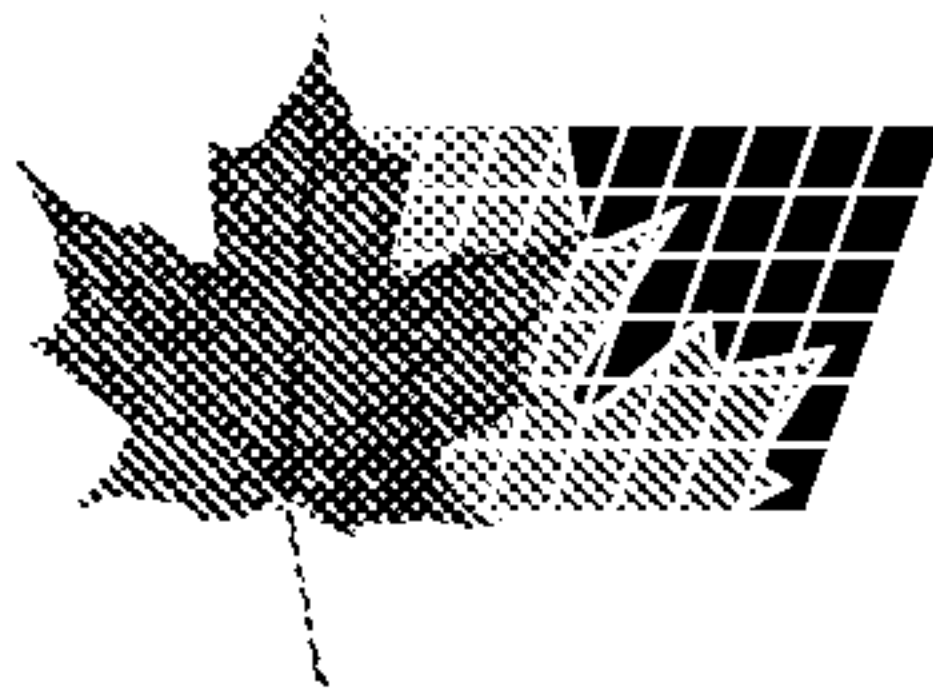




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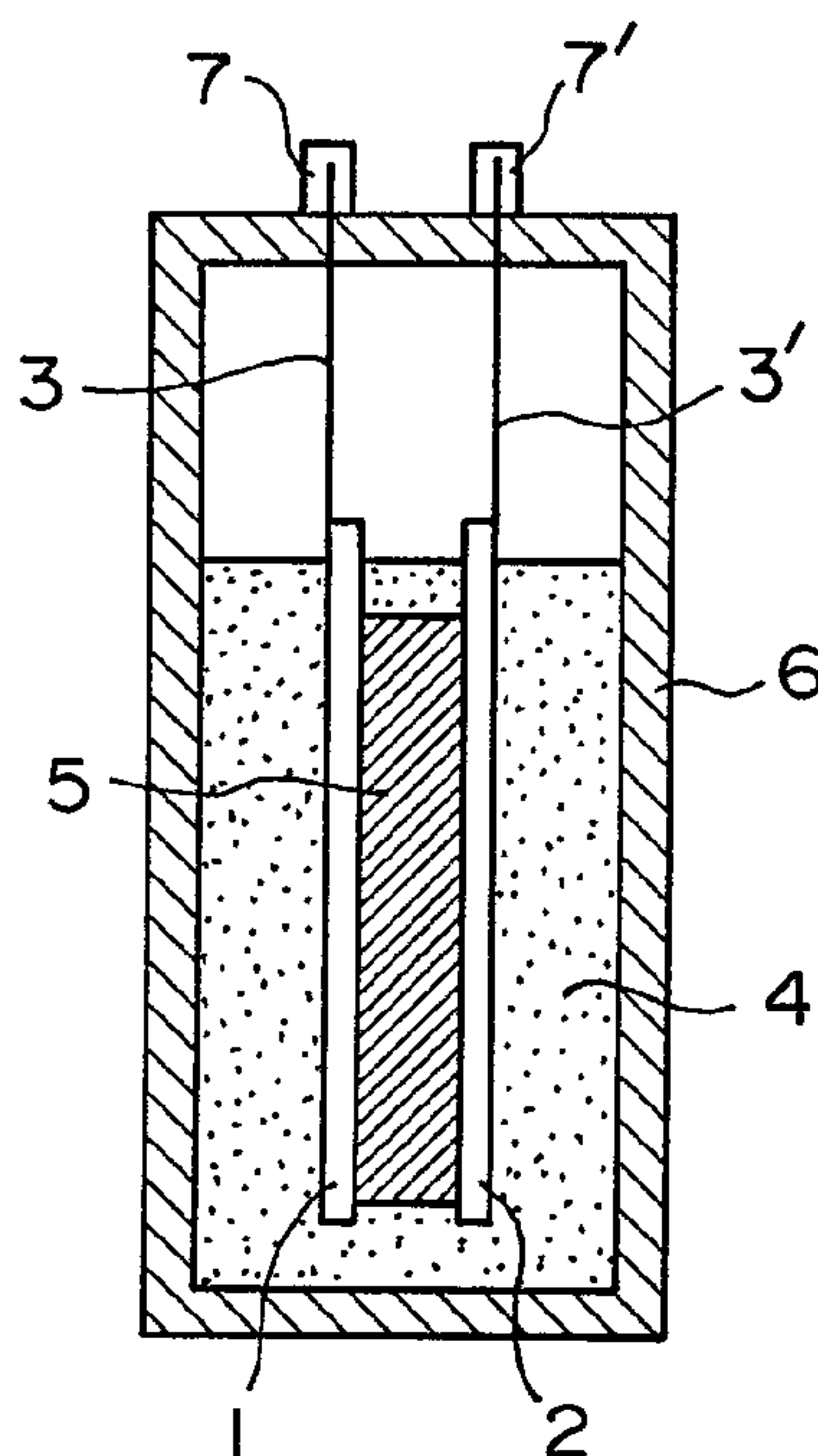
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(54) **ACCUMULATEUR A ELECTROLYTE ORGANIQUE**

(54) **ORGANIC ELECTROLYTIC CELL**



(57) There is provided an organic electrolytic cell equipped with a positive electrode, a negative electrode and a solution of a lithium salt in an aprotic organic solvent as an electrolytic solution, wherein (1) the positive electrode is one containing a metallic oxide, (2) the negative electrode is an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, said substrate being a heat-treated product of an aromatic condensation polymer, and (3) the total amount of lithium contained in the cell is 500 mAh/g or more, and the amount of lithium originating in the negative electrode is 100 mAh/g or more, based on the infusible, insoluble substrate (PAS) having a polyacene type skeletal structure as the negative electrode, the organic electrolytic cell being a secondary cell having high capacity and high voltage.

ABSTRACT

There is provided an organic electrolytic cell equipped with a positive electrode, a negative electrode and a solution of a lithium salt in an aprotic organic solvent as an electrolytic solution, wherein

(1) the positive electrode is one containing a metallic oxide,

(2) the negative electrode is an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, said substrate being a heat-treated product of an aromatic condensation polymer, and

(3) the total amount of lithium contained in the cell is 500 mAh/g or more, and the amount of lithium originating in the negative electrode is 100 mAh/g or more, based on the infusible, insoluble substrate (PAS) having a polyacene type skeletal structure as the negative electrode,

the organic electrolytic cell being a secondary cell having high capacity and high voltage.

DESCRIPTION

Title of the Invention

ORGANIC ELECTROLYTIC CELL

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DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to an organic electrolytic cell having high capacity and high voltage, wherein an infusible, insoluble substrate having a polyacene type skeletal structure is used as a negative electrode, and a metallic oxide is used as a positive electrode.

In recent years, a secondary cell wherein an electrically conductive macromolecule, an oxide of a transition metal or the like is used as a positive electrode, and metallic lithium or a lithium alloy is used as a negative electrode is proposed as a cell to be used in place of Ni-Cd storage cells and lead storage cells, because of its high energy density.

However, when such a secondary cell is subjected to repeated charge and discharge, its capacity is largely lowered due to deterioration of the positive electrode or negative electrode, and thus there still remains a problem in its practical aspect. Particularly by deterioration of the negative electrode, mossy lithium called dendrite is formed, and through repeats of charge and discharge, dendrite finally penetrates the separator and causes a short circuit, and in some case the cell is ruptured, and thus there has been a problem in its safety, too.

Recently, for solving the above problems, there has been proposed a cell wherein a carbonaceous material such as graphite is used as the negative electrode, and a lithium-containing metallic oxide such as LiCoO_2 is used as the positive electrode. The cell is a so-called rocking chair-type cell wherein after assembly

of the cell, lithium is supplied from the lithium-containing metallic oxide as the positive electrode to the negative electrode through charge, and lithium of the negative electrode is returned to the positive electrode through discharge. Although the cell is characterized by high voltage and high capacity, its capacity is at most in the order of 80 to 90 mAh/cc (based on the total volume of the electrodes, the separator and the current collectors); high energy density which is a characteristic of lithium cells has not been obtained.

On the other hand, an infusible, insoluble substrate having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, which is a heat-treated product of an aromatic condensation polymer, can be doped with a larger amount of lithium, compared with general carbonaceous materials. However, when a cell was assembled using the infusible, insoluble substrate, its capacity was not adequately satisfactory.

Thus, the first object of the present invention lies in providing a secondary cell having high capacity and high voltage.

Another object of the invention lies in providing such a secondary cell that charge and discharge are possible over a long term, and are safe.

Still another object of the invention lies in providing a secondary cell which is easy to prepare.

Still another object of the invention will be apparent from the following description.

The present inventors found that for attaining the above objects and advantages, it is important to use a metallic oxide as the positive electrode and an infusible, insoluble substrate having a polyacene type skeletal structure as the negative electrode, and control the

amount of lithium in the cell appropriately.

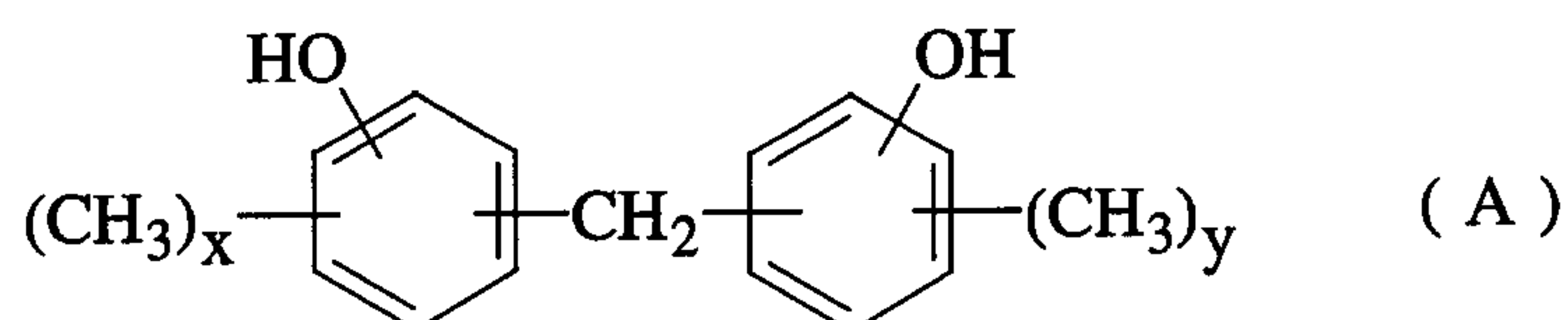
More specifically, the present invention provides an organic electrolyte cell comprising a positive electrode, a negative electrode and a solution of a lithium salt in an aprotic organic solvent, wherein

(1) the positive electrode contains a metallic oxide,

(2) the negative electrode is made of an infusible, insoluble substrate having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, the substrate being a heat-treated product of an aromatic condensation polymer (hereinafter referred to as PAS), and

(3) the total amount of lithium contained in the cell is 500 mAh/g or more, and the amount of lithium originating in the negative electrode is 100 mAh/g or more, based on the negative electrode PAS.

The aromatic condensation polymer used for producing the infusible and insoluble substrate used in the invention is a condensate of an aromatic hydrocarbon compound with an aldehyde. As the aromatic hydrocarbon compound, so-called phenols such as, for example, phenol, cresol and xylenol are preferred. It can, for example, be a methylene-bisphenol represented by the following formula (A)



3a

wherein x and y are independently 0, 1 or 2, or it can also be a hydroxy-biphenyl or a hydroxy-naphthalene. Among them, phenols, particularly phenol are preferred.

