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ELECTROCHEMICAL APPARATUS AND ELECTRONIC APPARATUS

Abstract

An electrochemical apparatus includes a positive electrode plate, the positive electrode plate includes a positive electrode current collector and a positive electrode active material layer disposed on one or two sides of the positive electrode current collector, and the positive electrode active material layer includes a positive electrode active material. A ratio of $D_{\text{sub.v99}}$ of the positive electrode active material to thickness $H_{\text{sub.1}}$ of the positive electrode active material layer on a single side of the positive electrode current collector satisfies $0.5 \leq D_{\text{sub.v99}}/H_{\text{sub.1}} \leq 0.9$. Agglomeration regions with a bright spot having a diameter $D_{\text{sub.0}}$ greater than or equal to $20 \mu\text{m}$ are observed on a surface of the positive electrode active material layer under a scanning electron microscope, and the agglomeration regions have a number density less than or equal to 5 pcs/cm^{sup.2} per unit area on the surface of the positive electrode active material layer.

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Background/Summary

CROSS-REFERENCE TO RELATED APPLICATIONS [0001] This application is a continuation of International Application No. PCT/CN2023/123525, filed on Oct. 9, 2023, which claims priority to Chinese Patent application Ser. No. 20/221,1353337.1, filed on Nov. 1, 2022, and entitled “ELECTROCHEMICAL APPARATUS AND ELECTRONIC APPARATUS”, which is incorporated herein by reference in its entirety.

TECHNICAL FIELD

[0002] This application relates to the field of electrochemical apparatus technologies, and specifically, to an electrochemical apparatus and an electronic apparatus.

BACKGROUND

[0003] Consumer electronics, electric vehicles, and other devices have become an indispensable part of people's lives, and these devices require electrochemical apparatuses to provide electrical energy for them, thereby driving the rapid development of electrochemical apparatuses represented by lithium-ion secondary batteries.

[0004] However, in the related technology, the electrochemical apparatus has poor high-temperature cycling performance, which limits its application in some devices.

SUMMARY

[0005] This application provides an electrochemical apparatus and an electronic apparatus, the electrochemical apparatus having good high-temperature cycling performance.

[0006] According to a first aspect, some embodiments of this application provide an electrochemical apparatus including a positive electrode plate, where the positive electrode plate includes a positive electrode current collector and a positive electrode active material layer disposed on one or two sides of the positive electrode current collector, and the positive electrode active material layer includes a positive electrode active material. A ratio of D_{99} of the positive electrode active material to thickness H_{10} of the positive electrode active material layer on a single side of the positive electrode current collector satisfies $0.5 \leq D_{99}/H_{10} \leq 0.9$. Agglomeration regions with a bright spot having a diameter D_0 greater than or equal to 20 μm are observed on a surface of the positive electrode active material layer under a scanning electron microscope, and the agglomeration regions have a number density less than or equal to 5 pcs/ cm^2 per unit area on the surface of the positive electrode active material layer.

[0007] In the electrochemical apparatus provided in the embodiment of this application, a ratio of D_{99} of the positive electrode active material to thickness H_{10} of the positive electrode active material layer on a single side of the positive electrode current collector satisfies $0.5 \leq D_{99}/H_{10} \leq 0.9$. This allows for the thinning of the positive electrode plate, reducing the ion transport path, thereby reducing ion polarization and the resistance of the electrochemical apparatus. In addition, agglomeration regions with a bright spot having a diameter D_0 greater

than or equal to 20 μm on surface of the positive electrode active material layer observed under a scanning electron microscope have a number density less than or equal to 5 pcs/cm.² per unit area on the surface of the positive electrode active material layer, such that an amount of the positive electrode active material layer crushed during a rolling process is reduced, thereby improving high-temperature cycling performance of the electrochemical apparatus.

[0008] According to any one of the preceding embodiments of the first aspect of this application, D.sub.v99 of the positive electrode active material satisfies $15\ \mu\text{m} \leq D_{\text{sub.v99}} \leq 30\ \mu\text{m}$, and thickness H1 of the positive electrode active material layer on one single side satisfies $10\ \mu\text{m} \leq H_{\text{sub.1}} \leq 35\ \mu\text{m}$.

[0009] According to any one of the preceding embodiments of the first aspect of this application, a ratio of D.sub.v50 of the positive electrode active material and D.sub.v10 of the positive electrode active material satisfies $1.0 \leq D_{\text{sub.v50}}/D_{\text{sub.v10}} \leq 2.5$.

[0010] According to any one of the preceding embodiments of the first aspect of this application, a ratio of D.sub.v50 of the positive electrode active material and D.sub.v10 of the positive electrode active material satisfies $1.0 \leq D_{\text{sub.v50}}/D_{\text{sub.v10}} \leq 2.0$.

[0011] According to any one of the preceding embodiments of the first aspect of this application, the positive electrode active material includes a nickel-cobalt-manganese ternary material.

[0012] According to any one of the preceding embodiments of the first aspect of this application, a molar fraction of nickel in the nickel-cobalt-manganese ternary material is greater than or equal to 60%.

[0013] According to any one of the preceding embodiments of the first aspect of this application, thickness H.sub.2 of the positive electrode current collector satisfies at least one of the following characteristics:

[00001](I) $0.3 \leq H_2 / D_{\text{v99}} \leq 1$; (II) $1 \leq H_1 / H_2 \leq 4$; or (III) $7\ \mu\text{m} \leq H_2 \leq 20\ \mu\text{m}$.

[0014] According to any one of the preceding embodiments of the first aspect of this application, thickness H2 of the positive electrode current collector satisfies $8\ \mu\text{m} \leq H_{\text{sub.2}} \leq 12\ \mu\text{m}$.

[0015] According to any of the preceding embodiments of the first aspect of this application, the surface density of the positive electrode active material layer is 80 mg/1540.25 mm² to 180 mg/1540.25 mm.².

[0016] According to any one of the preceding embodiments of the first aspect of this application, the positive electrode active material layer has a compacted density of 3.3 g/cm.³ to 3.7 g/cm.³.

[0017] According to any one of the preceding embodiments of the first aspect of this application, the positive electrode active material layer has a compacted density of 3.5 g/cm.³ to 3.65 g/cm.³.

[0018] According to any one of the preceding embodiments of the first aspect of this application, the positive electrode active material layer further includes a binder, the binder has a corresponding Fourier transform infrared spectral characteristic peak in the range of 1655 cm⁻¹ to 1705 cm⁻¹.

[0019] According to any one of the preceding embodiments of the first aspect of this application, the binder includes a polyvinylidene fluoride copolymer.

[0020] According to any one of the preceding embodiments of the first aspect of this application, the binder has a structural formula represented by formula (A):



[0021] In formula (A), VDF is a structural unit of polyvinylidene fluoride, TFE is a structural unit of polytetrafluoroethylene, HFP is a structural unit of polyhexafluoropropylene, PVP is a structural unit of polyvinylpyrrolidone, $0.35 \leq m \leq 1$, $0 \leq n \leq 0.4$, $0 \leq r \leq 0.2$, $0 \leq x \leq 0.2$, and $m+n+r+x=1$.

[0022] According to a second aspect, some embodiments of this application provide an electronic apparatus including the electrochemical apparatus described in any one of the preceding embodiments of the first aspect of this application.

[0023] The foregoing description is merely an overview of the technical solution of this application. For a better understanding of the technical means in this application such that they can be implemented according to the content of the specification, and to make the above and other objectives, features, and advantages of this application more obvious and easier to understand, the following describes specific embodiments of this application.

Description

BRIEF DESCRIPTION OF DRAWINGS

[0024] A person of ordinary skill in the art can clearly understand various other advantages and benefits by reading the detailed description of the embodiments below. The accompanying drawings are merely intended to illustrate some embodiments and are not intended to limit this application. In addition, in all the accompanying drawings, same parts are denoted by same reference signs. In the accompanying drawings:

[0025] FIG. 1 is a schematic structural diagram of an electrochemical apparatus according to an embodiment of this application.

[0026] FIG. 2 is a schematic diagram of an exploded structure of the electrochemical apparatus shown in FIG. 1.

[0027] FIG. 3 is an SEM image of a positive electrode plate in the electrochemical apparatus according to Example 1 of this application.

