.2 g of NMC622 was replaced with NMC532 (obtained from

Tianjin Bamo Technology Co., Ltd., China) of the same weight.

Preparation of positive electrode of Example 20

[00289] The positive electrode of Example 20 was prepared in the same manner as in Example 1, except that the 28.2 g of NMC622 was replaced with $LiCoO_2$ (obtained from Tianjin Bamo Technology Co., Ltd., China) of the same weight.

Preparation of positive electrode of Example 21

[00290] The positive electrode of Example 21 was prepared in the same manner as in Example 1, except that the 28.2 g of NMC622 was replaced with $LiFePO_4$ (obtained from Xiamen Tungsten Industry Co., Ltd., China) of the same weight.

Preparation of negative electrode of Examples 2-25 and Comparative Examples 1-10

[00291] The negative electrodes of Examples 2-25 and Comparative Examples 1-10 were prepared in the same manner as in Example 1.

Assembling of coin cells of Examples 2-25 and Comparative Examples 1-10

[00292] The coin cells of Examples 2-25 and Comparative Examples 1-10 were assembled in the same manner as in Example 1.

Electrochemical measurements of Examples 2-25 and Comparative Examples 1-10

[00293] The electrochemical performance of the coin cells of Examples 2-25 and Comparative Examples 1-10 were measured in the same manner as in Example 1 and the test results are shown in Table 2 below. Only the capacity retention of the coin cells after 100 cycles of Examples 1-11, 13, 15-16, 19-21 and Comparative Examples 1-10 were measured and the test results are shown in Table 2 below.

Table 1

	Monomers added in the preparation of binder composition					
	AN*	AA*	AM*	Other carboxylic acid group-containing monomer		MA*
	Proportion of monomer (wt%)			Type	Proportion of monomer (wt%)	
Example 1	68.82	17.00	14.18	-	0.00	0.00
Example 2	72.00	10.00	18.00	-	0.00	0.00
Example 3	60.00	28.00	12.00	-	0.00	0.00
Example 4	68.82	13.08	14.18	MAA*	3.92	0.00
Example 5	67.30	14.25	14.19	MAA	4.26	0.00
Example 6	64.60	19.32	10.32	MAA	5.76	0.00
Example 7	60.00	19.32	14.92	MAA	5.76	0.00
Example 8	61.50	19.32	13.42	MAA	5.76	0.00
Example 9	63.01	19.31	11.92	MAA	5.76	0.00
Example 10	73.00	7.00	15.00	MAA	5.00	0.00
Example 11	73.00	13.00	8.00	MAA	6.00	0.00
Example 12	60.00	27.00	10.00	MAA	3.00	0.00
Example 13	70.00	5.00	20.00	MAA	5.00	0.00
Example 14	63.00	24.00	8.00	MAA	5.00	0.00
Example 15	68.82	13.08	14.18	2-ethylacrylic acid	3.92	0.00
Example 16	68.82	13.08	14.18	crotonic acid	3.92	0.00
Example 17	67.30	14.25	14.19	MAA	4.26	0.00
Example 18	64.60	19.32	10.32	MAA	5.76	0.00
Example 19	68.82	13.08	14.18	MAA	3.92	0.00
Example 20	68.82	13.08	14.18	MAA	3.92	0.00
Example 21	68.82	13.08	14.18	MAA	3.92	0.00
Example 22	68.82	13.08	14.18	MAA	3.92	0.00
Example 23	68.82	13.08	14.18	MAA	3.92	0.00

Example 24	68.82	17.00	14.18	-	0.00	0.00
Example 25	68.82	17.00	14.18	-	0.00	0.00
Comparative Example 1	73.68	6.33	19.99	-	0.00	0.00
Comparative Example 2	62.16	31.81	6.03	-	0.00	0.00
Comparative Example 3	55.26	20.25	18.80	MAA	5.69	0.00
Comparative Example 4	80.14	8.96	6.31	MAA	4.59	0.00
Comparative Example 5	83.78	3.90	11.54	MAA	0.78	0.00
Comparative Example 6	61.83	23.56	5.96	MAA	8.66	0.00
Comparative Example 7	71.64	18.71	3.69	MAA	5.96	0.00
Comparative Example 8	61.30	5.02	28.66	MAA	5.02	0.00
Comparative Example 9	44.45	28.47	13.48	MAA	13.60	0.00
Comparative Example 10	62.18	14.48	11.90	MAA	4.32	7.12

*AN refers to acrylonitrile, AA refers to acrylic acid, AM refers to acrylamide, MAA refers to methacrylic acid and MA refers to methyl acrylate.