As the aromatic condensation polymer,

there can also be used a modified aromatic condensation polymer wherein part of the aromatic hydrocarbon compound having phenolic hydroxyl group(s) is replaced with an aromatic hydrocarbon compound having no phenolic hydroxyl group such as, for example, xylene, toluene or aniline, for example a condensate of phenol, xylene and formaldehyde, and further, there can also be used a modified aromatic polymer wherein the above part is replaced with melamine or urea. Further, furan resins are also preferred.

Further, as the aldehyde, it is possible to use aldehydes such as formaldehyde, acetaldehyde and furfural, but formaldehyde is preferred. A phenol-formaldehyde condensate can be any of a novolak one, a resol one and a mixture thereof.

The infusible, insoluble substrate can be obtained by heat treating the above aromatic polymer, and there can be used all of the infusible, insoluble substrates having a polyacene type skeletal structure described in Japanese Patent Publication No. 44212/1989 (U.S. Patent No. 4,601,849, EP 67444), Japanese Patent Publication No. 24024/1991 (U.S. Patent No. 4,615,960, EP 149497), etc., and such an infusible, insoluble substrate can also be prepared as follows.

An infusible, insoluble substrate having a hydrogen/carbon atomic ratio (hereinafter referred to as H/C) of 0.50 to 0.05, preferably 0.35 to 0.10 can be obtained by gradually heating the aromatic condensation polymer up to a proper temperature of 400°C to 800°C in a non-oxidizing atmosphere (including vacuum) such as nitrogen or argon.

It is also possible to obtain an infusible, insoluble substrate having a specific surface area, measured by the BET method, of 600 m²/g or more according

to the method described in Japanese Patent Publication No. 24024/1991 (U.S. Patent No. 4,615,960, EP 149497), or the like. For example, an infusible, insoluble substrate having the above H/C and having a specific surface area, measured by the BET method, of, e.g., 600 m²/g or more can also be obtained, for example, by preparing a solution containing an initial condensate of an aromatic condensation polymer and an inorganic salt such as zinc chloride; heating the solution to cure it in a mold; gradually heating the cured matter, in a non-oxidizing atmosphere (including vacuum), up to a temperature of 350°C to 800°C, preferably up to a proper temperature of 400°C to 750°C; and then sufficiently washing it with water, diluted hydrochloric acid or the like.

As to an infusible, insoluble substrate used in the invention, according to X-ray diffraction (CuK α), the main peak is observed at $2\theta = 24^\circ$ or less, and besides the main peak, broad, another peak is observed between $2\theta = 41^\circ$ and $2\theta = 46^\circ$.

Namely, it is suggested that the infusible, insoluble substrate has a polyacene type skeletal structure wherein an aromatic polycyclic structure was moderately developed, and takes an amorphous structure, and thus the substrate can be doped stably with lithium, and therefore, is useful as an active material for cells.

When H/C is above 0.50, the aromatic polycyclic structure does not sufficiently develop, and thus it is impossible to conduct doping and undoping of lithium smoothly, and when a cell is assembled, charge and discharge efficiency is lowered. On the other hand, when H/C is less than 0.05, the capacity of the cell of the invention is lowered, which is undesirable.

The negative electrode of the invention is composed of the infusible, insoluble substrate (hereinafter referred to as PAS), and practically, it is desir-

able to use a form obtained by forming PAS in an easily formable form such as a powdery form, a granular form or a short fiber form with a binder.

As the binder, a fluorine binder is preferred, and a fluorine binder having a fluorine/carbon atomic ratio (hereinafter referred to as F/C) of under 1.5 but 0.75 or more is further preferred, and a fluorine-containing polymer binder having a F/C atomic ratio of under 1.3 but 0.75 or more is particularly preferred.

As the fluorine binder, there can, for example, be mentioned polyvinylidene fluoride, a vinylidene fluoride-ethylene trifluoride copolymer, an ethylene-ethylene tetrafluoride copolymer, a propylene-ethylene tetrafluoride copolymer, etc., and further, it is also possible to use a fluorine-containing polymer wherein hydrogens at the principal chain are replaced with alkyl groups. In the case of poly-vinylidene fluoride, F/C is 1, and in the case of the vinylidene fluoride-ethylene trifluoride copolymer, when the molar fractions of vinylidene fluoride are 50 % and 80 %, F/C values become 1.25 and 1.1, respectively, and in the case of the propylene-ethylene tetrafluoride copolymer, when the molar fraction of propylene is 50 %, F/C becomes 0.75. Among them, polyvinylidene fluoride, and a vinylidene fluoride-ethylene trifluoride copolymer wherein the molar fraction of vinylidene fluoride is 50 % or more are preferred, and practically, polyvinylidene fluoride is preferred.

When these binders are used, it is possible to adequately utilize doping ability (capacity) with lithium which PAS has.

As the positive electrode of the organic electrolytic cell of the invention, there can, for example, be used lithium-containing metallic oxides capable of electrochemical doping with lithium and electrochemical undoping of lithium, which can be represented by the

general formula Li_xMyO_z (M is a metal capable of taking plural valences, and can be two or more metals) such as Li_xCoO_2 , Li_xNiO_2 , Li_xMnO_2 or Li_xFeO_2 , or oxides of transition metals such as cobalt, manganese and nickel.
5 Particularly, a lithium-containing oxide having a voltage of 4V or more vs Li/Li^+ is preferred. Among them, lithium-containing cobalt oxides and lithium-containing nickel oxides are preferred.

10 The positive electrode in the invention is one made by forming the metallic oxide, if necessary in addition of an electrically conductive material and a binder, and the kinds, compositions, etc. of the electrically conductive material and binder can suitably be specified.

15 As to the kind of the electrically conductive material, it can be powder of metal such as metallic nickel, but particularly preferred are carbonaceous ones such as, for example, active carbon, carbon black, acetylene black and graphite. Its mixing ratio is
20 varied depending on the electric conductivity of the active substance, the shape of the electrode, etc., but it is suitable to add it in an amount of 2 to 40 % based on the active substance.

Further, as to the kind of the binder, any
25 binder can be used so long as it is insoluble in the later-described electrolytic solution which is used in the invention, and there can, for example, preferably be used rubber binders such as SBR, fluorine-containing resins such as polyethylene tetrafluoride and poly-
30 vinylidene fluoride, and thermoplastic resins such as polypropylene and polyethylene, and its mixing ratio is preferably 20 % or less.

The positive electrode and negative electrode used in the invention can take various shapes, for exam-
35 ple, plate, film and cylinder, and such a shape that the electrode is formed on metallic foil.

Particularly, an electrode obtained by forming the positive electrode or the negative electrode on metallic foil so as to be a film or plate is preferred because such an electrode can be applied to various cells as an electrode which is formed on a current collector.

In the cell of the invention, it is possible to increase its capacity greatly, compared with usual cells, by using the PAS as the negative electrode and appropriately controlling the amount of lithium contained in the cell.

In the invention, the total amount of lithium contained in the cell is the total of lithium originating in the positive electrode, lithium originating in the electrolytic solution, and lithium originating in the negative electrode.

Lithium originating in the positive electrode is lithium contained in the positive electrode at the time of assembly of the cell, and part or all of the lithium is supplied to the negative electrode by an operation (charge or the like) of sending a current from an outer circuit. Lithium originating in the electrolytic solution is lithium in the electrolytic solution contained in the separator, the positive electrode, the negative electrode, etc. Further, lithium originating in the negative electrode is lithium carried on the negative electrode PAS of the invention (lithium other than lithium originating in the positive electrode and lithium originating in the electrolytic solution).

A method for carrying lithium on the negative electrode PAS is not particularly limited, and there can, for example, be mentioned a method which comprises previously doping, before assembly of a cell, the negative electrode PAS with lithium in an electrochemical cell wherein metallic lithium is used as a counter electrode, and then assembling a cell; a method which comprises making electric current conduct between the

negative electrode PAS and metallic lithium in a cell by a method, e.g. of laminating metallic lithium on the negative electrode PAS, and then doping PAS with lithium in the cell; etc.

5 The total amount of lithium in the cell in the invention is 500 mAh/g or more, preferably 600 mAh/g or more based on the negative electrode PAS, and in the case of under 500 mAh/g, adequate capacity cannot be obtained.

10 Further, the amount of lithium originating in the negative electrode is 100 mAh/g or more, preferably 150 mAh/g or more based on the negative electrode PAS, and in the case of under 100 mAh/g, adequate capacity cannot be obtained even if the total amount of lithium
15 is 500 mAh/g or more based on the negative electrode PAS.

 Although as to the amount of lithium originating in the positive electrode and the amount of lithium originating in the electrolytic solution, it is sufficient if the above conditions are satisfied, it is
20 preferred that the amount of lithium originating in the positive electrode is 300 mAh/g or more based on the negative electrode PAS, namely that the amount of lithium originating in the positive electrode is 300 mAh or
25 more per g of the negative electrode PAS.

 Further when a lithium-containing oxide is used as the positive electrode, it is advantageous to make the amount of lithium originating in the negative
30 electrode 600 mAh/g or less based on the negative electrode PAS.

 As the solvent constituting the electrolytic solution used in the invention is used an aprotic organic solvent. As the aprotic organic solvent, there can, for example, be mentioned ethylene carbonate, propylene
35 carbonate, dimethyl carbonate, diethyl carbonate, γ -butyrolactone, acetonitrile, dimethoxyethane,

tetrahydrofuran, dioxolane, methylene chloride, sulfolane, etc, and a mixed solvent of two or more of these aprotic organic solvents can also be used.

Further, as an electrolyte to be dissolved in the mixed or single solvent, any of electrolytes capable of forming lithium ions can be used. As the electrolytes, there can, for example, be mentioned LiI, LiClO₄, LiAsF₆, LiBF₄, LiPF₆, LiHF₂, etc.

The electrolyte and the solvent are mixed in a state of sufficient dehydration to give an electrolytic solution, and for making the internal resistance of the electrolytic solution small, it is preferred to make the concentration of the electrolyte in the electrolytic solution at least 0.1 mole/liter or more, and usually, it is further preferred to make it 0.2 to 1.5 moles/liter.

As the current collector for taking current outside the cell, there can, for example, be used carbon, platinum, nickel, stainless steel, aluminum, copper, etc., and when a foil-like or netlike current collector is used, an electrode can be given as an electrode which is formed on a current collector by forming the electrode on the current collector.

A preferred embodiment of the invention is described below referring to the drawings. Fig. 1 is a cross-sectional side view describing the basic structure of the cell according to the invention. In Fig. 1, (1) is a positive electrode, and (2) is a negative electrode. (3) and (3') are current collectors connected to the respective electrodes and respective outside terminals (7) and (7') so that voltage drop does not arise. (4) is an electrolytic solution, and therein an aforesaid compound capable of forming ions with which the electrodes can be doped, is dissolved in an aprotic organic solvent. The electrolytic solution is usually liquid, but for preventing leakage of the solution, it

can also be used after being made a gel or a solid.
(5) is a separator disposed for preventing contact between both positive and negative electrodes and holding the electrolytic solution.

5 The separator is made of a porous material which is durable against the electrolytic solution, the electrode active substance, etc., has open pores and is electrically non-conductive. It can usually be a cloth, a non-woven cloth, a porous material, or the like composed of glass fiber, polyethylene, polypropylene or the like. To decrease the internal resistance of the cell, the separator is preferably as thin as possible. Its thickness, however, is determined by considering the amount of the electrolytic solution held, its permeability, its strength, etc. The positive and negative electrodes and the separator are fixed in position within a cell casing (6) in a manner not to give rise to any problem in use. The shape, size, etc. of the electrodes can properly be determined according to the shape and
10
15
20 performance of the desired cell.

The shape of the cell of the invention can be coin-type, cylindrical-type, rectangular-type, box-type, etc., but is not particularly limited.

As described above, as to the characteristics and advantages of the organic electrolytic cell of the invention, the organic electrolytic cell is such a cell that PAS is used as the negative electrode, a metallic oxide is used as the positive electrode, both of the amount of lithium in the cell and the amount of lithium originating in the negative electrode PAS are appropriately controlled, and it has high capacity and high voltage.
25
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As described above, the basic structure of the organic electrolytic cell of the invention is to control the total amount of lithium contained in the cell to 500 mAh/g or more, and control the amount of
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lithium originating in the negative electrode to 100 mAh/g or more, but, further, preferred embodiments of the invention are described below.

(1) In the cell of the invention, its capacity can be greatly increased, compared with usual cells, by appropriately controlling the amount of lithium contained in the cell, as described above, and, at the same time, controlling the pore structure of PAS used as the negative electrode so as to be described below.