[0028] FIG. 4 is an SEM image of a positive electrode plate in the electrochemical apparatus according to Comparative example 1 of this application.

DESCRIPTION OF REFERENCE SIGNS

[0029] 5. electrochemical apparatus, 51. housing, 52. electrode assembly, and 53. top cover assembly.

DETAILED DESCRIPTION

[0030] The following describes in detail the embodiments of technical solutions of this application with reference to the accompanying drawings. The following embodiments are merely intended for a clearer description of the technical solutions of this application and therefore are used as just examples which do not constitute any limitations on the protection scope of this application.

[0031] Unless otherwise defined, all technical and scientific terms used herein shall have the same meanings as commonly understood by those skilled in the art to which this application relates. The terms used herein are intended to merely describe the specific embodiments rather than to limit this application. The terms “include”, “comprise”, and “have” and any other variations thereof in the specification, claims, and brief description of drawings of this application are intended to cover non-exclusive inclusions.

[0032] In the description of the embodiments of this application, the terms “first”, “second”, and the like are merely intended to distinguish between different objects, and shall not be understood as any indication or implication of relative importance or any implicit indication of the number, sequence or primary-secondary relationship of the technical features indicated. In the description of this application, “a plurality of” means at least two unless otherwise specifically stated.

[0033] In this specification, reference to “embodiment” means that specific features, structures or characteristics described with reference to the embodiment may be incorporated in at least one embodiment of this application. The word “embodiment” appearing in various places in the specification does not necessarily refer to the same embodiment or an independent or alternative embodiment that is exclusive of other embodiments. It is explicitly or implicitly understood by persons skilled in the art that the embodiments described herein may be combined with other embodiments.

[0034] In the description of the embodiments of this application, the term “and/or” is only an

associative relationship for describing associated objects, indicating that three relationships may be present. For example, A and/or B may indicate the following three cases: presence of only A, presence of both A and B, and presence of only B. In addition, the character “/” in this specification generally indicates an “or” relationship between contextually associated objects.

[0035] In the description of some embodiments of this specification, it should be noted that, unless otherwise stated, “more than” and “less than” are inclusive of the present number, and “more” in “more than one piece” or “more than one type” means more than two.

[0036] The grouping of alternative elements or embodiments disclosed herein should not be construed as a limitation. Each member of the group may be adopted and claimed separately, or be adopted and claimed in any combination with other members of the group or other elements found herein. It is foreseeable that, one or more members of a group may be included in or excluded from the group for reasons of convenience and/or patentability. When any such inclusion or exclusion occurs, the specification is hereby considered to contain a modified group, thereby satisfying the written description for all Markush groups used in the claims.

[0037] Various modifications and changes can be made within the scope of this application without departing from the protection of this application, which would be apparent to persons skilled in the art. Therefore, this application is intended to cover modifications and changes to this application that fall within the corresponding claims (scope of protection claimed) and their equivalents. It should be noted that the embodiments provided in this application may be combined unless a contradiction occurs.

[0038] Before explaining the scope of protection provided in the embodiments of this application, this application first specifies the problems in the related art for a better understanding of the embodiments of this application.

[0039] Electrochemical apparatuses, represented by lithium-ion secondary batteries, are widely used in consumer electronics, electric vehicles, and other devices due to their high energy density, good cycling performance, and rechargeability. With the rapid development of secondary battery technology, the development of fast charging and discharging technology has been promoted, making lithium-ion secondary batteries that support fast charging and discharging one of the most popular electrochemical apparatuses in the market.

[0040] In related art, lithium-ion secondary batteries generate a large amount of heat during high-rate charging and discharging, causing the temperature of the lithium-ion secondary batteries to rise, making them prone to polarization. To reduce the polarization of lithium-ion secondary batteries, the thickness of the electrode plate can be reduced to shorten the ion transport path. However, the reduction of the thickness of the electrode plate causes the agglomerated slurry applied on the surface to form bumps, and the active material within the bumps is prone to crushing during the rolling process. This causes the temperature of the lithium-ion secondary battery to rise during the high-rate charging and discharging, resulting in a decrease in the cycling performance of the lithium-ion secondary battery.

[0041] In view of this, some embodiments of this application provide an electrochemical apparatus and an electronic apparatus, the electrochemical apparatus having good high-temperature cycling performance.

[0042] The electrochemical apparatus according to this application includes any apparatus in which an electrochemical reaction takes place. Specific examples of the apparatus include all types of primary batteries, secondary batteries, fuel batteries, solar batteries, or capacitors. For example, the electrochemical apparatus is a lithium secondary battery, the lithium secondary battery including a lithium metal secondary battery, a lithium-ion secondary battery, a lithium polymer secondary battery, or a lithium-ion polymer secondary battery.

Electrochemical Apparatus

[0043] Some embodiments of this application provide an electrochemical apparatus including a positive electrode plate, where the positive electrode plate includes a positive electrode current

collector and a positive electrode active material layer disposed on one or two sides of the positive electrode current collector, and the positive electrode active material layer includes a positive electrode active material. A ratio of D_{v99} of the positive electrode active material to thickness H_1 of the positive electrode active material layer on a single side of the positive electrode current collector satisfies $0.5 \leq D_{v99}/H_1 \leq 0.9$. Agglomeration regions with a bright spot having a diameter D_0 greater than or equal to 20 μm are observed on a surface of the positive electrode active material layer under a scanning electron microscope, and the agglomeration regions have a number density less than or equal to 5 pcs/ cm^2 per unit area on the surface of the positive electrode active material layer.

[0044] In this application, after the bumps on the surface of the electrode plate are rolled, agglomeration regions having a diameter D_0 greater than or equal to 20 μm are formed. The number of agglomeration regions within a unit area is reduced by decreasing the number of bumps on the surface of the electrode plate, thereby improving the uniformity of the electrode plate surface and reducing the decrease in cycling performance caused by the crushing of bumps during the rolling process. Fewer agglomeration regions mean fewer bumps on the surface of the electrode plate before rolling. Therefore, the agglomeration regions on the electrode plate after rolling should be as few as possible.

[0045] In the volume cumulative distribution of the positive electrode active material, D_{v99} is a particle size when the volume cumulative amount reaches 99% as counted from the small particle size side.

[0046] The diameter D_0 is the maximum measurable distance in the region without a clearly defined shape.

[0047] Agglomerated particles in the positive electrode slurry forming the positive electrode active material layer are applied to the surface of the positive electrode current collector and protrude from the surface of the positive electrode active material layer. After being rolled, these agglomerated particles form the agglomeration regions.

[0048] In the electrochemical apparatus provided in the embodiment of this application, a ratio of D_{v99} of the positive electrode active material to thickness H_1 of the positive electrode active material layer on a single side of the positive electrode current collector satisfies $0.5 \leq D_{v99}/H_1 \leq 0.9$. This allows for the thinning of the positive electrode plate, reducing the ion transport path, thereby reducing ion polarization and the resistance of the electrochemical apparatus. In addition, agglomeration regions with a bright spot having a diameter D_0 greater than or equal to 20 μm on surface of the positive electrode active material layer observed under a scanning electron microscope have a number density less than or equal to 5 pcs/ cm^2 per unit area on the surface of the positive electrode active material layer, such that an amount of the positive electrode active material layer crushed during a rolling process is reduced, thereby improving high-temperature cycling performance of the electrochemical apparatus.

[0049] In some embodiments of this application, an appropriate ratio of D_{v99} of the positive electrode active material to the thickness H_1 of the positive electrode active material layer on a single side of the positive electrode current collector not only enables the positive electrode plate to be thinner and reduces defects occurring in the processing process but also reduces the occurrence of crushing of the positive electrode active material.

[0050] In some embodiments of this application, D_{v99} of the positive electrode active material satisfies $15 \mu\text{m} \leq D_{v99} \leq 30 \mu\text{m}$, and the thickness of the positive electrode active material layer on a single side of the positive electrode current collector H_1 satisfies $10 \mu\text{m} \leq H_1 \leq 35 \mu\text{m}$.