Table 2

	Physical properties of binder composition			Electrolyte swelling (%)	Adhesive strength (N/cm)	0.5C Initial discharging capacity (mAh/g)	Capacity retention after 50 cycles (%)	Capacity retention after 100 cycles (%)
	pH	Solid content (%)	Viscosity (mPa·s)					
Example 1	7.42	14.85	17,380	3.40	3.73	128	92.58	88.13
Example 2	7.61	15.62	12,920	3.28	3.66	129	92.88	88.24
Example 3	7.44	14.88	36,410	3.23	3.41	131	93.42	87.96
Example 4	7.56	15.47	16,570	3.21	3.89	132	94.01	90.54
Example 5	7.90	16.54	18,250	3.44	3.75	133	93.94	89.92
Example 6	7.80	17.47	26,000	2.49	3.51	130	94.61	91.02
Example 7	7.65	14.58	43,000	3.72	2.93	136	95.02	90.67
Example 8	7.77	14.71	35,000	2.96	3.02	128	93.99	89.44
Example 9	7.24	14.76	19,420	2.40	3.39	135	93.57	88.79
Example 10	7.06	13.74	14,030	2.89	3.74	141	94.77	84.20

Example 11	7.31	15.64	19,280	3.52	3.88	129	95.92	86.57
Example 12	7.51	17.93	37,400	2.80	3.63	130	93.72	-
Example 13	7.48	13.62	11,360	3.30	3.49	134	95.23	84.83
Example 14	7.71	15.02	36,950	3.01	3.20	139	93.75	-
Example 15	7.38	15.52	15,830	2.97	3.31	136	93.94	90.01
Example 16	7.41	14.85	18,400	3.46	3.26	134	94.33	90.48
Example 17	8.43	16.33	17,970	3.38	3.79	133	94.02	-
Example 18	8.51	17.54	25,800	2.53	3.60	131	94.53	-
Example 19	7.55	16.78	21,530	3.24	3.90	128	93.60	88.65
Example 20	7.64	17.22	16,480	3.21	3.83	164	93.42	89.86
Example 21	7.57	13.25	17,540	3.27	3.72	152	93.37	89.45
Example 22	7.30	14.56	16,560	3.55	3.62	134	94.04	-
Example 23	7.42	15.77	18,420	3.27	2.85	130	93.95	-
Example 24	7.17	13.98	14,840	3.50	3.41	133	93.88	-
Example 25	7.63	16.82	19,600	3.24	3.68	133	93.93	-
Comparative Example 1	7.45	14.93	8,100	2.38	1.22	104	84.20	75.79
Comparative Example 2	7.36	16.41	52,300	6.36	1.62	111	83.84	75.88
Comparative Example 3	7.62	12.70	40,270	2.62	1.57	109	86.73	78.23
Comparative Example 4	7.41	16.84	13,830	5.50	2.33	112	85.77	78.46
Comparative Example 5	7.11	13.02	6,210	2.82	2.56	106	85.31	77.05
Comparative Example 6	7.47	15.12	62,770	5.39	1.12	109	85.94	78.14
Comparative Example 7	7.98	15.87	35,660	6.13	1.21	110	88.35	80.41
Comparative Example 8	7.65	12.09	14,060	1.12	1.35	110	89.25	80.04
Comparative Example 9	7.53	17.61	74,530	2.55	1.22	106	85.04	77.62
Comparative Example 10	7.62	17.89	22,790	2.62	3.34	105	84.32	74.95

[00294] While the invention has been described with respect to a limited number of embodiments, the specific features of one embodiment should not be attributed to other embodiments of the invention. In some embodiments, the methods may include numerous steps not mentioned herein. In other embodiments, the methods do not include, or are substantially free of, any steps not enumerated herein. Variations and modifications from the described embodiments exist. The appended claims intend to cover all those modifications and variations as falling within the scope of the invention.

CLAIMS

1. A binder composition for a secondary battery electrode comprising a copolymer and a dispersion medium, wherein the copolymer comprises a structural unit (a) derived from a carboxylic acid group-containing monomer, a structural unit (b) derived from an amide group-containing monomer and a structural unit (c) derived from a nitrile group-containing monomer; wherein the proportion of structural unit (a) derived from a carboxylic acid group-containing monomer in the copolymer is from 7% to 25% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition; wherein the proportion of structural unit (b) derived from an amide group-containing monomer in the copolymer is from 4% to 17% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition; wherein the proportion of structural unit (c) derived from a nitrile group-containing monomer in the copolymer is from 65% to 80% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition; and wherein the binder composition is free of a structural unit derived from an ester group-containing monomer.