The amount of nitrogen gas adsorbed on PAS in the invention can be measured as follows. Namely, 0.035 g of PAS fine particles having an average particle size of 15 μm obtained by grinding using a disc mill is put in the sample cell of a volumetric apparatus (made by Yuasa Ionics, Ltd. AutoSorb-1*), and nitrogen gas is adsorbed at a liquid nitrogen temperature of 77° K. From the adsorption isotherm obtained, the amount (cc/g) of the gas adsorbed is plotted against the thickness t (Å) of the layer of the gas adsorbed. t (Å) is calculated by the following equation (1).

$$t \text{ (Å)} = \left(\frac{13.99}{\log(P/P_0) + 0.034} \right)^{1/2} \quad (1)$$

wherein P/P_0 is the relative pressure of nitrogen.

In the invention, it is preferred to control the pore structure of PAS used as the negative electrode so that the amount of the gas adsorbed at the thickness of adsorbed nitrogen of 10 Å found from the above nitrogen adsorption isotherm can be 100 cc/g or less, particularly 80 cc/g or less.

In the invention, when the amount of the gas adsorbed on PAS at the thickness of adsorbed nitrogen of 10 Å is above 100 cc/g, adequate capacity cannot be obtained. Further, in the invention, the total amount of

* Trade-mark

lithium in the cell is 500 mAh/g or more, preferably 600 mAh/g or more, based on the negative electrode PAS, and in the case of under 500 mAh/g, adequate capacity cannot be obtained.

5 Further, the amount of lithium originating in the negative electrode in the invention is 100 mAh/g or more, preferably 150 mAh/g or more, based on the negative electrode PAS, and in the case of under 100 mAh/g, adequate capacity cannot be obtained even if the total
10 lithium amount is 500 mAh/g or more based on the negative electrode PAS.

Although as to the amount of lithium originating in the positive electrode and the amount of lithium originating in the electrolytic solution, it is sufficient if the above conditions are satisfied, it is
15 preferred that the amount of lithium originating in the positive electrode is 300 mAh/g or more based on the negative electrode PAS.

(2) The second preferred embodiment of the
20 invention is described below.

As already stated, in the invention, it is preferred to use as the negative electrode a form obtained by forming fine particles (e.g., granular, powdery, short fiber, etc.) of an infusible, insoluble substrate (hereinafter referred to as PAS) having a
25 polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, the substrate being a heat-treated product of an aromatic condensation polymer, using a binder, preferably a fluorine binder.

30 Thus, in the invention, it is advantageous that a form of fine particles of the infusible, insoluble substrate (PAS), obtained by forming PAS fine particles in an easy to make form such as a powdery form or a granular form with a binder, is used as the negative
35 electrode, and further that the fine particles PAS are such that they have an average particle size of 20 μ m or

less, and when it is supposed that the 50 % size is $2a$ μm , the rate of particles having a particle size of $1a$ μm or less is 10 % or more by volume ratio, and the rate of particles having a particle size of $4a$ μm or more is 10 % or more by volume ratio, and further preferably, when it is supposed that the 50 % size is $2a$ μm , the rate of particles having a particle size of $1a$ μm or less is 20 % or more by volume ratio, and the rate of particles having a particle size of $4a$ μm or more is 10 % or more by volume ratio, and particularly preferably, when it is supposed that the 50 % size is $2a$ μm , the rate of particles having a particle size of $1a$ μm or less is 20 % or more by volume ratio, and the rate of particles having a particle size of $4a$ μm or more is 20 % or more by volume ratio. In the invention, it is advantageous, for obtaining a cell having high capacity, that the fine particles has a wide particle size distribution and an average particle size not more than 20 μm . When the average particle size is above 20 μm , or when even if the average particle size is 20 μm or less, in the case where when it is supposed that the 50 % size is $2a$ μm , the rate of particles having a particle size of $1a$ μm or less is under 10 % by volume ratio, or the rate of particles having a particle size of $4a$ μm or more is under 10 % by volume ratio, the capacity of the resultant cell is low, which is undesirable.

In this connection, the average particle size is a volume average particle size, and the 50 % size is a particle size corresponding to 50 % of an integration curve of particle volume (see the following literature).

"Funtai Riron to Oyo" (Fine Particle Theory and Its Application), edited by Kiichiro Kubo and others, pages 450 to 453, published by Maruzen Co., Ltd. on May 12, 1969.

PAS as fine particles can be obtained by grinding an infusible, insoluble substrate obtained by

heat treating a form of an aromatic polymer. The method of grinding is not particularly limited, but it is efficient to use a grinder having both grinding mechanisms of impact and friction, for example, such a ball mill as a pot mill or a vibrating mill. Further, in some case, PAS as fine particles can also be obtained by classifying the powder obtained, or by mixing two or more of PAS powders having different particle size distributions.

As the binder used in the negative electrode of the invention, a fluorine binder is preferred, as stated already, and further preferred is a fluorine binder having a fluorine/carbon atomic ratio (hereinafter referred to as F/C) of under 1.5 but 0.75 or more, and particularly preferred is a fluorine binder having a F/C of under 1.3 but 0.75 or more.

As the fluorine binder, there can, for example, be mentioned polyvinylidene fluoride, a vinylidene fluoride-ethylene trifluoride copolymer, an ethylene-ethylene tetrafluoride copolymer, a propylene-ethylene tetrafluoride copolymer, etc., and further, it is also possible to use a fluorine-containing polymer wherein hydrogens at the principal chain are replaced with alkyl groups.

The negative electrode of the invention is a form obtained by forming fine particles of PAS with a binder, as stated already, and the porosity of the negative electrode is determined by impregnating the negative electrode with propylene carbonate at 25°C, and is preferably 40 % or less. When the porosity is above 40 %, even if the particle size of PAS is controlled as described above, as to cells obtained therefrom, adequate capacity is hard to get.

It is preferred, as stated above, that the porosity of the negative electrode used in the invention is 40 % or less, and according to experience of the

present inventors, cells of high capacity can be obtained, unexpectedly, even when the porosity of the negative electrode is in the order of 25 %. In the light of this fact, it is believed that there is no
5 problem even if the porosity of the negative electrode is in the order of 20 %.

(3) The third preferred embodiment of the invention is described below.

As the negative electrode of organic electro-
10 lytic cell of the invention is preferred

a negative electrode being obtained by forming an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, the substrate
15 being a heat-treated product of an aromatic condensation polymer, on metallic foil using a thermoplastic binder, and then heating the resultant form at a temperature equal to or higher than the melting point of the thermoplastic binder.

20 As stated already, as this thermoplastic binder are preferred fluorine-containing polymer binders, particularly, a fluorine-containing polymer having a fluorine/carbon atomic ratio of under 1.5 but 0.75 or more, particularly polyvinylidene fluoride.

25 When such a binder is used, it is possible to sufficiently utilize the ability (capacity) of PAS to be doped with lithium.

When, as to the negative electrode of the invention, the PAS is formed with a binder and the
30 resultant form is heat treated at a temperature equal to or higher than the melting point of the thermoplastic binder, the method of heat treatment is not particularly limited, but it is preferred to conduct the heat treatment in a non-oxidizing atmosphere in the range of from
35 a temperature higher by 5°C than the melting point to a temperature higher by 100°C than it. When the heat

treatment is not conducted, for example when the negative electrode PAS formed on metallic foil is doped with lithium in an electrochemical cell wherein metallic lithium is used as a counter electrode, and then a cell
5 is assembled using the doped negative electrode PAS, there arise phenomena, e.g. that the flexural strength of the electrode is weakened, peeling of the electrode is liable to occur, and further, the internal resistance of the assembled cell increases, and as a result, it
10 becomes difficult to get sufficient capacity.

When, in the invention, an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure is formed on metallic foil using a thermoplastic binder, preferably a fluorine-containing polymer
15 binder, the forming is conducted, for example by sufficiently mixing the infusible, insoluble substrate, the fluorine-containing polymer, and a solvent or dispersion medium, and then forming the mixture. The content of the fluorine-containing polymer varies depending on the
20 shape and particle size of the infusible, insoluble substrate, the strength and shape of the desired electrode, etc., but is preferably 2 % to 50 %, more preferably 5 % to 30 % by weight based on the infusible, insoluble substrate. As the solvent are preferred
25 solvents capable of dissolving the fluorine-containing polymer such as N,N-dimethylformamide, N-methylpyrrolidone and N,N-dimethylacetamide. In the above mixture, it does not become a particular problem whether the fluorine-containing polymer is completely dissolved,
30 or only part thereof is dissolved, but it is preferred for obtaining a homogeneous electrode that the fluorine-containing polymer is completely dissolved. Further, the viscosity of the mixture can be controlled by the amount of the solvent, and for example, it is
35 possible to form the mixture adjusted to a high viscosity into a sheet using a roller or the like, and it is

also possible to obtain an extremely thin electrode having a thickness, e.g. of 100 μm or less by applying a mixed slurry adjusted to a low viscosity onto metallic foil, drying it and if necessary, pressing it. Particularly, when excellent flexibility is desired, an application forming method is desirable.

The positive electrode and negative electrode used in the invention can take various shapes, for example, plate, film and cylinder shapes, and such a shape that the electrode is formed on metallic foil. Particularly, an electrode obtained by forming the positive electrode or the negative electrode on metallic foil is preferred because such an electrode can be applied to various cells as an electrode which is formed on a current collector.

(4) The fourth preferred embodiment of the invention is described below.

Further, in the organic electrolytic cell of the invention, it is preferred to use one wherein

- i) the negative electrode is an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, the substrate being a heat-treated product of an aromatic condensation polymer,
- ii) the total amount of lithium contained in the cell is 500 mAh/g or more, and the amount of lithium originating in the negative electrode is 100 mAh/g or more, based on the negative electrode PAS, and
- iii) lithium originating in the negative electrode is previously carried on PAS before assembly of the cell.

Lithium originating in the negative electrode in the invention is lithium carried on the negative electrode PAS of the invention (lithium other than lithium originating in the positive electrode and lithium originating in the electrolytic solution).

In the above preferred embodiment of the invention, a method for carrying lithium on the negative electrode PAS is not particularly limited so long as it is possible by the method to carry lithium on the negative electrode PAS previously before assembly of the cell. For example, it is possible to previously carry lithium on the negative electrode PAS by passing a constant current or applying a constant voltage in an electrochemical cell wherein metallic lithium is used as a counter electrode. When lithium is carried on the negative electrode PAS after assembly of the cell, for example, when a method is taken which comprises making current conduct between the negative electrode PAS and metallic lithium in a cell by a method, e.g. of laminating metallic lithium on the negative electrode PAS, and then doping PAS with lithium in the cell, not only capacity as a practical cell is lowered, but, e.g., the internal resistance of the cell increases, which is undesirable.

Even in this case, it is advantageous that the total amount of lithium contained in the cell is 500 mAh/g or more, preferably 600 mAh/g or more, based on the negative electrode PAS, and in the case of under 500 mAh/g, sufficient capacity cannot be obtained.

Further, the amount of lithium originating in the negative electrode in the invention is 100 mAh/g or more, preferably 150 mAh/g or more, based on the negative electrode PAS, and in the case of under 100 mAh/g, adequate capacity cannot be obtained even if the total lithium amount is 500 mAh/g or more based on the negative electrode PAS. Further, when a lithium-containing oxide is used in the positive electrode, it is practical to make the amount of lithium originating in the negative electrode 600 mAh/g or less based on the negative electrode PAS, as stated already.

Although as to the amount of lithium originat-

ing in the positive electrode and the amount of lithium originating in the electrolytic solution, it is sufficient if the above conditions are satisfied, it is preferred that the amount of lithium originating in the positive electrode is 300 mAh/g or more based on the negative electrode PAS.

An example of still other embodiments of the invention is described below according to a drawing. Fig. 2 is a partially cross-sectional perspective view of the basic structure of the cell according to the invention. In Fig. 2, (1) is a positive electrode, and (2) is a negative electrode. (3) and (3') are current collectors, and each electrode is formed on each current collector. Lead terminals (10) and (10') are connected to the respective current collectors so that voltage drop does not arise, and the other ends are connected to a cell casing (6) and a top cap (9). (5) is a separator impregnated with an electrolytic solution, and in the electrolytic solution, an aforesaid compound capable of forming ions with which doping is made is dissolved in an aprotic organic solvent. The electrolytic solution is usually liquid, and impregnated into the separator, but it can also be used, without any separator, after being made a gel or a solid for preventing leakage of the solution. (8) is an insulating packing disposed for preventing contact between both positive and negative electrodes (the cell casing and the top cap).