[0051] In these embodiments described above, an appropriate ratio of D_{v99} of the positive electrode active material to the thickness H_1 of the positive electrode active material layer on a single side of the positive electrode current collector can reduce the occurrence of defects such as scratches in the processing of the positive electrode active material layer and can further reduce the amount of the positive electrode active material that is crushed in the rolling process, thereby

helping improve the high-temperature cycling performance.

[0052] In some examples, the D_{v99} of the positive electrode active material may be, but is not limited to, 10 μm , 11 μm , 12 μm , 13 μm , 14 μm , 15 μm , 16 μm , 17 μm , 18 μm , 19 μm , 20 μm , 21 μm , 22 μm , 23 μm , 24 μm , 25 μm , 26 μm , 27 μm , 28 μm , 29 μm , or 30 μm . The thickness H_1 of the positive electrode active material layer on a single side of the positive electrode current collector may be, but is not limited to, 10 μm , 11 μm , 12 μm , 13 μm , 14 μm , 15 μm , 16 μm , 17 μm , 18 μm , 19 μm , 20 μm , 21 μm , 22 μm , 23 μm , 24 μm , 25 μm , 26 μm , 27 μm , 28 μm , 29 μm , 30 μm , 31 μm , 32 μm , 33 μm , 34 μm , or 35 μm .

[0053] In addition, other volumetric particle size distributions of the positive electrode active material within an appropriate range can also reduce the occurrence of crushing of the positive electrode active material, thereby reducing the electrical resistance of the positive electrode plate and the expansion rate of the electrochemical apparatus.

[0054] In some embodiments of this application, a ratio of D_{v50} of the positive electrode active material to D_{v10} of the positive electrode active material satisfies $1.0 \leq D_{v50}/D_{v10} \leq 2.5$.

[0055] In the volume cumulative distribution of the positive electrode active material, D_{v50} is a particle size when the volume cumulative amount reaches 50% as counted from the small particle size side.

[0056] In the volume cumulative distribution of the positive electrode active material, D_{v10} is a particle size when the volume cumulative amount reaches 10% as counted from the small particle size side.

[0057] In these embodiments, with the ratio of D_{v50} to D_{v10} of the positive electrode active material satisfying the relationship described above, the crushing of the positive electrode active material can be reduced, thereby decreasing the electrical resistance of the positive electrode plate and the expansion rate of the electrochemical apparatus to further improve the high-temperature cycling performance of the electrochemical apparatus.

[0058] In some embodiments of this application, a ratio of D_{v50} of the positive electrode active material to D_{v10} of the positive electrode active material satisfies $1.0 \leq D_{v50}/D_{v10} \leq 2.0$.

[0059] In these embodiments described above, with the ratio of D_{v50} of the positive electrode active material to D_{v10} satisfying the relationship described above, the crushing of the positive electrode active material can be further reduced to improve the high-temperature cycling performance of the electrochemical apparatus.

[0060] In some examples, a ratio of the diameter D_{v50} of the positive electrode active material to D_{v10} of the positive electrode active material may be, but is not limited to, 1.0, 1.1, 1.2, 1.3, 1.4, 1.5, 1.6, 1.7, 1.8, 1.9, 2.0, 2.1, 2.2, 2.3, 2.4, or 2.5.

[0061] In some embodiments of this application, the thickness of the positive electrode current collector also affects the compacted density and processing of the positive electrode active material layer. In some embodiments of this application, thickness H_2 of the positive electrode current collector satisfies at least one of the following characteristics:

[00002](I) $0.3 \leq H_2 / D_{v99} \leq 1$; (II) $1 \leq H_1 / H_2 \leq 4$; or (III) $7 \mu\text{m} \leq H_2 \leq 20 \mu\text{m}$.

[0062] In these embodiments described above, the thickness H_2 of the positive electrode current collector, H_2/D_{v99} , and H_1/H_2 being within an appropriate range can further improve the compacted density of the positive electrode active material layer, and also reduce the occurrence of breaking of the positive electrode plate during rolling and the occurrence of overpressure crushing of the positive electrode active material. This can lower the resistance of the positive electrode plate and the expansion rate of the electrochemical apparatus, thereby improving the high-temperature cycling performance of the electrochemical apparatus.

[0063] In some embodiments of this application, the thickness H_2 of the positive electrode current collector may be in the range of 8 μm -12 μm . The thickness H_2 of the positive

electrode current collector being in this range may further increase the compacted density of the positive electrode active material and reduce the resistance of the positive electrode plate, thereby improving the high-temperature cycling performance of the electrochemical apparatus.

[0064] In some examples, a ratio of the thickness $H_{sub.2}$ of the positive electrode current collector to $D_{sub.v99}$ of the positive electrode active material may be, but is not limited to, 0.3, 0.4, 0.5, 0.6, 0.7, 0.8, 0.9, or 1. A ratio of the thickness $H_{sub.1}$ of the positive electrode active material layer to the thickness $H_{sub.2}$ of the positive electrode current collector may be, but is not limited to, 1, 1.1, 1.2, 1.3, 1.4, 1.5, 1.6, 1.7, 1.8, 1.9, 2.0, 2.1, 2.2, 2.3, 2.4, 2.5, 2.6, 2.7, 2.8, 2.9, 3.0, 3.1, 3.2, 3.3, 3.4, 3.5, 3.6, 3.7, 3.8, 3.9, or 4.0. The thickness $H_{sub.2}$ of the positive electrode current collector may be, but is not limited to, 7 μm , 8 μm , 9 μm , 10 μm , 11 μm , 12 μm , 13 μm , 14 μm , 15 μm , 16 μm , 17 μm , 18 μm , 19 μm , or 20 μm .

[0065] The embodiments of this application do not specifically limit the material of the positive electrode current collector, and the positive electrode current collector may be a metal foil or a porous metal plate, for example, a metal foil or porous metal plate made of aluminum, copper, nickel, titanium, iron, or another metal, or an alloy thereof. In some embodiments of this application, the positive electrode current collector is an aluminum foil.

[0066] In some embodiments of this application, the positive electrode active material layer has a compacted density of 3.3 g/cm³ to 3.7 g/cm³.

[0067] In these embodiments described above, setting the compacted density of the positive electrode active material layer within the above range can help improve the high-temperature cycling performance of the electrochemical apparatus.

[0068] In other embodiments of this application, the compacted density of the positive electrode active material layer is 3.5 g/cm³ to 3.65 g/cm³.

[0069] In some examples, the compacted density of the positive electrode active material layer may be, but is not limited to, 3.3 g/cm³, 3.31 g/cm³, 3.32 g/cm³, 3.33 g/cm³, 3.34 g/cm³, 3.35 g/cm³, 3.36 g/cm³, 3.37 g/cm³, 3.38 g/cm³, 3.39 g/cm³, 3.40 g/cm³, 3.41 g/cm³, 3.42 g/cm³, 3.43 g/cm³, 3.44 g/cm³, 3.45 g/cm³, 3.46 g/cm³, 3.47 g/cm³, 3.48 g/cm³, 3.49 g/cm³, 3.50 g/cm³, 3.51 g/cm³, 3.52 g/cm³, 3.53 g/cm³, 3.54 g/cm³, 3.55 g/cm³, 3.56 g/cm³, 3.57 g/cm³, 3.58 g/cm³, 3.59 g/cm³, or 3.6 g/cm³.

[0070] The positive electrode active material can directly affect the energy density and cycling performance of the electrochemical apparatus. Therefore, in some embodiments of this application, the positive electrode active material includes a nickel-cobalt-manganese ternary material. The nickel-cobalt-manganese ternary material can help improve the energy density and cycling performance of the electrochemical apparatus.

[0071] The nickel-cobalt-manganese ternary material can be a ternary structure material such as NCM811, NCM622, NCM613, NCM523, NCM111, and the like.

[0072] In NCM811, N represents nickel, C represents cobalt, M represents manganese, and 811 represents a ratio of moles of nickel, cobalt, and manganese in the ternary material, that is, the mole ratio of the nickel element: the cobalt element: the manganese element is 8:1:1, and in this case, the proportion of moles of nickel to total moles of nickel, cobalt, and manganese in the nickel-cobalt-manganese ternary material is 80%. In NCM523, the mole ratio of the nickel element: the cobalt element: the manganese element is 5:2:3, and the proportion of moles of nickel to the total moles of nickel-cobalt-manganese in the nickel-cobalt-manganese ternary material is 50%.