2. The binder composition according to claim 1, wherein the carboxylic acid group-containing monomer is selected from the group consisting of acrylic acid, methacrylic acid, crotonic acid, 2-butyl crotonic acid, cinnamic acid, maleic acid, maleic anhydride, fumaric acid, itaconic acid, itaconic anhydride, tetraconic acid, 2-ethylacrylic acid, isocrotonic acid, cis-2-pentenoic acid, trans-2-pentenoic acid, angelic acid, tiglic acid, 3,3-dimethyl acrylic acid, 3-propyl acrylic acid, trans-2-methyl-3-ethyl acrylic acid, cis-2-methyl-3-ethyl acrylic acid, 3-isopropyl acrylic acid, trans-3-methyl-3-ethyl acrylic acid, cis-3-methyl-3-ethyl acrylic acid, 2-isopropyl acrylic acid, trimethyl acrylic acid, 2-methyl-3,3-diethyl acrylic acid, 3-butyl acrylic acid, 2-butyl acrylic acid, 2-pentyl acrylic acid, 2-methyl-2-hexenoic acid, trans-3-methyl-2-hexenoic acid, 3-methyl-3-propyl acrylic acid, 2-ethyl-3-propyl acrylic acid, 2,3-diethyl acrylic acid, 3,3-diethyl acrylic acid, 3-methyl-3-hexyl acrylic acid, 3-methyl-3-tert-butyl acrylic acid, 2-methyl-3-pentyl acrylic acid, 3-methyl-3-pentyl acrylic acid, 4-methyl-2-hexenoic acid, 4-ethyl-2-hexenoic acid, 3-methyl-2-ethyl-2-hexenoic acid, 3-tert-butyl acrylic acid, 2,3-dimethyl-3-ethyl acrylic acid, 3,3-dimethyl-2-ethyl acrylic acid, 3-methyl-3-isopropyl acrylic acid, 2-methyl-3-isopropyl acrylic acid, trans-2-octenoic acid, cis-2-octenoic acid, trans-2-decenoic acid, α-acetoxyacrylic acid, β-trans-aryloxyacrylic acid, α-chloro-β-E-methoxyacrylic acid, methyl maleic acid, dimethyl maleic acid, phenyl maleic acid, bromo maleic acid, chloromaleic acid, dichloromaleic acid, fluoromaleic acid, difluoro maleic acid, nonyl hydrogen maleate, decyl hydrogen maleate, dodecyl hydrogen maleate, octadecyl hydrogen maleate, fluoroalkyl hydrogen maleate, maleic anhydride, methyl maleic anhydride, dimethyl maleic anhydride, acrylic anhydride, methacrylic anhydride, methacrolein, methacryloyl chloride, methacryloyl fluoride, methacryloyl bromide and combinations thereof.

3. The binder composition according to claim 1, wherein the secondary battery is a lithium-ion battery.

4. The binder composition according to claim 1, wherein the amide group-containing monomer is selected from the group consisting of methacrylamide, N-methyl methacrylamide, N-ethyl methacrylamide, N-n-propyl methacrylamide, N-isopropyl methacrylamide, N-n-butyl methacrylamide, N-isobutyl methacrylamide, N,N-dimethyl acrylamide, N,N-dimethyl methacrylamide, N,N-diethyl acrylamide, N,N-diethyl methacrylamide, N-methylol methacrylamide, N-(methoxymethyl)methacrylamide, N-(ethoxymethyl)methacrylamide, N-(propoxymethyl)methacrylamide, N-(butoxymethyl)methacrylamide, N,N-dimethylaminopropyl methacrylamide, N,N-dimethylaminoethyl methacrylamide, N,N-dimethylol methacrylamide, diacetone methacrylamide, methacryloyl morpholine and combinations thereof.

5. The binder composition according to claim 1, wherein the nitrile group-containing monomer is selected from the group consisting of acrylonitrile, α-halogenoacrylonitrile, α-alkylacrylonitrile, α-chloroacrylonitrile, α-bromoacrylonitrile, α-fluoroacrylonitrile, methacrylonitrile, α-ethylacrylonitrile, α-isopropylacrylonitrile, α-n-hexylacrylonitrile, α-methoxyacrylonitrile, 3-methoxyacrylonitrile, 3-ethoxyacrylonitrile, α-acetoxyacrylonitrile, α-phenylacrylonitrile, α-tolylacrylonitrile, α-(methoxyphenyl)acrylonitrile, α-(chlorophenyl)acrylonitrile, α-(cyanophenyl)acrylonitrile, vinylidene cyanide and combinations thereof.