The separator is made of a porous material which is durable against the electrolytic solution, the electrode active substance, etc., has open pores and is electrically non-conductive. It can usually be a cloth, a non-woven cloth, a porous material, or the like composed of glass fiber, polyethylene, polypropylene or the like. To decrease the internal resistance of the cell, the separator is preferably as thin as possible. Its thickness, however, is determined by considering the

amount of the electrolytic solution held, its permeability, its strength, etc. The positive and negative electrodes and the separator are fixed in position within a cell casing (6) in a manner not to give rise to any
5 problem in use. The shape, size, etc. of the electrodes can properly be determined according to the shape and performance of the desired cell.

The shape of the cell of the invention is not limited to the cylindrical shape as exemplified above,
10 but can also be coin-type, rectangular-type, box-type, etc., and is not particularly limited.

(5) The fifth preferred embodiment of the invention is described below.

In the invention, in the case of preparing an
15 organic electrolytic cell equipped, as stated already, with a positive electrode, a negative electrode and a solution of a lithium salt in an aprotic organic solvent as an electrolytic solution, wherein

(1) the positive electrode contains a metallic
20 oxide,

(2) the negative electrode is an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, said substrate being a heat-treated
25 product of an aromatic condensation polymer, and

(3) the total amount of lithium contained in the cell is 500 mAh/g or more, and the amount of lithium originating in the negative electrode is 100 mAh/g or more, based on the negative electrode PAS,

30 it is preferred to electrochemically carry lithium originating in the negative electrode on the negative electrode by applying an electric potential equal to or less than the electric potential of metallic Li to the negative electrode.

35 In the invention, as stated already, the total amount of lithium contained in the cell is the total of

lithium originating in the positive electrode, lithium originating in the electrolytic solution, and lithium originating in the negative electrode.

Lithium originating in the positive electrode
5 is lithium contained in the positive electrode at the time of assembly of the cell, and part or all of the lithium is supplied to the negative electrode by an operation (charge or the like) of sending a current from an outer circuit. Lithium originating in the electro-
10 lytic solution is lithium in the electrolytic solution contained in the separator, the positive electrode, the negative electrode, etc. Further, lithium originating in the negative electrode is lithium carried on the negative electrode PAS of the invention (lithium other
15 than lithium originating in the positive electrode and lithium originating in the electrolytic solution).

In the invention, as to a method of carrying lithium on the negative electrode PAS, it is possible to previously carry lithium on the negative electrode PAS
20 by, before assembly of the cell, passing a constant current or applying a constant voltage or using a combination thereof, in an electrochemical cell wherein metallic lithium is used as a counter electrode.

In the invention, in previously carrying
25 lithium on the negative electrode PAS, it is particularly preferred to apply an electric potential equal to or less than the electric potential of metallic lithium, at least once, to the negative electrode PAS, based on the electric potential of metallic lithium. The voltage to
30 be applied is varied depending on the desired amount of lithium originating in the negative electrode, PAS, the kind and shape of the electrodes, and the kind and shape of the electrolytic cell, but is preferably 0 mV to -1,000 mV, more preferably -10 mV to -300 mV based on
35 the electric potential of metallic lithium. An important thing is to select such a voltage as low as possi-

ble that metallic lithium does not electrodeposit, and in some case, it is possible to carry lithium first at an electric potential equal to or less than the electric potential of metallic lithium, gradually increase the voltage, and finally complete the carrying at a positive electric potential, or it is also possible to conduct the carrying first at a positive electric potential and then at an electric potential equal to or less than the electric potential of metallic lithium.

10 The total amount of lithium contained in the cell is 500 mAh/g or more, preferably 600 mAh/g or more, based on the negative electrode PAS, and in the case of under 500 mAh/g, sufficient capacity cannot be obtained.

15 Further, the amount of lithium originating in the negative electrode in the invention is 100 mAh/g or more, preferably 150 mAh/g or more, based on the negative electrode PAS, and in the case of under 100 mAh/g, adequate capacity cannot be obtained even if the total lithium amount is 500 mAh/g or more based on the negative electrode PAS. Further, when a lithium-containing oxide is used in the positive electrode, it is practical to make the amount of lithium originating in the negative electrode 600 mAh/g or less based on the negative electrode PAS.

25 Although as to the amount of lithium originating in the positive electrode and the amount of lithium originating in the electrolytic solution, it is sufficient if the above conditions are satisfied, it is preferred that the amount of lithium originating in the positive electrode is 300 mAh/g or more based on the negative electrode PAS.

(6) Further, the sixth preferred embodiment of the invention is described below.

35 In the invention, in previously carrying, as stated in the above (4), lithium originating in the negative electrode on PAS before assembly of the cell,

it is particularly preferred to electrochemically carry lithium originating in the negative electrode using a solution of a lithium salt in a cyclic carbonate solvent.

5 In this connection, the lithium salt is an electrolyte capable of forming lithium ions such as, for example, LiClO_4 , LiAsF_6 , LiBF_4 and LiPF_6 , and as the cyclic carbonate solvent, there can be used a single solvent such as ethylene carbonate or propylene carbonate, or a mixed solvent of two or more thereof. The
10 above electrolyte and solvent are mixed in a state of sufficient dehydration to give an electrolytic solution, and for making the internal resistance due to the electrolytic solution small, it is preferred to make the
15 concentration of the electrolyte in the electrolytic solution at least 0.1 mole/liter, and usually, it is further preferred to make the concentration 0.2 to 1.5 moles/liter. A method for previously carrying lithium on the negative electrode PAS is not particularly limited
20 so long as thereby it is possible to accomplish the purpose, and for example, it is possible to previously carry lithium on the negative electrode PAS by using the electrolytic solution, and passing a constant current or applying a constant voltage in an electrochemical cell
25 wherein metallic lithium is used as a counter electrode.

 According to the invention described above, there is provided an organic electrolytic cell which is easy to prepare, and is usable as a secondary cell having high capacity and high voltage.

30 Further, the organic electrolytic cell of the invention can be such a secondary cell that charge and discharge is possible over a long period and it is excellent in safety.

 A secondary cell having particularly high
35 capacity is provided by combining one or more of the above-mentioned first, second, third, fourth, fifth and

sixth preferred embodiments of the invention.

Further, the third and fourth preferred embodiments of the invention are advantageous because thereby are provided secondary cells having low internal resistance and high capacity.

Further, the above fifth preferred embodiment of the invention has an advantage of making it possible to prepare the above secondary cell having high capacity and high voltage more easily.

The invention is described below by examples. However, the following examples are for specifically describing the embodiments of the invention, and the invention is not limited to the following examples at all, or is not restricted thereby at all.

The basic structure of the cell according to the invention shown in the attached drawing Fig. 1 is as follows.

- 1 Positive electrode
- 2 Negative electrode
- 20 3, 3' Current collector
- 4 Electrolytic solution
- 5 Separator
- 6 Cell casing
- 7, 7' Outside terminals

Further, the basal constitution of the cell according to the invention shown in the attached drawing Fig. 2 is as follows.

- 1 Positive electrode
- 2 Negative electrode
- 30 3 Current collector (positive electrode)
- 3' Current collector (negative electrode)
- 8 Insulating packing
- 5 Separator
- 6 Cell casing
- 35 9 Top cap
- 10 Terminal (positive electrode)

10' Terminal (negative electrode)

Example 1

A phenol-formaldehyde resin forming plate 0.5 mm thick was put in a silicon carbide heating elements, the inside temperature was increased at a rate of 10°C /hour in an atmosphere of nitrogen, and heat treatment was conducted up to 650°C to synthesize an infusible, insoluble substrate (referred to as PAS). The resultant PAS plate was ground using a disc mill to give PAS powder having an average particle size of 15 μm . Its H/C ratio was 0.22.

Then, 100 weight parts of the PAS powder, and 100 weight parts of a solution of 10 weight parts of polyvinylidene fluoride powder in 90 weight parts of N,N-dimethylformamide were adequately mixed to give a slurry. The slurry was applied onto copper foil 10 μm thick (a current collector for the negative electrode) using an applicator, heat-dried, and pressed to give a PAS negative electrode 110 μm thick.

50 weight parts of a solution of 10 weight parts of polyvinylidene fluoride powder in 90 weight parts of N,N-dimethylformamide was adequately mixed with 100 parts of commercially available LiCoO_2 (made by Strem Chemicals Co.) and 5 parts of graphite to give a slurry. The slurry was applied onto aluminum foil 20 μm thick (a current collector for the positive electrode) using an applicator, dried, and pressed to give a PAS positive electrode 1 having a thickness of 150 μm .

The above negative electrode was doped with lithium in amounts of 150 mAh/g, 200 mAh/g and 300 mAh/g based on the negative electrode PAS at a constant current (a current was sent whereby 30 mAh/g of lithium could be carried per hour on the negative electrode PAS), using lithium as a counter electrode and using a solution of LiPF_6 at a concentration of 1 mole/liter in

a 1 : 1 (weight ratio) mixed solvent of propylene carbonate and diethyl carbonate as an electrolytic solution, whereby the lithium was carried thereon (lithium originating in the negative electrode). The resultant
5 negative electrodes were designated Negative electrodes 1, 2 and 3, respectively.

Three kinds of cells as shown in Fig. 1 were assembled using the above positive electrode and Negative electrodes 1, 2 and 3 (each $1 \times 1 \text{ cm}^2$). As the
10 separator, a polypropylene separator $25 \text{ }\mu\text{m}$ thick was used. Further, as the electrolytic solution was used a solution of LiPF_6 at a concentration of 1 mole/liter in a 1 : 1 (weight ratio) mixed solvent of propylene carbonate and diethyl carbonate. The total lithium amount
15 in the cell based on the negative electrode PAS is shown in Table 1.

Each of the above cells was charged at a constant current of 0.25 mA/cm^2 until the cell voltage became 4.3 V, and successively, discharged at a constant
20 current of 0.25 mA/cm^2 until the cell voltage became 2.5 V. This 4.3 V-2.5 V cycle was repeated, and in the third discharge, evaluation was made using volume capacity (mAh/cc). As the volume standard was used the total of the electrode volumes, the separator volume and the
25 current collector volumes. The results are shown in the following Table 1.

Table 1

5	Positive electrode	Negative electrode	Total lithium amount in the cell (mAh/g)	Lithium amount originating in the negative electrode (mAh/g)	Volume capacity (mAh/cc)
	1	1	890	150	131
	1	2	940	200	141
10	1	3	1040	300	155

Example 2

15 Positive electrodes having thicknesses of 100 μm and 200 μm were obtained in the same manner as in Example 1, and designated Positive electrode 2 and Positive electrode 3, respectively.

20 Cells were assembled in the same manner as in Example 1 using combinations of Positive electrode 2 with Negative electrode 2, and Positive electrode 3 with Negative electrode 3, and evaluation was made using volume capacity. The results are shown in the following Table 2.

Table 2

30	Positive electrode	Negative electrode	Total lithium amount in the cell (mAh/g)	Lithium amount originating in the negative electrode (mAh/g)	Volume capacity (mAh/cc)
	2	2	680	200	121
	3	2	1190	200	146

Comparative example 1

Positive electrode 4 having a thickness of 260 μm was obtained in the same manner as in Example 1. As the negative electrode was used Negative electrode 4 on which lithium was not previously carried. Cells were assembled in the same manner as in Example 1 using combinations between Positive electrodes 1, 3 or 4 and Negative electrode 4, and evaluation was made using volume capacity. The results are shown in the following Table 3.

Table 3

Positive electrode	Negative electrode	Total lithium amount in the cell (mAh/g)	Lithium amount originating in the negative electrode (mAh/g)	Volume capacity (mAh/cc)
1	4	740	0	95
3	4	990	0	104
4	4	1350	0	105

When the lithium amount originating in the negative electrode is 0, sufficient capacity was not obtained however larger the total lithium amount was made.

Comparative example 2

Negative electrode 5 was obtained in the same manner as in Example 1 except that the lithium amount to be previously carried was changed to 50 mAh/g. A cell was assembled in the same manner as in Example 1 using a combination between Positive electrode 1 and Negative electrode 5, and evaluation was made using volume capacity. The results are shown in the following Table 4.

Table 4

Positive electrode	Negative electrode	Total lithium amount in the cell (mAh/g)	Lithium amount originating in the negative electrode (mAh/g)	Volume capacity (mAh/cc)
1	5	790	50	103

When the lithium amount originating in the negative electrode (in this example, lithium with which the previous doping was made) was small, sufficient capacity was not obtained even if the total lithium amount in the cell is sufficient.