[0073] In some embodiments of this application, a proportion of moles of nickel to total moles of nickel, cobalt, and manganese in the nickel-cobalt-manganese ternary material is greater than or equal to 60%.

[0074] In some other embodiments of this application, the positive electrode active material may further include at least one of a lithium cobaltate material, a lithium manganate material, another metal oxide supporting lithium deintercalation or intercalation, or an olivine structure material such

as lithium manganese iron phosphate, lithium iron phosphate, and lithium manganese phosphate.
[0075] The positive electrode active material layer is formed by coating the surface of the positive electrode current collector with a positive electrode slurry, and the surface density of the positive electrode active material layer has an impact on the energy density and cycling performance of the electrochemical apparatus.

[0076] In this application, the surface density refers to a mass of the positive electrode active material per unit area.

[0077] In some embodiments of this application, the surface density of the positive electrode active material layer is 80 mg/1540.25 mm² to 180 mg/1540.25 mm².

[0078] In these embodiments described above, setting the surface density of the positive electrode active material layer within the appropriate range described above may help components in the positive electrode active material layer to be uniformly dispersed on the surface of the positive electrode current collector, reducing the formation of gels, thereby improving the high-temperature cycling performance of the electrochemical apparatus.

[0079] In some examples, the surface density of the positive electrode active material layer may be, but is not limited to, 80 mg/1540.25 mm²-90 mg/1540.25 mm², 91 mg/1540.25 mm²-100 mg/1540.25 mm², 101 mg/1540.25 mm²-110 mg/1540.25 mm², 111 mg/1540.25 mm²-120 mg/1540.25 mm², 121 mg/1540.25 mm²-130 mg/1540.25 mm², 131 mg/1540.25 mm²-140 mg/1540.25 mm², 141 mg/1540.25 mm²-150 mg/1540.25 mm², 151 mg/1540.25 mm²-160 mg/1540.25 mm², 161 mg/1540.25 mm²-170 mg/1540.25 mm², or 171 mg/1540.25 mm²-180 mg/1540.25 mm².

[0080] In some embodiments of this application, the positive electrode active material layer further includes a binder, the binder has the following Fourier-transform infrared spectral characteristic peak in the range of 1655 cm⁻¹ to 1705 cm⁻¹.

[0081] In these embodiments, the binder can improve the stability of slurry viscosity and, particularly in the case of the positive electrode active material layer including a high-nickel positive electrode active material, can further reduce the formation of gels, thereby contributing to the formation of a uniform and thinned positive electrode active material layer.

[0082] In some embodiments of this application, the binder includes a polyvinylidene fluoride copolymer.

[0083] In some embodiments of this application, the binder has the structural formula represented by formula (A):



[0084] In formula (A), VDF is a structural unit of polyvinylidene fluoride, TFE is a structural unit of polytetrafluoroethylene, HFP is a structural unit of polyhexafluoropropylene, PVP is a structural unit of polyvinylpyrrolidone, $0.35 \leq m \leq 1$, $0 \leq n \leq 0.4$, $0 \leq r \leq 0.2$, $0 \leq x < 0.2$, and $m+n+r+x=1$.

[0085] In these embodiments described above, the binder having the structural formula represented by formula (A) can contribute to the dispersion of the slurry components such as the conductive agent and the positive electrode active material, to inhibit the formation of gels. This further reduces the occurrence of abnormal agglomeration between the components, thereby reducing the number of bumps on the surface of the electrode plate as well as the number of agglomeration regions formed by rolling the bumps. Especially in the high-nickel positive electrode active material system, the formation of gels in the slurry is further inhibited, thereby further improving the high-temperature cycling performance of the electrochemical apparatus.

[0086] In an embodiment of this application, the binder having formula (A) can be produced by the following preparation method:

[0087] After the reactor with a volume of 25 L is vacuumed and nitrogen is pumped to replace oxygen, 18 kg of deionized water, 200 g of perfluorooctanoic acid sodium solution with a mass

concentration of 5%, and 80 g of paraffin wax (with a melting point of 60° C.) are put into the reactor, then, the stirring speed is adjusted to 130 rpm/min, and the temperature of the reactor is increased to 85° C.

[0088] Vinylidene fluoride monomer, tetrafluoroethylene monomer, hexafluoropropylene monomer, and vinylpyrrolidone monomer are added proportionally to the reactor with a reactor pressure of 5.0 MPa, and then 1.15 g of initiator dioctyl peroxydicarbonate is added to carry out the polymerization reaction.

[0089] Then, vinylidene fluoride monomer is added to maintain the reactor pressure at 5.0 MPa, 0.01 g of initiator is added every 10 min, and chain transfer agent HFC-4310 is added in four batches at conversion rates of 20%, 40%, 60%, and 80% respectively, with 5 g added each time, where a total of 5 kg of vinylidene fluoride monomer is added to the reaction.

[0090] When the pressure in the reactor is decreased to 4.0 MPa after 140 minutes of the reaction time, gas is released to collect materials, followed by centrifugation, washing, and drying, to produce a binder having formula (A).

[0091] In some embodiments of this application, the binder satisfies at least one of the following characteristics: [0092] (I) weight-average molecular weight M_w of the binder satisfies $900000 \leq M_w \leq 1200000$; or [0093] (II) weight-average molecular weight M_w and number-average molecular weight M_n of the binder satisfy $1.8 \leq M_w/M_n \leq 2.4$.

[0094] In these embodiments above, the binder satisfying at least one of (I) or (II) can contribute to the dispersion of the slurry components such as the conductive agent and the positive electrode active material, to inhibit the formation of gels. This further reduces the occurrence of abnormal agglomeration between the components. Especially in the high-nickel positive electrode active material system, the formation of gels in the slurry is further inhibited, thereby further improving the high-temperature cycling performance of the electrochemical apparatus.

[0095] In some embodiments of this application, based on the weight of the positive electrode active material layer, weight percentage X of the binder satisfies $0.5\% \leq X \leq 3\%$.

[0096] In these embodiments, the weight percentage X of the binder being within the above appropriate range enables a high bonding strength between the positive electrode active material layer and the positive electrode current collector, so that the stability of the positive electrode active material layer is improved.

[0097] In some other embodiments of this application, the weight percentage X of the binder satisfies $0.7\% \leq X \leq 1.5\%$.

[0098] In some examples, the weight percentage X of the binder may be 0.5%, 0.6%, 0.7%, 0.8%, 0.9%, 1%, 1.1%, 1.2%, 1.3%, 1.4%, 1.5%, 1.6%, 1.7%, 1.8%, 1.9%, 2%, 2.1%, 2.2%, 2.3%, 2.4%, 2.5%, 2.6%, 2.7%, 2.8%, 2.9%, or 3%.

[0099] In some embodiments of this application, the positive electrode active material layer further includes a conductive agent, and the conductive agent contributes to improving the electronic conductivity of the positive electrode plate. For example, the conductive agent is selected from at least one of a carbon-based material, a metal-based material, a conductive polymer, or a mixture thereof. For example, the carbon-based material is selected from carbon black, acetylene black, Ketjen black, carbon fiber, carbon nanotubes, or any combination thereof. The metal-based material is selected from metal powder, metal fiber, copper, nickel, aluminum, or silver. The conductive polymer is a polyphenylene derivative.

[0100] In some embodiments of this application, bonding strength F of the positive electrode active material layer satisfies $10 \text{ N/m} \leq F \leq 40 \text{ N/m}$.

[0101] In these embodiments above, the binder can help make the bonding strength F of the positive electrode active material layer satisfy the above relationship. This is beneficial for meeting the bonding requirements during the processing of the positive electrode plate, and can also delay the increase speed of expansion rate of the positive electrode plate during cycling, resulting in better high-temperature cycling performance of the electrochemical apparatus.