6. The binder composition according to claim 1, wherein the dispersion medium is water.

7. The binder composition according to claim 6, wherein the dispersion medium further comprises a hydrophilic solvent selected from the group consisting of ethanol, isopropanol, n-propanol, tert-butanol, n-butanol, dimethylacetamide (DMAc), dimethylformamide (DMF), N-methylpyrrolidone (NMP), methyl ethyl ketone (MEK), ethyl acetate (EA), butyl acetate (BA) and combinations thereof.

8. The binder composition according to claim 1, wherein the proportion of the sum of structural unit (a) derived from a carboxylic acid group-containing monomer and structural unit (b) derived from an amide group-containing monomer in the copolymer is from 18% to 35% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition.

9. The binder composition according to claim 1, wherein the molar ratio of the structural unit (c) derived from a nitrile group-containing monomer to the sum of the structural unit (a) derived from a carboxylic acid group-containing monomer and structural unit (b) derived from an amide group-containing monomer in the copolymer is from 1.5 to 4; and wherein the molar ratio of the sum of the structural unit (c) derived from a nitrile group-containing monomer and structural unit (a) derived from a carboxylic acid group-containing monomer to the structural

unit (b) derived from an amide group-containing monomer in the copolymer is from 5 to 15.

10. The binder composition according to claim 1, wherein the pH of the binder composition is from 7 to 9; wherein the viscosity of the binder composition as measured at 25 °C with a rotational viscosity meter is from 10,000 mPa·s to 50,000 mPa·s; wherein the electrolyte swelling of the binder composition as measured by the ratio of weight change from soaking a previously-dried binder composition strip in electrolyte ($LiPF_6$ (1 M) in a mixture of ethylene carbonate(EC), ethyl methyl carbonate (EMC) and dimethyl carbonate (DMC) at a volume ratio of 1:1:1) at 25 ºC for 3 days to the weight of the dry strip is from 2% to 4%; and wherein the solid content of the binder composition is from 12% to 18% by weight, based on the total weight of the binder composition.

11. An electrode for a secondary battery, comprising an electrode active material, a conductive agent and the binder composition according to claim 1.

12. The electrode according to claim 11, wherein the electrode active material is selected from the group consisting of $LiCoO_2$, $LiNiO_2$, $LiNi_xMn_yO_2$, $Li_{1+z}Ni_xMn_yCo_{1-x-y}O_2$, $LiNi_xCo_yAl_zO_2$, LiV_2O_5, $LiTiS_2$, $LiMoS_2$, $LiMnO_2$, $LiCrO_2$, $LiMn_2O_4$, Li_2MnO_3, $LiFeO_2$, $LiFePO_4$, $Li_4Ti_5O_{12}$, $LiNi_{0.92}Mn_{0.04}Co_{0.04}O_2$, $Li_{1+d}Ni_aMn_bCo_cAl_{(1-a-b-c)}O_2$ and combinations thereof; wherein each x is independently from 0.2 to 0.9; each y is independently from 0.1 to 0.45; wherein each z is independently from 0 to 0.2; and wherein $-0.2 \leq d \leq 0.2$, $0 \leq a < 1$, $0 \leq b < 1$, $0 \leq c < 1$, and $a+b+c \leq 1$.

13. A binder composition for a secondary battery electrode comprising a copolymer, wherein the copolymer comprises a structural unit (a) derived from a carboxylic acid group-containing monomer, a structural unit (b) derived from an amide group-containing monomer and a structural unit (c) derived from a nitrile group-containing monomer; wherein the proportion of structural unit (a) derived from a carboxylic acid group-containing monomer in the copolymer is from 7% to 25% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition; wherein the proportion of structural unit (b) derived from an amide group-containing monomer in the copolymer is from 4% to 17% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition; wherein the proportion of structural unit (c) derived from a nitrile group-containing monomer in the copolymer is from 65% to 80% by mole, based on the total number of moles of monomeric units in the copolymer in the binder composition; and wherein the binder composition is free of a structural unit derived from an ester group-containing monomer.

14. The binder composition according to claim 13, wherein the adhesive strength between the binder composition and a current collector is from 2 N/cm to 4 N/cm.