(1) Examples of the first preferred embodiment of the invention are set forth below.

Example 3

50 weight parts of a xylene resin (made by Lignte Co.), 50 weight parts of novolak (made by Showa Kobunshi Co. Ltd.) and 0.1 weight part of xylenesulfonic acid were heated at 100°C to give a xylene-modified novolak resin. 10 weight parts of hexamethylenetetramine was mixed with 100 weight parts of the resin, and the mixture was ground and then formed by thermal pressing into a formed plate.

The formed plate of the xylene-modified novolak resin was put in a silicon carbide heating elements, the inside temperature was increased at a rate of 10°C /hour in an atmosphere of nitrogen, and heat treatment was conducted up to 650°C to synthesize an infusible, insoluble substrate (referred to as PAS). The resultant PAS plate was ground using a disc mill to give PAS powder having an average particle size of 15 μm . Its H/C ratio was 0.22. The amount of gas adsorbed on the PAS fine particles at a nitrogen adsorption thickness of 10 Å was 29 cc/g.

Then, 100 weight parts of the PAS powder, and 100 weight parts of a solution of 10 weight parts of polyvinylidene fluoride powder in 90 weight parts of N,N-dimethylformamide were adequately mixed to give a
5 slurry. The slurry was applied onto copper foil 10 μm thick (a current collector for the negative electrode) using an applicator, dried, and pressed to give a PAS negative electrode 110 μm thick.

50 weight parts of a solution of 10 weight
10 parts of polyvinylidene fluoride powder in 90 weight parts of N,N-dimethylformamide was adequately mixed with 100 parts of commercially available LiCoO_2 (made by Strem Chemicals Co.) and 5 parts of graphite to give a
15 slurry. The slurry was applied onto aluminum foil 20 μm thick (a current collector for the positive electrode) using an applicator, dried, and pressed to give a PAS positive electrode 5 having a thickness of 165 μm .

The above negative electrode was doped with lithium in amounts of 150 mAh/g, 200 mAh/g and 300 mAh/g
20 based on the negative electrode PAS at a constant current (a current was sent whereby 30 mAh/g of lithium could be carried per hour on the negative electrode PAS), using lithium as a counter electrode and using a solution of LiPF_6 at a concentration of 1 mole/liter in
25 a 1 : 1 (weight ratio) mixed solvent of propylene carbonate and diethyl carbonate as an electrolytic solution, whereby the lithium was carried thereon (lithium originating in the negative electrode). The resultant negative electrodes were designated Negative electrodes
30 6, 7 and 8, respectively. Three kinds of cells as shown in Fig. 1 were assembled using the above positive electrode and Negative electrodes 6, 7 and 8 (each 1 x 1 cm^2). As the separator, a polypropylene separator 25 μm thick was used. Further, as the electrolytic solution
35 was used a solution of LiPF_6 at a concentration of 1 mole/liter in a 1 : 1 (weight ratio) mixed solvent of

propylene carbonate and diethyl carbonate. The total lithium amount in the cell based on the negative electrode PAS is shown in Table 5.

Each of the above cells was charged at a constant current of 0.25 mA/cm^2 until the cell voltage became 4.3 V, and successively, discharged at a constant current of 0.25 mA/cm^2 until the cell voltage became 2.5 V. This 4.3 V-2.5 V cycle was repeated, and in the third discharge, evaluation was made using volume capacity (mAh/cc). As the volume standard was used the total of the electrode volumes, the separator volume and the current collector volumes. The results are shown in the following Table 5.

Table 5

Negative electrode	Nitrogen adsorption amount at an adsorption thickness of 10Å (cc/g)	Total lithium amount in the cell (mAh/g)	Lithium amount originating in the negative electrode (mAh/g)	Volume capacity (mAh/cc)
6	29	990	150	151
7	29	1020	200	157
8	29	1130	300	171

Example 4

Negative electrodes were obtained in the same manner as in Example 3 except that the composition of the raw materials for PAS was changed to that of 30 weight parts of the xylene resin and 70 weight parts of the novolak, or that of 10 weight parts of the xylene resin and 90 weight parts of the novolak. The resultant negative electrodes were doped respectively with lithium in amounts of 300 mAh/g based on the negative electrode PAS, whereby the lithium was carried thereon to give Negative electrode 9 and Negative electrode 10, respectively.

Cells were assembled in the same manner as in Example 3, and evaluation was made using volume capacity. The results are shown in the following Table 6.

Table 6

Negative electrode	Nitrogen adsorption amount at an adsorption thickness of 10Å (cc/g)	Total lithium amount in the cell (mAh/g)	Lithium amount originating in the negative electrode (mAh/g)	Volume capacity (mAh/cc)
9	60	1130	300	162
10	83	1130	300	151

Comparative example 3

Two the same negative electrodes were obtained in the same manner as in Example 3 except that the composition of the raw materials for PAS was changed to that of 30 weight parts of the xylene resin and 70 weight parts of the novolak. As to one of the negative electrodes, lithium was not previously carried thereon, and the other negative electrode was doped with lithium in amounts of 50 mAh/g based on the negative electrode PAS, whereby the lithium was carried thereon. The resultant negative electrodes were designated Negative electrode 11 and Negative electrode 12, respectively.

Cells were assembled in the same manner as in Example 3, and evaluation was made using volume capacity. The results are shown in the following Table 7.

Table 7

Negative electrode	Nitrogen adsorption amount at an adsorption thickness of 10Å (cc/g)	Total lithium amount in the cell (mAh/g)	Lithium amount originating in the negative electrode (mAh/g)	Volume capacity (mAh/cc)
11	60	820	0	109
12	60	870	50	120

Comparative example 4

Negative electrodes were obtained in the same manner as in Example 3 except that the composition of the raw materials for PAS was changed to a composition of 100 weight parts of the novolak and 10 weight parts of hexamethylenetetramine, or such a composition that only powder resol ("ResiTop" made by Showa Kobunshi Co., Ltd.) was used as the raw material. These negative electrode were doped with lithium in an amount of 300 mAh/g based on the negative electrode PAS, whereby the lithium was carried thereon. The resultant negative electrodes were designated Negative electrode 13 and Negative electrode 14, respectively.

Cells were assembled in the same manner as in Example 3, and evaluation was made using volume capacity. The results are shown in the following Table 8.

Table 8

Negative electrode	Nitrogen adsorption amount at an adsorption thickness of 10Å (cc/g)	Total lithium amount in the cell (mAh/g)	Lithium amount originating in the negative electrode (mAh/g)	Volume capacity (mAh/cc)
13	120	1140	300	130
14	132	1120	300	131

(2) Examples of the second preferred embodiment of the invention are described below.

Example 5

A formed plate of phenol resin 0.5 mm thick
5 was put in a silicon carbide heating elements, the
inside temperature was increased at a rate of 10°C /hour
in an atmosphere of nitrogen, and heat treatment was
conducted up to 650°C to synthesize an infusible, insol-
uble substrate (referred to as PAS). Its H/C ratio was
10 0.22. The resultant PAS plate was ground in an
alumina-made pot mill with changed grinding times to
give PAS powders respectively having the particle size
distributions shown in Table 9 (No. 1, No. 2, No. 3 and
No. 4). The particle size distribution was measured by
15 dispersing the resultant fine particles into water using
ultrasonic waves and then using a laser diffraction-type
particle size distribution measuring apparatus.

Table 9

PAS fine particle No.	Electrode raw material fine particles					Electrode	
	Average particle size (μm)	50% size (μm)	2a _u	Particles having a particle size of a or less of a _u (%)	Particles having a particle size of 4a or more of 4a _u (%)	Negative electrode No.	Porosity (%)
1	10.2	9.5		26	14	15	38
2	6.0	4.3		28	28	16	33
3	3.6	2.4		31	25	17	29
4	2.6	1.6		23	27	18	30

Then, 100 weight parts of any one of the above PAS powders, and 100 weight parts of a solution of 10 weight parts of polyvinylidene fluoride powder in 90 weight parts of N,N-dimethylformamide were adequately
5 mixed to give a slurry. The slurry was applied onto copper foil 10 μm thick (a current collector for the negative electrode) using an applicator, dried, and pressed to give a PAS negative electrode 110 μm thick. The porosity of the negative electrode was determined by
10 impregnating the negative electrode with propylene carbonate at 25°C. The density of the used propylene carbonate was 1.20 g/cc (measured by a pycnometer). The results are shown together in Table 9. 50 weight parts of a solution of 10 weight parts of polyvinylidene
15 fluoride powder in 90 weight parts of N,N-dimethylformamide was adequately mixed with 100 parts of commercially available LiCoO_2 (made by Strem Chemicals Co.) and 5 parts of graphite to give a slurry. The slurry was applied onto aluminum foil 20 μm thick (a current
20 collector for the positive electrode) using an applicator, dried, and pressed. Thereby, Positive electrodes 6 and 7 were obtained having thicknesses of 160 μm and 180 μm , respectively.

Each of the above negative electrodes was
25 doped with lithium in an amount of 300 mAh/g based on the negative electrode PAS at a constant current (a current was sent whereby 30 mAh/g of lithium could be carried per hour on the negative electrode PAS), using lithium as a counter electrode and using a solution of
30 LiPF_6 at a concentration of 1 mole/liter in a 1 : 1 (weight ratio) mixed solvent of propylene carbonate and diethyl carbonate as an electrolytic solution, whereby the lithium was carried thereon (lithium originating in the negative electrode).

35 The above positive electrodes 6 and 7 were combined with the negative electrodes 15, 16, 17 and 18

(each $1 \times 1 \text{ cm}^2$) so that the total lithium amount in the cell became about 1,100 mAh/g based on the negative electrode PAS, and five kinds of cells as shown in Fig. 1 were assembled. As the separator, a polypropylene separator 25 μm thick was used. Further, as the electrolytic solution was used a solution of LiPF_6 at a concentration of 1 mole/liter in a 1 : 1 (weight ratio) mixed solvent of propylene carbonate and diethyl carbonate. The total lithium amount in the cell based on the negative electrode PAS is shown in Table 10.

Each of the above cells was charged at a constant current of 0.25 mA/cm^2 until the cell voltage became 4.3 V, and successively, discharged at a constant electric current of 0.25 mA/cm^2 until the cell voltage became 2.5 V. This 4.3 V-2.5 V cycle was repeated, and in the third discharge, evaluation was made using volume capacity (mAh/cc). As the volume standard was used the total of the electrode volumes, the separator volume and the current collector volumes. The results are also shown in Table 10.

Table 10

Negative electrode	Positive electrode	Total lithium amount in the cell (mAh/g)	Lithium amount originating in the negative electrode (mAh/g)	Volume capacity (mAh/cc)
15	6	1100	300	155
16	7	1170	300	173
17	7	1105	300	173
18	7	1120	300	168

Example 6

Positive electrodes 8, 9 and 10 having thicknesses of 240 μm , 210 μm and 200 μm , respectively, were obtained in the same manner as in Example 5. The amounts of lithium originating in the negative electrode of Negative electrode 17 were made to be 0 mAh/g (comparative), 150 mAh/g and 200 mAh/g (the invention), and the resultant negative electrodes were combined with the above positive electrodes, in the same manner as in Example 5, and cells were assembled and evaluated using volume capacity, in the same manner as in Example 5. The results are shown in Table 11.

Table 11

Negative electrode	Positive electrode	Total lithium amount in the cell (mAh/g)	Lithium amount originating in the negative electrode (mAh/g)	Volume capacity (mAh/cc)
17	8	1080	0	119
17	9	1095	150	148
17	10	1100	200	161

When the lithium amount originating in the negative electrode is 0, sufficient capacity was not obtained.

Comparative example 5

A formed plate of phenol resin 0.5 mm thick was put in a silicon carbide heating elements, the inside temperature was increased at a rate of 10°C /hour in an atmosphere of nitrogen, and heat treatment was conducted up to 650°C to synthesize a PAS. Its H/C ratio was 0.22. The resultant PAS plate was ground

using a jet mill to give PAS powders respectively having the particle size distributions shown in Table 4 (No. 5, No. 6 and No. 7). Negative electrodes (No. 19, No. 20 and No. 21) were obtained and porosities of them were
5 determined, in the same manner as in Example 5. The results are shown in Table 12.