[0102] The bonding strength in these embodiments described above can be obtained by testing using a test method as follows: [0103] (1) taking a dried positive electrode plate and cut it into a sample with a width of 30 mm and a length of 100 mm-160 mm using a blade; [0104] (2) attaching a special double-sided tape to a steel plate, where the double-sided tape has a width of 20 mm and a length of 90 mm-150 mm; [0105] (3) attaching the sample to the double-sided tape with the testing surface facing down to fit the double-sided tape; [0106] (4) inserting a paper strip with the same width as the sample and a length greater than 80 mm below the sample, and fixing it with crepe adhesive to create the test sample; [0107] (5) turning on the tensile testing machine (brand: SAA, model: Instron 3365), and after the indicator lights, adjusting the limit block to an appropriate position; [0108] (6) fixing the test sample to the testing platform, folding the paper strip upward, fixing it with a fixture, and starting to pull the paper strip at a speed of 10 mm/min within a testing range from 0 mm to 40 mm at a 90° angle to separate the positive electrode active material layer adhered to the surface of the double-sided tape from the positive electrode current collector until the test is completed; and [0109] (7) saving test data according to the software prompts to obtain data of bonding strength between the positive electrode active material layer and the positive electrode current collector, and after the test is completed, removing the sample and turning off the instrument.

[0110] The positive electrode plate of this application can be prepared according to conventional methods in the art. For example, the active material, the conductive material, and the binder are dispersed in N-methyl pyrrolidone (NMP) to produce a homogeneous positive electrode slurry, the positive electrode slurry is applied on the positive electrode current collector, followed by drying, cold pressing, sheet cutting, slitting, and re-drying, to produce the positive electrode plate.

[0111] In some embodiments of this application, the electrochemical apparatus further includes a negative electrode plate, a separator, and an electrolyte.

[0112] The negative electrode plate includes a negative electrode current collector and a negative electrode active material layer disposed on at least one surface of the negative electrode current collector. It should be understood that the negative electrode plate may be provided with the negative electrode active material layer on one surface of the negative electrode current collector or may be provided with a negative electrode active material layer on two surfaces of the negative electrode current collector. This is not limited in the embodiments of this application.

[0113] The negative electrode current collector may be a metal foil or a porous metal plate, for example, a metal foil or porous metal plate made of copper, nickel, titanium, iron, or another metal, or an alloy thereof. In some embodiments of this application, the negative electrode current collector is a copper foil.

[0114] This application does not limit the type of the negative electrode active material in the negative electrode active material layer, which can be selected according to requirements. For example, another negative electrode active material includes, but is not limited to, at least one of natural graphite, artificial graphite, a mesocarbon microbead (MCMB), hard carbon, soft carbon, silicon, a silicon--carbon composite, SiO, a Li—Sn alloy, a Li—Sn—O alloy, Sn, SnO, SnO.sub.2, a spinel-structure Li.sub.4Ti.sub.5O.sub.12, a Li—Al alloy, and the like.

[0115] In some embodiments of this application, the negative electrode active material layer further includes a binder. The binder may be selected from at least one of styrene-butadiene rubber (SBR), polyacrylic acid (PAA), polyacrylic acid sodium (PAAS), polyacrylamide (PAM), polyvinyl alcohol (PVA), sodium alginate (SA), polymethacrylic acid (PMAA), and carboxymethyl chitosan (CMCS).

[0116] In some embodiments of this application, the negative electrode active material layer further includes a conductive agent. The conductive agent may be selected from at least one of superconducting carbon, acetylene black, carbon black, Ketjen black, carbon dots, carbon nanotubes, graphene, and carbon nanofiber.

[0117] In some embodiments of this application, the negative electrode active material layer may

further include other promoters such as a thickener (for example, sodium carboxymethyl cellulose (CMC-Na)).

[0118] However, this application is not limited to the above-described materials, and the negative electrode plate of this application may alternatively be made of other well-known materials that may be used as the negative electrode active material, the conductive agent, the binder, and the thickener.

[0119] The negative electrode plate of this application can be prepared according to conventional methods in the art. For example, the negative electrode active material, the conductive agent, the binder, and the thickener are dispersed in a solvent, where the solvent may be N-methyl pyrrolidone (NMP) or deionized water, to form a uniform negative electrode slurry. The negative electrode slurry is applied to the negative electrode current collector, and processes such as drying and cold pressing are performed to produce the negative electrode active material layer and the negative electrode plate.

[0120] The separator is sandwiched between the positive electrode plate and the negative electrode plate to mainly prevent short circuits between the positive and negative electrodes and allow active ions to pass through. The separator is not limited to any particular type in this application and may be any well-known porous separator with good chemical stability and mechanical stability.

[0121] In some embodiments of this application, a material of the separator may be selected from, but is not limited to, one or more of glass fiber, non-woven fabric, polyethylene, polypropylene, and polyvinylidene fluoride. Optionally, the material of the separator may include polyethylene and/or polypropylene. The separator may be a single-layer thin film or a multi-layer composite thin film. When the separator is a multi-layer composite thin film, each layer is made of the same or different materials. In other embodiments of this application, the separator may also be provided with a ceramic coating or a metal oxide coating.

[0122] In the electrochemical apparatus, the electrolyte is a carrier for ion transfer, which can play a role of conducting ions between the positive electrode plate and the negative electrode plate, and guarantees that the electrochemical apparatus obtains advantages such as good cycling performance.

[0123] In some embodiments of this application, the electrolyte includes an organic solvent, a lithium salt, and optional additives. Types of the organic solvent, the lithium salt, and the additives are not specifically limited and can be selected as required.

[0124] In some embodiments of this application, the lithium salt includes, but is not limited to, at least one of lithium hexafluorophosphate (LiPF₆), lithium tetrafluoroborate (LiBF₄), lithium perchlorate (LiClO₄), lithium bis (fluorosulphonyl) amide (LiFSI), lithium bitrifluoromethanesulphonyl amide (LiTFSI), lithium trifluoromethanesulfonate (LiTFS), lithium difluoro (oxalato) borate (LiDFOB), lithium dioxalate borate (LiBOB), lithium difluorophosphate (LiPOF₂), lithium difluoro (dioxalato) phosphate (LiDFOP), or lithium tetrafluoro oxalato phosphate (LiTFOP). One of the foregoing lithium salts may be used alone, or two or more of them may be used.

[0125] In some embodiments of this application, the organic solvent includes, but is not limited to, one or more of ethylene carbonate (EC), propylene carbonate (PC), ethyl methyl carbonate (EMC), diethyl carbonate (DEC), dimethyl carbonate (DMC), dipropyl carbonate (DPC), methyl propyl carbonate (MPC), ethyl propyl carbonate (EPC), butylene carbonate (BC), fluoroethylene carbonate (FEC), methyl formate (MF), methyl acetate (MA), ethyl acetate (EA), propyl acetate (PA), methyl propionate (MP), ethyl propionate (EP), propyl propionate (PP), methyl butyrate (MB), ethyl butyrate (EB), 1,4-butyrolactone (GBL), sulfolane (SF), methyl sulfonyl methane (MSM), methyl ethyl sulfone (EMS), and diethyl sulfone (ESE). The foregoing organic solvents may be used alone, or two or more of them may be used together.

[0126] In some embodiments of this application, the additive may include a negative electrode film-forming additive or a positive electrode film-forming additive or may include an additive

capable of improving some performance of the battery, for example, an additive for improving overcharge performance of the battery, or an additive for improving high-temperature performance or low-temperature performance of the battery.

[0127] In some examples, the additives include, but are not limited to, at least one of fluoroethylene carbonate (FEC), vinyl carbonate (VC), vinyl ethylene carbonate (VEC), vinyl sulfate (DTD), propylene sulfate, ethylene sulfite (ES), 1,3-propanesulfonate lactone (PS), 1,3-propylene sulfonate lactone (PST), sulfonate cyclic quaternary ammonium salt, succinic anhydride, succinonitrile (SN), adiponitrile (AND), tris (trimethylsilane) phosphate (TMSP), or tris (trimethylsilane) borate (TMSB).

[0128] The electrolyte may be prepared using a conventional method in the art. For example, the organic solvent, the lithium salt, and the optional additive may be mixed to produce a uniform electrolyte. A sequence of adding the materials is not particularly limited. For example, the lithium salt and the optional additive may be simultaneously added to the organic solvent to be mixed well to produce the electrolyte; alternatively, the lithium salt is added to the organic solvent, then the optional additive is added to the organic solvent, and the mixture is stirred to uniform to produce the electrolyte.

[0129] In some embodiments of this application, a positive electrode plate, a negative electrode plate, and a separator may be made into an electrode assembly through winding or lamination.

[0130] The electrochemical apparatus in some embodiments of this application further includes an outer package for encapsulating the electrode assembly and the electrolyte. In some embodiments of this application, the outer package may be a hard shell, for example, a hard plastic shell, an aluminum shell, or a steel shell; or a soft pack, for example, a soft pouch. A material of the soft pack may be plastic, for example, at least one of polypropylene (PP), polybutylene terephthalate (PBT), and polybutylene succinate (PBS).

[0131] The embodiments of this application do not impose special limitations on the shape of the electrochemical apparatus, and the electrochemical apparatus may be cylindrical, rectangular, or of any other shape. FIG. 1 shows a rectangular electrochemical apparatus as an example.

[0132] In some embodiments of this application, referring to FIG. 2, the outer package may include a housing 51 and a cover assembly 53. The housing 51 includes a base plate and side plates connected to the base plate, and the base plate and the side plates enclose an accommodating cavity. The housing 51 has an opening communicating with the accommodating cavity, and the cover assembly 53 can cover the opening to seal the accommodating cavity. The positive electrode plate, the negative electrode plate, and the separator may be made into an electrode assembly 52 through winding or lamination. The electrode assembly 52 is enclosed in the accommodating cavity. The electrolyte infiltrates into the electrode assembly 52. The electrochemical apparatus 5 may include one or more electrode assemblies 52, and persons skilled in the art may make choices according to actual requirements.

[0133] After the electrolyte infiltrates the electrode assembly, processes such as vacuum encapsulation, standing, formation, and vacuum molding are performed to produce the electrochemical apparatus.

Electronic Apparatus

[0134] A second aspect of this application provides an electronic apparatus, including the electrochemical apparatus according to the first aspect of this application. The electrochemical apparatus provided in this application has good high-temperature cycling performance and high energy density, so that the electronic apparatus provided in this application has good high-temperature cycling performance.

[0135] The electronic apparatus in some embodiments of this application is not particularly limited and may be any known electronic apparatus used in the prior art. In some embodiments of this application, the electronic apparatus may include but is not limited to a notebook computer, a pen-input computer, a mobile computer, an electronic book player, a portable telephone, a portable fax

machine, a portable copier, a portable printer, a stereo headset, a video recorder, a liquid crystal television, a portable cleaner, a portable CD player, a mini-disc, a transceiver, an electronic notebook, a calculator, a storage card, a portable recorder, a radio, a standby power source, a motor, an automobile, a motorcycle, a motor bicycle, a bicycle, a lighting appliance, a toy, a game console, a clock, an electric tool, a flash lamp, a camera, a large household battery, or a lithium-ion capacitor.

[0136] Content disclosed in this application is described in detail in the following examples. These examples are only for illustrative purposes because various modifications and changes made without departing from the scope of the content disclosed in this application are apparent to persons skilled in the art. Unless otherwise stated, all parts, percentages, and ratios reported in the following examples are based on weight, all reagents used in the examples are commercially available or synthesized in a conventional manner, and can be used directly without further treatment, and all instruments used in the examples are commercially available.

[0137] The following embodiments describe in detail the electrochemical apparatus and the method for manufacturing the electrochemical apparatus using a lithium-ion secondary battery as an example for convenience.

EXAMPLE 1-1

Preparation of Positive Electrode Plate

[0138] Conductive carbon black and a binder were mixed in accordance with the mass ratio of 1:1.5 to get the conductive binder mixture, and then the conductive binder mixture was added to NMP, followed by stirring for mixing, to make a conductive paste (with a solid content of 7%);

[0139] a positive electrode active substance lithium nickel cobalt manganese oxide (NCM) was added to the conductive paste, stirring continued using a vacuum mixer until the mixture was homogeneous, to produce a positive electrode slurry with a solid content of about 75%, where the mass ratio of NCM811, carbon nano tube, and the conductive binder is 97:1:2; and

[0140] the positive electrode slurry was applied onto an aluminum foil with a thickness of 10 μm , followed by drying, cold pressing, cutting, and tab welding, to produce a positive electrode plate.

Preparation of Negative Electrode Plate

[0141] Artificial graphite, styrene-butadiene rubber, and sodium carboxymethyl cellulose were mixed at a mass ratio of 96:2:2 with deionized water and additives, the mixture was stirred to uniform to produce a negative electrode slurry; and the negative electrode slurry was applied onto a copper foil with a thickness of 12 μm , followed by drying, cold pressing, cutting, and tab welding, to produce a negative electrode plate.

Preparation of Electrolyte

[0142] Under a dry argon environment, EC, PC, and DEC (based on a weight ratio of 1:1:1) were mixed, and LiPF₆ was added. The mixture was made even to obtain a base electrolyte, where the concentration of LiPF₆ was 1.15 mol/L.

Preparation of Separator

[0143] A polyethylene (PE) porous polymer film was used as a separator.

Preparation of Lithium-Ion Secondary Battery

[0144] The positive electrode plate, the separator, and the negative electrode plate were sequentially stacked to form an electrode assembly. Then, the electrode assembly was placed into an outer package, and the electrolyte was then injected into the outer package to infiltrate the electrode assembly, followed by processes such as packaging, standing, and formation to produce a lithium-ion secondary battery.

EXAMPLE 1-2 to Example 1-6

[0145] The preparation method was similar to the preparation method of Example 1-1, except that some parameters of the positive electrode plate were different. Refer to Table 1 for details.

COMPARATIVE EXAMPLE 1-1 TO COMPARATIVE EXAMPLE 1-4

[0146] The preparation method was similar to the preparation method of Example 1-1, except that

some parameters of the positive electrode plate were different. Refer to Table 1 for details.

EXAMPLE 2-1 to Example 2-7

[0147] The preparation method was similar to the preparation method of Example 1-1, except that some parameters of the positive electrode plate were different. Refer to Table 2 for details.

EXAMPLE 3-1 TO EXAMPLE 3-9

[0148] The preparation method was similar to the preparation method of Example 1-1, except that some parameters of the positive electrode plate were different. Refer to Table 3 for details.

EXAMPLE 4-1 TO EXAMPLE 4-8

[0149] The preparation method was similar to the preparation method of Example 1-1, except that some parameters of the positive electrode plate were different. Refer to Table 4 for details.

EXAMPLE 5-1 TO EXAMPLE 5-5

[0150] The preparation method was similar to the preparation method of Example 1-1, except that some parameters of the positive electrode plate were different. Refer to Table 5 for details.

TESTING

1. Testing for Surface Density of Positive Electrode ctive Material Layer

[0151] Ten small circular samples were punched from the positive electrode plate coated with the positive electrode active material layer. Each sample has a surface area of 1540.25 mm². The punched positive electrode plate samples were weighed to obtain the average mass m₁. Then, ten small circular samples were punched in the same way from the positive electrode current collector and weighed using a balance to obtain the average mass m₂. The surface density is calculated as (m₁-m₂)/1540.25 mm².

2. Testing for Compacted Density of Positive Electrode Active Material Layer

[0152] The discharged lithium-ion secondary battery under test was disassembled to take out the positive electrode plate, and the positive electrode plate was immersed in DMO (dimethyl oxalate) for 30 min to remove the electrolyte and by-products on the surface of the positive electrode plate, and then it was dried in a fume hood for 4 h. The dried positive electrode plate was taken out and measured using a micrometer to obtain the thickness H_{sub.1} of the positive electrode active material layer of the positive electrode plate. Then, a unit area of the positive electrode active material layer was scraped off from the positive electrode plate using a scraper and measured using a balance to obtain the mass m₃ of the positive electrode active material layer per unit area of the positive electrode plate. The compacted density of the positive electrode active material layer is calculated as m₃/H_{sub.1}.

3. Testing for D_{sub.v50}, D_{sub.v10}, and D_{sub.v99} of Positive Electrode Active Material

[0153] (1) Equipment startup: Firstly, the sample introduction system of the laser diffraction/scattering particle size analyzer (Master Sizer 3000) was turned on, and then the optical system and computer were turned on to preheat the equipment for 30 minutes.