Table 12

PAS fine particle No.	Electrode raw material fine particles				Electrode	
	Average particle size (μm)	50% size (μm) 2a	Particles having a particle size of a or less (%)	Particles having a particle size of 4a or more (%)	Negative electrode No.	Porosity (%)
5	22	21	24	22	19	38
6	8.5	7.7	4	6	20	46
7	2.5	2.1	12	9	21	45

Positive electrodes 11 and 12 having thicknesses of 150 μm and 130 μm , respectively, were obtained in the same manner as in Example 5. Thereafter, 300 mAh/g of lithium originating in the negative electrode 5 was carried on each of the negative electrodes of the above No. 19 to 21 in the same manner as in Example 5, and evaluation was conducted in the same manner as in Example 5. The results are shown in Table 13.

10

Table 13

15	Negative electrode	Positive electrode	Total lithium amount in the cell (mAh/g)	Lithium amount originating in the negative electrode (mAh/g)	Volume capacity (mAh/cc)
	19	11	1090	300	116
	20	12	1120	300	124
	21	12	1100	300	126

20

Comparative example 6

PAS No. 2 of Example 5 was used, and 100 weight parts of the PAS powder was adequately mixed with 110 weight parts of a solution of 10 weight parts of polyvinylidene fluoride powder in 90 weight parts of N,N-dimethylformamide to give a slurry. The slurry was applied onto copper foil 10 μm thick (a current collector for the negative electrode) using an applicator, and dried to give a PAS negative electrode 110 μm thick. The porosity of the negative electrode was determined by impregnating it with propylene carbonate at 25°C, and was 46 %.

35

A positive electrode 130 μm thick was obtained in the same manner as in Example 1. Thereafter, 300

mAh/g of lithium originating in the negative electrode was carried on the negative electrode in the same manner as in Example 5, and evaluation was conducted in the same manner as in Example 5. The volume capacity of the resultant cell was 131 mAh/g.

(3) An examples of the third preferred embodiment of the invention is described below.

Example 7

A formed plate of phenol resin 0.5 mm thick was put in a silicon carbide heating elements, the inside temperature was increased at a rate of 10°C /hour in an atmosphere of nitrogen, and heat treatment was conducted up to 650°C to synthesize an infusible, insoluble substrate (referred to as PAS). The resultant PAS plate was ground in a pot mill to give PAS powder having an average particle size of about 3 μ m. Its H/C ratio was 0.22.

Then, 100 weight parts of the above PAS powder, and 100 weight parts of a solution of 10 weight parts of polyvinylidene fluoride powder having a melting point of 172°C in 90 weight parts of N,N-dimethylformamide were adequately mixed to give a slurry. The slurry was applied onto copper foil 10 μ m thick (a current collector for the negative electrode) using an applicator, dried, and pressed to give a PAS negative electrode 210 μ m thick wherein both sides of the foil were coated with PAS. The PAS negative electrode was heat treated in vacuo at 100°C, 160°C, 190°C or 220°C to give Negative electrodes 22, 23, 24 and 25.

50 weight parts of a solution of 10 weight parts of polyvinylidene fluoride powder in 90 weight parts of N,N-dimethylformamide was adequately mixed with 100 parts of commercially available LiCoO_2 (made by Strem Chemicals Co.) and 5 parts of graphite to give a slurry. The slurry was applied onto aluminum foil 20 μ m thick (a current collector for the positive electrode)

using an applicator, dried, and pressed. Thereby was obtained a positive electrode 340 μm thick wherein both sides of the aluminum foil were coated with LiCoO_2 .

Each of the above negative electrodes was
5 doped with lithium in an amount of 300 mAh/g based on the negative electrode PAS at a constant current (a current was sent whereby 30 mAh/g of lithium could be carried per hour on the negative electrode PAS), using lithium as a counter electrode and using a solution of
10 LiPF_6 at a concentration of 1 mole/liter in a 1 : 1 (weight ratio) mixed solvent of propylene carbonate and diethyl carbonate as an electrolytic solution, whereby the lithium was carried thereon (lithium originating in the negative electrode). The resultant negative elec-
15 trodes were designated Negative electrodes 22, 23, 24 and 25.

A cylindrical cell as shown in Fig. 2 was assembled using the above positive electrode 1, and any one of Negative electrodes 22, 23, 24 and 25 (each 4 x
20 35 cm^2). As the separator, a polypropylene separator 25 μm thick was used. As the positive terminal was used an aluminum terminal 150 μm thick and 5 mm wide, and as the negative terminal was used a nickel terminal having the same size as the aluminum terminal, and these terminals
25 were attached to the ends of both electrodes, respectively. Further, as the electrolytic solution was used a solution of LiPF_6 at a concentration of 1 mole/liter in a 1 : 1 (weight ratio) mixed solvent of propylene carbonate and diethyl carbonate. The total lithium
30 amount in the cell based on the negative electrode PAS was 1,170 mAh/g on each cell. In this connection, when Negative electrode 1 of 100°C treatment was used, the electrode peeled off from the metallic foil at the time of winding of the electrode, and a cell could not be
35 obtained.

Each of the resultant cells was charged at a

constant current of 0.25 mA/cm^2 until the cell voltage became 4.3 V, and the internal resistance thereof was measured, and then, successively, the cell was discharged at a constant current of 0.25 mA/cm^2 until the cell voltage became 2.5 V. This 4.3 V-2.5 V cycle was repeated, and in the third discharge, evaluation was made using volume capacity (mAh/cc). As the volume standard was used the total of the electrode volumes, the separator volumes and the current collector volumes. The results are also shown in Table 14.

Table 14

Negative electrode No.	Heat treatment (°C)	Internal resistance (mΩ)	Volume capacity (mAh/cc)
22	100	Peeling of the electrode at the time of winding	
23	160	230	132
24	190	120	168
25	220	120	164

25

In the cases of the negative electrodes 24 and 25 which are preferred examples of the invention, the internal resistances are lower and the volume capacities are higher, compared with the cases of the negative electrodes 22 and 23.

30

Comparative example 7

Positive electrode 2 having a thickness of 460 μm was obtained in the same manner as in Example 7. The sizes of the positive electrode and negative electrode were made to be equally $4 \times 30 \text{ cm}^2$.

35

The amount of lithium originating in any of

the negative electrodes was made to be 0 mAh/g, and the above positive electrode was combined with any one of the negative electrodes 22, 23, 24 and 25. Then, cells were assembled and their evaluation was made by their volume capacities, in the same manner as in Example 7. The total lithium amount in each cell based on the negative electrode PAS was 1,090 mAh/g. The results are shown in Table 15.

Table 15

Negative electrode No.	Heat treatment (°C)	Internal resistance (mΩ)	Volume capacity (mAh/cc)
22	100	230	99
23	160	225	101
24	190	160	116
25	220	155	118

(4) Examples of the fourth preferred embodiment of the invention are described below.

Example 8

A formed plate of phenol resin 0.5 mm thick was put in a silicon carbide heating elements, the inside temperature was increased at a rate of 10°C /hour in an atmosphere of nitrogen, and heat treatment was conducted up to 650°C to synthesize an infusible, insoluble substrate (referred to as PAS). The resultant PAS plate was ground in a disc mill to give PAS powder having an average particle size of about 15 μm. Its H/C ratio was 0.22.

Then, 100 weight parts of the above PAS powder, and 100 weight parts of a solution of 10 weight

parts of polyvinylidene fluoride powder in 90 weight parts of N,N-dimethylformamide were adequately mixed to give a slurry. The slurry was applied onto copper foil 10 μm thick (a current collector for the negative electrode) using an applicator, dried, and pressed to give a PAS negative electrode 210 μm thick wherein both sides of the foil were coated with PAS.

50 weight parts of a solution of 10 weight parts of polyvinylidene fluoride powder in 90 weight parts of N,N-dimethylformamide was adequately mixed with 100 parts of commercially available LiCoO_2 (made by Strem Chemicals Co.) and 5 parts of graphite to give a slurry. The slurry was applied onto aluminum foil 20 μm thick (a current collector for the positive electrode) using an applicator, dried, and pressed. Thereby was obtained a positive electrode 280 μm wherein both sides of the foil were coated with LiCoO_2 .

The above negative electrode was doped with lithium in an amount of 300 mAh/g based on the negative electrode PAS at a constant current (a current was sent whereby 30 mAh/g of lithium could be carried per hour on the negative electrode PAS), using lithium as a counter electrode and using a solution of LiPF_6 at a concentration of 1 mole/liter in a 1 : 1 (weight ratio) mixed solvent of propylene carbonate and diethyl carbonate as an electrolytic solution, whereby the lithium was carried thereon (lithium originating in the negative electrode).

A cylindrical cell as shown in Fig. 2 was assembled using the above positive electrode and negative electrode (each $4 \times 35 \text{ cm}^2$). As the separator, a polypropylene separator 25 μm thick was used. As the positive terminal was used an aluminum terminal 150 μm thick and 5 mm wide, and as the negative terminal was used a nickel terminal having the same size as the aluminum terminal, and these terminals were attached to

the ends of both electrodes, respectively. Further, as the electrolytic solution was used a solution of LiPF_6 at a concentration of 1 mole/liter in a 1 : 1 (weight ratio) mixed solvent of propylene carbonate and diethyl carbonate. The total lithium amount in the cell based on the negative electrode PAS was 1,040 mAh/g.

The resultant cell was charged at a constant current of 0.25 mA/cm^2 until the cell voltage became 4.3 V, and the internal resistance thereof was measured, and then, successively, the cell was discharged at a constant current of 0.25 mA/cm^2 until the cell voltage became 2.5 V. This 4.3 V-2.5 V cycle was repeated, and in the third discharge, evaluation was made using volume capacity (mAh/cc). As the volume standard was used the total of the electrode volumes, the separator volumes and the current collector volumes. The results are also shown in Table 16.

Comparative example 8

A positive electrode 380 μm thick was obtained in the same manner as in Example 8. The sizes of the positive electrode and negative electrode were made to be equally $4 \times 30 \text{ cm}^2$.

The amount of lithium originating in the negative electrode was made to be 0 mAh/g, a cell was assembled in the same manner as in Example 8, and its evaluation was made by volume capacity. The total lithium amount in the cell based on the negative electrode PAS was 1,010 mAh/g. The results are shown in Table 16.

Comparative example 9

In Example 8, metallic lithium in an amount of 300 mAh/g (about 12 μm) was stuck on the negative electrode PAS, and then, two cylindrical cells as in Example 8 were assembled. The cells were left alone at room temperature for 3 days, and when one of them was decomposed, the metallic lithium completely disappeared. Evaluation was made by volume capacity in the same

manner as in Example 8. The total lithium amount in this cell was 1,040 mAh/g. The results are shown in Table 16.

When the lithium amount originating in the negative electrode was 0, sufficient capacity could not be obtained, and when lithium originating in the negative electrode was carried in the cell, the internal resistance of the cell was increased and the cell capacity was lowered.

Example 9

In Example 8, metallic lithium (about 200 μm) was stuck on the negative electrode PAS, the resultant matter was held between polypropylene plates each 2mm thick, and lithium originating in the negative electrode was carried on the negative electrode PAS in the same electrolytic solution as in Example 1. When about 40 minutes later, the metallic lithium was peeled from the PAS negative electrode, the PAS negative electrode was doped with 300 mAh/g of lithium. Thereafter, a cylindrical cell as in Example 8 was assembled, and evaluation was made by volume capacity in the same manner as in Example 8. The total lithium amount in this cell was 1,040 mAh/g. The results are shown in Table 16.

Example 10

In Example 8, lithium originating in the negative electrode was carried on the negative electrode PAS by short-circuiting the counter electrode metallic lithium (about 200 μm) and the negative electrode PAS. Thereby, the negative electrode PAS could be doped with 300 mAh/g of lithium in about 35 minutes. Thereafter, a cylindrical cell as in Example 8 was assembled, and evaluation was made by volume capacity in the same manner as in Example 8. The total lithium amount in this cell was 1,040 mAh/g. The results are shown in Table 16.

Table 16

Negative electrode No.	Lithium originating in the negative electrode	Lithium amount originating in the negative electrode (mAh/g)	Volume capacity (mAh/cc)	Internal resistance (mΩ)
Example 8	Before assembly	300	151	160
Comparative example 8	Before assembly	0	98	200
Comparative example 9	After assembly	300	117	360
Example 9	Before assembly	300	149	150
Example 10	Before assembly	300	153	150

(5) An example of the fifth preferred embodiment of the invention is described below.

Example 11

A formed plate of phenol resin 0.5 mm thick
5 was put in a silicon carbide heating elements, the
inside temperature was increased at a rate of 10°C /hour
in an atmosphere of nitrogen, and heat treatment was
conducted up to 650°C to synthesize an infusible, insol-
uble substrate (referred to as PAS). . The resultant PAS
10 plate was ground in a disc mill to give PAS powder
having an average particle size of about 15 µm. Its H/C
ratio was 0.22.