[0154] (2) Cleaning of the sample introduction system: The sample introduction vessel was added with water, and the rotation speed was adjusted to the maximum to clean the vessel for 5 seconds. Then, the rotation speed was adjusted to 0. Cleaning of the sample introduction vessel was repeatedly 3 times to ensure the cleanliness of the sample introduction vessel.

[0155] (3) After a “Manual Measurement” interface was displayed, the material name, refractive index, material type, test time, and test repetitions were set in sequence;

[0156] (4) Light and background light were measured by clicking “start” at the electrode;

[0157] (5) The positive electrode active material was dispersed in aqueous solution (10 mL) to make a sample, the sample was added to the feeder, where the opacity increased with the amount of the sample added. When the opacity reached 8% to 12%, no more samples were added. After the opacity became stable (generally after 30 seconds), “start” was tapped to measure the particle size. After the test was completed, D_{sub.v50}, D_{sub.v10}, and D_{sub.v99} were obtained.

[0158] (6) Three parallel samples were tested to calculate the averages of D_{sub.v50}, D_{sub.v10}, and D_{sub.v99}.

4. Measurement of Thickness H.sub.2 of Positive Electrode Current Collector

[0159] The thickness H.sub.2 of the positive electrode current collector was measured using a micrometer.

5. Measurement of Weight-Average Molecular Weight Mw and Number-Average Molecular Weight Mn of Binder

[0160] GB/T 21863-2008 Gel Permeation Chromatography was referred to for measurement of molecular weight and molecular weight distribution, with the ultra-high performance polymer chromatograph ACQUITY APC and the detector ACQUITY Differential Refractive Detector used. The test steps were as follows:

[0161] (1) Power on and preheating: The chromatographic column and pipeline were installed, the control console, power supply, and the like were turned on, and the testing software Empower was opened.

[0162] (2) Parameter setting: sample volume: 0 μ L to 50 μ L (depending on the concentration of the sample); pump flow rate: 0.2 mL/min; mobile phase: 30 mol/L for NMP solution containing LiBr; sealing cleaning solution: isopropanol; guard column: PL gel 10 μ m MiniMIX B Guard (size: 50 mm \times 4.6 mm \times 2); analytical column: PL gel 10 μ m MiniMIX B (size: 250 mm \times 4.6 mm); standard sample: polystyrene kit; run time: 30 minutes; detector: ACQUITY Differential Refractive Index (RI) Detector; column oven temperature: 90° C.; and detector temperature: 55° C.

[0163] (3) Sample testing: a. Standard and test samples were prepared: 0.002 g to 0.004 g of standard/test samples were weighed and added to 2 mL of mobile phase liquid to prepare a 0.1% to 0.5% mixed standard/test solution. The solution was put into a refrigerator for more than 8 hours. b. Standard solution/samples were tested: The sample group under test was edited, the established sample group method was selected, and after the baseline became stabilized, a run queue was tapped to start testing the samples.

[0164] (4) Data processing: A calibration curve was established using the chemical workstation software according to the relationship between retention time and molecular weight. Integration quantification was performed on the sample chromatograms, and the chemical workstation software automatically generated the molecular weight and molecular weight distribution results.

6. Testing for Diameter D.SUB.0 of Agglomeration Region and Number Density Per Unit Area

[0165] (1) A sample with a width of 5 cm and a length of 5 cm was cut from a dried positive electrode plate using a blade and fixed on a microscope test bench;

[0166] (2) The microscope test device was turned on, the test magnification was adjusted to X100, and adjustment was performed until the interface was clear;

[0167] (3) Measurement was performed: an observation range of 1.0 cm \times 1.0 cm was selected as the observation area, white agglomeration regions in the observation area were counted, and the maximum diameter of the agglomeration regions was recorded as D.sub.0.

[0168] FIGS. 3 and 4 show SEM images of the positive electrode plate in Example 1 and Comparative example 1, respectively.

7. Testing for Resistance of Positive Electrode Plate

[0169] The test was performed using a resistance meter. Details were as follows: the power supply of the equipment was maintained at 220 V, and the atmospheric pressure was greater than 0.7 MPa; the positive electrode plate was removed from the fully discharged battery, and the cut positive electrode plate (60 mm \times 80 mm) was flatly paced on a sample holder; and then the sample holder was placed in the testing chamber of the device to start the testing, where the atmospheric pressure was set to "0" throughout the entire testing process.

8. Testing for Capacity Retention Rate

[0170] At a test ambient temperature of 45° C., the lithium-ion battery after formation was charged to the cut-off voltage of 4.5 V at a current of 1.3 C in the constant current charging stopped. After fully charged, the battery was left standing for 5 min, and then discharged to 3.0 V at a current of 1.0 C. This was one charging and discharging cycle. The battery was charged and discharged for

189 79.20% 7.80% 3-2 Example 23.2 27.1 10 0.43 2.71 256 75.40% 9.80% 3-3 Example 23.2 32.5 10 0.43 3.25 189 79.30% 5.80% 3-4 Example 10.0 27.1 10 1.0 2.71 403 66.90% 12.50% 3-5 Example 15.6 27.1 15 0.96 1.81 310 68.00% 9.50% 3-6 Example 15.6 27.1 9 0.58 0.33 210 71.00% 9.80% 3-7 Example 8.0 27.1 10 1.25 2.71 433 64.20% 16.80% 3-8 Examples 23.2 32.5 8 0.34 4.06 410 65.60% 15.30% 3 to 9

[0177] According to Table 3, a comparison of test results between Examples 3-2, 3-3, 3-5, and 3-8 shows that the particle size distribution of the positive electrode active material D.sub.v99 and H.sub.2/D.sub.v99 being within the appropriate range can help to reduce the electrical resistance of the positive electrode plate as well as the expansion rate of the electrochemical apparatus, thereby increasing the high-temperature cycling performance of the electrochemical apparatus.

[0178] A comparison of test results between Examples 3-3 and 3-4 shows that the thickness H.sub.1 of the positive electrode active material layer and H.sub.1/H.sub.2 being within the appropriate range can contribute to lowering the resistance of the positive electrode plate and the expansion rate of the electrochemical apparatus, thereby improving the high-temperature cycling performance of the electrochemical apparatus.

[0179] A comparison of test results between Examples 3-2, 3-6, and 3-7 shows that the thickness H.sub.2 of the positive electrode current collector and H.sub.1/H.sub.2 being within the appropriate range can contribute to lowering the resistance of the positive electrode plate and the expansion rate of the electrochemical apparatus, thereby improving the high-temperature cycling performance of the electrochemical apparatus.

[0180] Table 4 lists the test results of the positive electrode active materials in Examples 4-1 to 4-8 at different surface densities and compacted densities.

TABLE-US-00004 TABLE 4 Capacity Positive electrode active retention Expansion material layer
 Resis- rate (after rate (after Surface Compacted tance 500 cycles 500 cycles No. density density
 (mΩ) at 45° C.) at 45° C.) Example 4-1 150 3.4 191 73.20% 7.80% Example 4-2 150 3.5 196
 75.30% 8.30% Example 4-3 150 3.6 168 84.50% 5.30% Example 4-4 150 3.7 208 74.70% 8.80%
 Example 4-5 180 3.6 189 79.30% 5.80% Example 4-6 80 3.6 295 78.40% 6.10% Example 4-7 200
 3.6 299 70.40% 11.30% Example 4-8 150 3.8 267 68.40% 12.60% Notes: The unit of the surface
 density is mg/1540.25 mm.sup.2; and the unit of the compacted density is g/cm.sup.3.

[0181] According to Table 4, a comparison of the test results between Examples 4-1 to 4-4 shows that the compacted density of the positive electrode active material layer being within the range of 3.3 g/cm.sup.3 to 3.7 g/cm.sup.3 can enable the positive electrode plate to have a lower electrical resistance as well as enable the electrochemical apparatus to have a lower expansion rate, thereby further improving the high-temperature cycling performance of electrochemical apparatus.

[0182] A comparison of the test results between Examples 4-5 to 4-7 shows that the surface density of the positive electrode active material layer being within the range of 80 mg/1540.25 mm.sup.2 to 180 mg/1540.25 mm.sup.2 can also enable the positive electrode plate to have a lower resistance and enable the electrochemical apparatus to have a lower expansion rate, thereby further improving the high-temperature cycling performance of the electrochemical apparatus.

TABLE-US-00005 TABLE 5 Capacity retention rate Expansion (after 500 rate (after Binder cycles
 at 500 cycles No. Mw Mw/Mn VDF TFE HFP —COOH PVP 45° C.) at 45° C.) Example 5-1 110
 w 2.3 ✓ ✓ ✓ x ✓ 80.5% 5.3% Example 5-2 110 w 2.3 ✓ ✓ ✓ x x 79.3% 5.5% Example 5-3 110
 w 2.3 ✓ ✓ x x x 60.5% 15.3% Example 5-4 110 w 2.3 ✓ x ✓ x x 61.5% 14.2% Example 5-5 105
 w 2.1 ✓ x x ✓ x 63.5% 14.3%

[0183] According to Table 5, it can be learned that in the electrochemical apparatus provided in the embodiments of this application, the binder contained in the positive electrode active material layer can effectively inhibit the formation of gels, helping improve the high-temperature cycling performance of the electrochemical apparatus.

[0184] In conclusion, it should be noted that the foregoing embodiments are for description of the technical solutions of this application only rather than for limiting this application. Although this

application has been described in detail with reference to the foregoing embodiments, persons of ordinary skill in the art should appreciate that they can still make modifications to the technical solutions described in the embodiments or make equivalent replacements to some or all technical features thereof without departing from the scope of the technical solutions of the embodiments of this application. All such modifications and equivalent replacements shall fall within the scope of claims and specification of this application. In particular, as long as there is no structural conflict, the various technical features mentioned in the embodiments can be combined in any manner. This application is not limited to the specific embodiments disclosed in this specification but includes all technical solutions falling within the scope of the claims.

Claims

1. An electrochemical apparatus comprising a positive electrode plate, the positive electrode plate comprises a positive electrode current collector and a positive electrode active material layer disposed on one or two sides of the positive electrode current collector, and the positive electrode active material layer comprises a positive electrode active material; wherein in the volume cumulative distribution of the positive electrode active material, $D_{.v99}$ is a particle size when the volume cumulative amount reaches 99% as counted from the small particle size side; a ratio of $D_{.v99}$ of the positive electrode active material to a thickness $H_{.1}$ of the positive electrode active material layer on a single side of the positive electrode current collector satisfies $0.5 \leq D_{.v99}/H_{.1} \leq 0.9$; and agglomeration regions with a bright spot having a diameter $D_{.0}$ greater than or equal to $20 \mu\text{m}$ are observed on a surface of the positive electrode active material layer under a scanning electron microscope, and the agglomeration regions have a number density less than or equal to 5 pcs/cm^2 per unit area on the surface of the positive electrode active material layer.
2. The electrochemical apparatus according to claim 1, wherein $15 \mu\text{m} \leq D_{.v99} \leq 30 \mu\text{m}$, and $10 \mu\text{m} \leq H_{.1} \leq 35 \mu\text{m}$.
3. The electrochemical apparatus according to claim 1, wherein in the volume cumulative distribution of the positive electrode active material, $D_{.v50}$ is a particle size when the volume cumulative amount reaches 50% as counted from the small particle size side, $D_{.v10}$ is a particle size when the volume cumulative amount reaches 10% as counted from the small particle size side; a ratio of $D_{.v50}$ to $D_{.v10}$ satisfies $1.0 < D_{.v50}/D_{.v10} \leq 2.5$.
4. The electrochemical apparatus according to claim 3, wherein $1.0 < D_{.v50}/D_{.v10} \leq 2.0$.
5. The electrochemical apparatus according to claim 1, wherein the positive electrode active material comprises a nickel-cobalt-manganese ternary material.
6. The electrochemical apparatus according to claim 5, wherein a proportion of moles of nickel to total moles of nickel, cobalt, and manganese in the nickel-cobalt-manganese ternary material is greater than or equal to 60%.
7. The electrochemical apparatus according to claim 1, wherein a thickness $H_{.2}$ of the positive electrode current collector satisfies at least one of the following characteristics: (I) $0.3 \leq H_2 / D_{.v99} \leq 1$; (II) $1 \leq H_1 / H_2 \leq 4$; or (III) $7 \mu\text{m} \leq H_2 \leq 20 \mu\text{m}$.
8. The electrochemical apparatus according to claim 7, wherein $8 \mu\text{m} \leq H_{.2} \leq 12 \mu\text{m}$.
9. The electrochemical apparatus according to claim 1, wherein the positive electrode active material layer has a surface density of $80 \text{ mg/1540.25 mm}^2$ to $180 \text{ mg/1540.25 mm}^2$.
10. The electrochemical apparatus according to claim 1, wherein the positive electrode active material layer has a compacted density of 3.3 g/cm^3 to 3.7 g/cm^3 .
11. The electrochemical apparatus according to claim 10, wherein the positive electrode active material layer has the compacted density of 3.5 g/cm^3 to 3.65 g/cm^3 .
12. The electrochemical apparatus according to claim 1, wherein the positive electrode active material layer further comprises a binder, the binder has a corresponding Fourier transform infrared

spectral characteristic peak in the range of 1655 cm.^{sup.}-1 to 1705 cm.^{sup.}-1.

13. The electrochemical apparatus according to claim 12, wherein the binder comprises a polyvinylidene fluoride copolymer.

14. The electrochemical apparatus according to claim 12, wherein the binder has a structural formula represented by formula (A):

$(VDF)_m(TFE)_n(HFP)_r(PVP)_x$ (A) wherein in formula (A), VDF is a structural unit of polyvinylidene fluoride, TFE is a structural unit of polytetrafluoroethylene, HFP is a structural unit of polyhexafluoropropylene, PVP is a structural unit of polyvinylpyrrolidone, $0.35 \leq m \leq 1$, $0 \leq n \leq 0.4$, $0 \leq r \leq 0.2$, $0 \leq x \leq 0.2$, and $m+n+r+x=1$.

15. An electronic apparatus comprising an electrochemical apparatus, the electrochemical apparatus comprising a positive electrode plate, the positive electrode plate comprises a positive electrode current collector and a positive electrode active material layer disposed on one or two sides of the positive electrode current collector, and the positive electrode active material layer comprises a positive electrode active material; wherein in the volume cumulative distribution of the positive electrode active material, $D_{.v99}$ is a particle size when the volume cumulative amount reaches 99% as counted from the small particle size side; a ratio of $D_{.v99}$ of the positive electrode active material to a thickness $H_{.1}$ of the positive electrode active material layer on a single side of the positive electrode current collector satisfies $0.5 \leq D_{.v99}/H_{.1} \leq 0.9$; and agglomeration regions with a bright spot having a diameter $D_{.0}$ greater than or equal to 20 μm are observed on a surface of the positive electrode active material layer under a scanning electron microscope, and the agglomeration regions have a number density less than or equal to 5 pcs/cm.^{sup.2} per unit area on the surface of the positive electrode active material layer.

16. The electronic apparatus according to claim 15, wherein $15 \mu\text{m} \leq D_{.v99} \leq 30 \mu\text{m}$, and $10 \mu\text{m} \leq H_{.1} \leq 35 \mu\text{m}$.

17. The electronic apparatus according to claim 15, wherein in the volume cumulative distribution of the positive electrode active material, $D_{.v50}$ is a particle size when the volume cumulative amount reaches 50% as counted from the small particle size side, $D_{.v10}$ is a particle size when the volume cumulative amount reaches 10% as counted from the small particle size side; a ratio of $D_{.v50}$ to $D_{.v10}$ satisfies $1.0 < D_{.v50}/D_{.v10} \leq 2.5$.

18. The electronic apparatus according to claim 17, wherein $1.0 < D_{.v50}/D_{.v10} \leq 2.0$.

19. The electronic apparatus according to claim 15, wherein the positive electrode active material comprises a nickel-cobalt-manganese ternary material.

20. The electronic apparatus according to claim 19, wherein a proportion of moles of nickel to total moles of nickel, cobalt, and manganese in the nickel-cobalt-manganese ternary material is greater than or equal to 60%.