Then, 100 weight parts of the above PAS pow-
der, and 100 weight parts of a solution of 10 weight
15 parts of polyvinylidene fluoride powder in 90 weight
parts of N,N-dimethylformamide were adequately mixed to
give a slurry. The slurry was applied onto copper foil
10 µm thick (a current collector for the negative elec-
trode) using an applicator, dried, and pressed to give a
20 PAS negative electrode 210 µm thick wherein both sides
of the foil were coated with PAS.

50 weight parts of a solution of 10 weight
parts of polyvinylidene fluoride powder in 90 weight
parts of N,N-dimethylformamide was adequately mixed with
25 100 parts of commercially available LiCoO₂ (made by
Strem Chemicals Co.) and 5 parts of graphite to give a
slurry. The slurry was applied onto aluminum foil 20 µm
thick (a current collector for the positive electrode)
using an applicator, dried, and pressed. Thereby was
30 obtained a positive electrode 280 µm thick wherein both
sides of the foil were coated with LiCoO₂.

An electrolytic cell having a lithium refer-
ence electrode was assembled, using the above negative
electrode, and lithium as a counter electrode, and
35 using, as the electrolytic solution, a solution of LiPF₆
at a concentration of 1 mole/liter in a 1 : 1 (weight

ratio) mixed solvent of propylene carbonate and diethyl carbonate. A constant voltage was applied so that the PAS negative electrode became +20 mV, 0 mV, -20 mV, -50 mV or -100 mV, based on the lithium reference electrode, and time needed for carrying 300 mAh/g of lithium originating in the negative electrode was measured. The results are shown in Table 17.

A cell as shown in Fig. 1 was assembled using the above positive electrode 1 and negative electrode (each $1 \times 1 \text{ cm}^2$). As the separator, a polypropylene separator 25 μm thick was used. The total lithium amount in the cell based on the negative electrode PAS was 1,040 mAh/g.

Each of the resultant cells was charged at a constant current of 0.25 mA/cm^2 until the cell voltage became 4.3 V, and then, successively, the cell was discharged at a constant current of 0.25 mA/cm^2 until the cell voltage became 2.5 V. This 4.3 V-2.5 V cycle was repeated, and in the third discharge, evaluation was made using volume capacity (mAh/cc). As the volume standard was used the total of the electrode volumes, the separator volumes and the current collector volumes. The results are also shown in Table 17.

Table 17

5	Voltage applied on the negative electrode lithium	Time needed for carrying lithium originating in the negative electrode (minute)	Volume capacity (mAh/cc)
	+20 mV Comparison	43 minutes (No deposition of lithium)	152
10	0 mV The invention	29 minutes (No deposition of lithium)	153
	-20 mV The invention	24 minutes (No deposition of lithium)	149
	-50 mV The invention	20 minutes (No deposition of lithium)	152
15	-100 mV The invention	16 minutes (No deposition of lithium)	150

20 Comparative example 10

In Example 11, metallic lithium (about 200 μm) was stuck on the negative electrode PAS, the resultant matter was held between polypropylene plates each 2mm thick, and lithium originating in the negative electrode was carried on the negative electrode PAS in the same electrolytic solution as in Example 1. When about 40 minutes later, the metallic lithium was peeled from the PAS negative electrode, the PAS negative electrode was doped with 300 mAh/g of lithium. It takes more time, compared with the case where a negative voltage was applied.

Comparative example 11

In Example 11, lithium originating in the negative electrode was carried on the negative electrode PAS by short-circuiting the counter electrode metallic lithium (about 200 μm) and the negative electrode PAS.

Thereby, the negative electrode PAS could be doped with 300 mAh/g of lithium in about 35 minutes. It takes more time, compared with the case where a negative voltage was applied.

5 Comparative example 12

A formed plate of phenol resin 0.5 mm thick was put in a silicon carbide heating elements, the inside temperature was increased at a rate of 10°C /hour in an atmosphere of nitrogen, and heat treatment was
10 conducted up to 1,000°C to give a carbonaceous material. The resultant PAS plate was ground in a disc mill to give powder of the carbonaceous material having an average particle size of about 13 μm. Its H/C ratio was 0.02.

15 The carbonaceous material was made into an electrode in the same manner as in Example 11, and lithium originating in the negative electrode was carried thereon in the same manner as in Example 11. In the case of +20 mV, time needed for the carrying was 50
20 minutes, and in the case of 0 mV, time needed for the carrying was 45 minutes, and when -20 mV, -50 mV and -100 mV were applied, respectively, metallic lithium was deposited on the negative electrode carbonaceous material. When the resultant negative electrodes were left
25 alone, the metallic lithium disappeared by about 30 hours after the start of the leaving alone, but these methods are not practical as methods for carrying lithium originating in the negative electrode.

Further, when a cell similar to that in Example
30 ple 11 was assembled using the negative electrode prepared by the application of +20 mV, and evaluated, a large amount of metallic lithium was deposited on the negative electrode after the cycle was conducted three times.

35 (6) Examples of the sixth preferred embodiment of the invention are described below.

Example 12

A formed plate of phenol resin 0.5 mm thick was put in a silicon carbide heating elements, the inside temperature was increased at a rate of 10°C /hour in an atmosphere of nitrogen, and heat treatment was conducted up to 650°C to synthesize an infusible, insoluble substrate (referred to as PAS). The resultant PAS plate was ground in a disc mill to give PAS powder having an average particle size of about 15 μm . Its H/C ratio was 0.22.

Then, 100 weight parts of the above PAS powder, and 100 weight parts of a solution of 10 weight parts of polyvinylidene fluoride powder in 90 weight parts of N,N-dimethylformamide were adequately mixed to give a slurry. The slurry was applied onto copper foil 10 μm thick (a current collector for the negative electrode) using an applicator, dried, and pressed to give a PAS negative electrode 210 μm thick wherein both sides of the foil were coated with PAS.

50 weight parts of a solution of 10 weight parts of polyvinylidene fluoride powder in 90 weight parts of N,N-dimethylformamide was adequately mixed with 100 parts of commercially available LiCoO_2 (made by Strem Chemicals Co.) and 5 parts of graphite to give a slurry. The slurry was applied onto aluminum foil 20 μm thick (a current collector for the positive electrode) using an applicator, dried, and pressed. Thereby was obtained a positive electrode 280 μm thick wherein both sides of the foil were coated with LiCoO_2 .

The above negative electrode was doped with lithium in an amount of 300 mAh/g based on the negative electrode PAS at a constant current (a current was sent whereby 30 mAh/g of lithium could be carried per hour on the negative electrode PAS), using lithium as a counter electrode and using a solution of LiPF_6 at a concentration of 1 mole/liter in propylene carbonate and diethyl

carbonate as an electrolytic solution, whereby the lithium was carried thereon (lithium originating in the negative electrode).

A cell as shown in Fig. 1 was assembled using the above positive electrode and negative electrode (each $1 \times 1 \text{ cm}^2$). As the separator, a polypropylene separator $25 \text{ }\mu\text{m}$ thick was used. Further, as the electrolytic solution was used a solution of LiPF_6 at a concentration of 1 mole/liter in a 1 : 1 (weight ratio) mixed solvent of propylene carbonate and diethyl carbonate. The total lithium amount in the cell based on the negative electrode PAS was 1,040 mAh/g.

The resultant cell was charged at a constant current of 0.25 mA/cm^2 until the cell voltage became 4.3 V, and then, successively, the cell was discharged at a constant electric current of 0.25 mA/cm^2 until the cell voltage became 2.5 V. This 4.3 V-2.5 V cycle was repeated, and when in the third discharge, evaluation was made using volume capacity (mAh/cc), it was 169 mAh/cc. As the volume standard was used the total of the electrode volumes, the separator volume and the current collector volumes.

Comparative example 13

A cell was assembled in the same manner as in Example 1 except that as to lithium originating in the negative electrode, a solution of LiPF_6 at a concentration of 1 mole/liter in a 1 : 1 (weight ratio) mixed solvent of propylene carbonate and diethyl carbonate was used, and evaluation was made using volume capacity, which was found to be 155 mAh/cc.

Example 13

A cell was assembled in the same manner as in Example 12 except that as to lithium originating in the negative electrode, a solution of LiPF_6 at a concentration of 1 mole/liter in a 1 : 1 (weight ratio) mixed solvent of propylene carbonate and diethyl carbonate was

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used, and evaluation was made using volume capacity, which was found to be 167 mAh/cc.

The volume capacity in this example is about 10 % higher than that in Comparative example 13.

THE EMBODIMENTS OF THE INVENTION IN WHICH AN EXCLUSIVE PROPERTY OR PRIVILEGE IS CLAIMED ARE DEFINED AS FOLLOWS:

1. An organic electrolytic cell comprising a positive electrode, a negative electrode and a solution of a lithium salt in an aprotic organic solvent as an electrolytic solution, wherein:

(1) the positive electrode contains a metallic oxide,

(2) the negative electrode is an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, the substrate being a heat-treated product of an aromatic condensation polymer, and

(3) the total amount of lithium contained in the cell is 500 mAh/g or more, and the amount of lithium originating in the negative electrode is 100 mAh/g or more, based on the infusible, insoluble substrate (PAS) having a polyacene type skeletal structure as the negative electrode.

2. The organic electrolytic cell according to claim 1 wherein the negative electrode is a form composed of the infusible, insoluble substrate having the polyacene type skeletal structure and a binder, and the binder is a fluorine-containing polymer having a fluorine/carbon atomic ratio of under 1.5 but 0.75 or more.

3. The organic electrolytic cell according to claim 2 wherein the fluorine-containing polymer is polyvinylidene fluoride.

4. The organic electrolytic cell according to claim 1 wherein a lithium-containing metallic oxide is used as the positive electrode.

5. An organic electrolytic cell comprising a positive electrode, a negative electrode and a solution of a lithium salt in an aprotic organic solvent as an electrolytic solution, wherein:

(1) the positive electrode contains a metallic oxide,

(2) the negative electrode is an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, the substrate being a heat-treated product of an aromatic condensation polymer, and

(3) the total amount of lithium contained in the cell is 500 mAh/g or more, and the amount of lithium originating in the negative electrode is 100 mAh/g or more, based on the negative electrode PAS,

the infusible, insoluble substrate (PAS) having a polyacene type skeletal structure being such that the amount of adsorbed gas at a nitrogen adsorption thickness of 10 Å obtained from the nitrogen adsorption isotherm is 100 cc/g or less.

6. An organic electrolytic cell comprising a positive electrode, a negative electrode and a solution of a lithium salt in an aprotic organic solvent as an electrolytic solution, wherein:

(1) the positive electrode contains a metallic oxide,

(2) the negative electrode is an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, the substrate being a heat-treated product of an aromatic condensation polymer, and

(3) the total amount of lithium contained in the cell is 500 mAh/g or more, and the amount of lithium originating in the negative electrode is 100 mAh/g or more, based on the negative electrode PAS,

the infusible, insoluble substrate (PAS) having a polyacene type skeletal structure being such that it has an average particle size of 20 µm or less, and when it is supposed that the 50 % size is 2a µm, the rate of particles having a particle size of 1a µm or

particles having a particle size of $4\mu\text{m}$ or more is 10 % or more by volume ratio, and the porosity of the negative electrode being 40 % or less.

7. An organic electrolytic cell comprising a positive electrode, a negative electrode and a solution of a lithium salt in an aprotic organic solvent as an electrolytic solution, wherein:

(1) the positive electrode contains a metallic oxide,

(2) the negative electrode is an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, the substrate being a heat-treated product of an aromatic condensation polymer, and

(3) the total amount of lithium contained in the cell is 500 mAh/g or more, and the amount of lithium originating in the negative electrode is 100 mAh/g or more, based on the negative electrode PAS,

the negative electrode being one obtained by forming the infusible, insoluble substrate (PAS) having a polyacene type skeletal structure on metallic foil using a thermoplastic binder, and then heating the resultant form at a temperature equal to or higher than the melting point of the thermoplastic binder

8. An organic electrolytic cell comprising a positive electrode, a negative electrode and a solution of a lithium salt in an aprotic organic solvent as an electrolytic solution, wherein:

(1) the positive electrode contains a metallic oxide,

(2) the negative electrode is an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, the substrate being a heat-treated product of an aromatic condensation polymer, and

(3) the total amount of lithium contained in

(3) the total amount of lithium contained in the cell is 500 mAh/g or more, and the amount of lithium originating in the negative electrode is 100 mAh/g or more, based on the negative electrode PAS,

lithium originating in the negative electrode being previously carried on PAS before assembly of the cell.

9. An organic electrolytic cell comprising a positive electrode, a negative electrode and a solution of a lithium salt in an aprotic organic solvent as an electrolytic solution, wherein:

(1) the positive electrode contains a metallic oxide,

(2) the negative electrode is an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, the substrate being a heat-treated product of an aromatic condensation polymer, and

(3) the total amount of lithium contained in the cell is 500 mAh/g or more, and the amount of lithium originating in the negative electrode is 100 mAh/g or more, based on the negative electrode PAS,

lithium originating in the negative electrode being electrochemically carried by applying an electric potential equal to or lower than the electric potential of metallic Li.

10. An organic electrolytic cell comprising a positive electrode, a negative electrode and a solution of a lithium salt in an aprotic organic solvent as an electrolytic solution, wherein:

(1) the positive electrode contains a metallic oxide,

(2) the negative electrode is an infusible, insoluble substrate (PAS) having a polyacene type skeletal structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05, the substrate being a heat-treated

product of an aromatic condensation polymer, and

(3) the total amount of lithium contained in the cell is 500 mAh/g or more, and the amount of lithium originating in the negative electrode is 100 mAh/g or more, based on the negative electrode PAS,

lithium originating in the negative electrode being electrochemically carried using a solution of a lithium salt in a cyclic carbonate solvent.

11. The organic electrolytic cell according to any of claims 3 to 10 wherein the negative electrode is formed from the infusible, insoluble substrate (PAS) and a binder; and the binder is a fluorine-containing polymer having a fluorine/carbon atomic ratio of under 1.5 but 0.75 or more.

12. The organic electrolytic cell according to any of claims 4 to 10 wherein the negative electrode is formed from the infusible, insoluble substrate (PAS) and a binder; and the binder is polyvinylidene fluoride.

13. The organic electrolytic cell according to any of claims 5 to 10 wherein the positive electrode is a lithium-containing metallic oxide.

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14. An organic electrolytic cell which comprises:

(i) a positive electrode comprising a lithium-containing metallic oxide capable of electrochemical doping and undoping of lithium, the positive electrode being electrically connected to a positive electrode current collector and a positive electrode lead terminal;

(ii) a negative electrode formed of a mixture of [a] fine particles of an infusible and insoluble heat-treated product of an aromatic condensation polymer, the heat-treated product having a polyacene skeleton structure and having a hydrogen/carbon atomic ratio of 0.5 to 0.05 and [b] a fluorine-containing thermoplastic polymer having a fluorine/carbon atomic ratio of 0.75 or more but under 1.5 as a binder, the negative electrode being electrically connected to a negative electrode current collector and a negative electrode lead terminal;

(iii) a solution of a lithium salt dissolved in an aprotic solvent as an electrolytic solution, the solution being liquid, gel or solid;

(iv) a separator for preventing contact of the positive and negative electrodes and for holding the electrolytic solution;

(v) a cell casing which contains therein the positive and negative electrodes, the electrolytic solution and the separator and which is electrically connected to the negative electrode lead terminal; and

(vi) a top cap being electrically insulated from the cell casing and being electrically connected to the positive electrode lead terminal;

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wherein the total amount of lithium contained in the cell originating in the positive and negative electrode and in the electrolytic solution is 500 mAh or more and the amount of lithium originating in the negative electrode is 100 mAh or more, each per gram of the infusible and insoluble heat treated product.

15. The organic electrolytic cell according to claim 14 wherein the fine particles of the infusible and insoluble heat-treated product has such a pore structure that the amount of nitrogen gas adsorbed at a liquid nitrogen temperature of 77°K at a thickness of 10 Å is 100 cc/g or less.

16. The organic electrolytic cell according to claim 14 or 15, wherein the fluorine-containing thermoplastic polymer has a fluorine/carbon atomic ratio of at least 0.75 but under 1.5 and is used in an amount of 2 to 50% by weight based on the infusible and insoluble heat-treated product.

17. The organic electrolytic cell according to any one of claims 14 to 16, wherein the amount of lithium originating in the negative electrode is 600 mAh or less per gram of the infusible and insoluble heat-treated product.

18. The organic electrolytic cell according to any one of claims 14 to 17, wherein the amount of lithium originating in the positive electrode is 300 mAh per gram of the infusible and insoluble heat-treated product.

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19. The organic electrolytic cell according to any one of claims 14 to 18, wherein the total amount of lithium contained in the cell is 600 to 1190 mAh and the amount of lithium originating in the negative electrode is 150 to 300 mAh, each per gram of the infusible and insoluble heat-treated product.

20. The organic electrolytic cell according to any one of claims 14 to 19, wherein the fine particles of the infusible and insoluble heat-treated product are derived from an aromatic condensation polymer of a phenol compound with an aldehyde and have an average particle size of 20 μm or less and a porosity of 40% or less.

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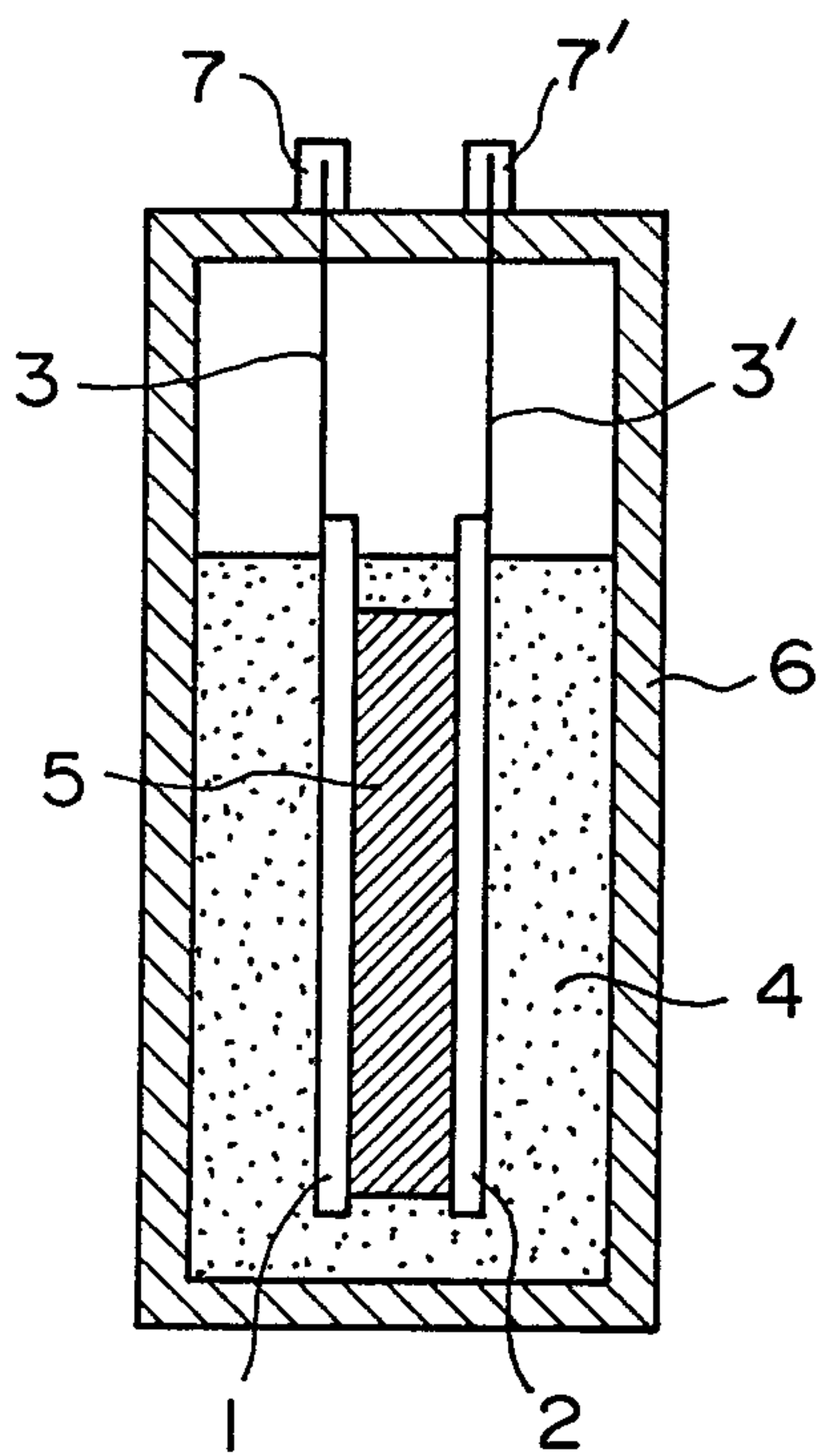


FIG. 1

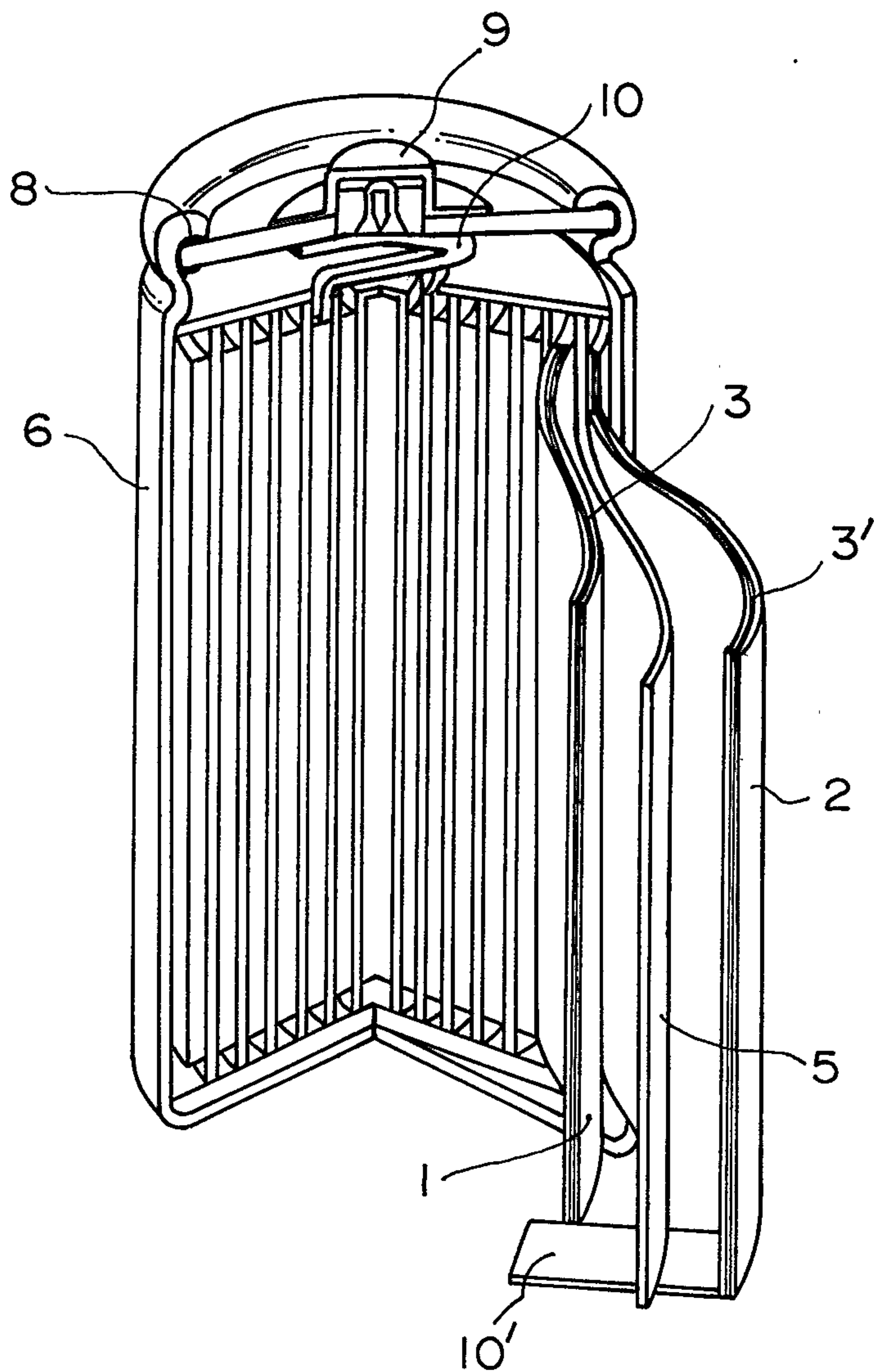


FIG. 2

