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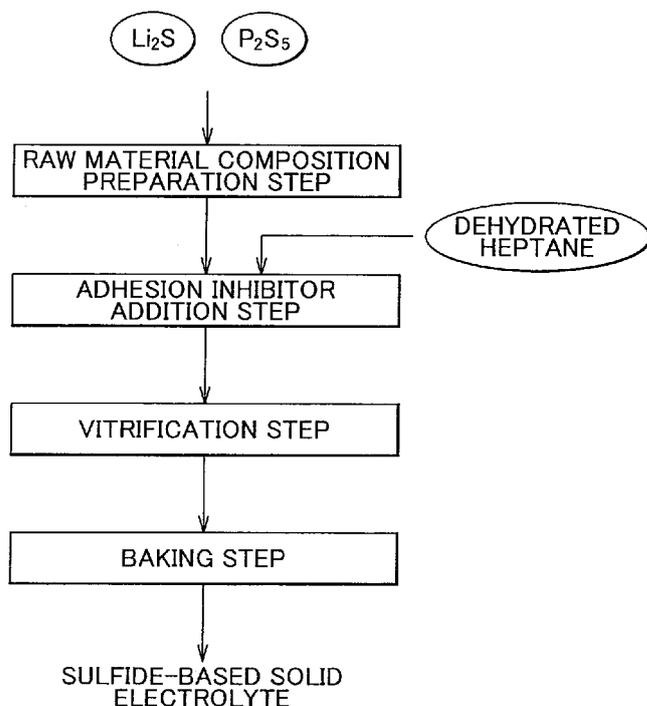
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(54) Title: PROCESS FOR PRODUCING SULFIDE-BASED SOLID ELECTROLYTE

FIG. 1



(57) Abstract: The invention provides a process for producing a sulfide-based solid electrolyte, the process having: a raw material composition preparation step for preparing a raw material composition containing at least sulfur (S), an adhesion inhibitor addition step for adding to the raw material composition an adhesion inhibitor that inhibits formation of adhered substances including the unreacted raw material composition, on the inner surface of a milling pot during mechanical milling, and a vitrification step for synthesizing sulfide-based glass by carrying out mechanical milling on the raw material composition to which the adhesion inhibitor has been added.

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PROCESS FOR PRODUCING SULFIDE-BASED SOLID ELECTROLYTE

BACKGROUND OF THE INVENTION

1. Field of the Invention

5 **[0001]** The invention relates to a process for producing a sulfide-based solid electrolyte useful as a solid electrolyte for, for example, all-solid-state lithium batteries, and more particularly, to a process for producing a sulfide-based solid electrolyte having superior production efficiency by eliminating the need to remove adhered substances formed on the inner surface of the milling pot during mechanical milling.

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2. Description of the Related Art

[0002] Accompanying the rapid proliferation of information-related devices and communication devices such as personal computers, video cameras and cell phones in recent years, increasing emphasis has been placed on the development of batteries for use
15 as power supplies of these devices. In addition, in the automobile industry as well, development is proceeding on high-output, high-capacity batteries for use in electric cars and hybrid vehicles. At present, among the various types of batteries available, lithium batteries are attracting attention from the viewpoint of their high energy density.

[0003] Since lithium batteries that are currently commercially available use
20 organic electrolytes that use a flammable organic solvent as a solvent, the attachment of a safety device for inhibiting temperature rises during a short-circuit as well as improvements are in terms of structure and materials to prevent short-circuits are required.

[0004] In contrast, since all-solid-state lithium batteries, which replace the
25 liquid electrolyte with a solid electrolyte to make the battery entirely solid state, do not use a flammable organic solvent therein, the safety device can be simplified thereby leading to reduced production costs and superior productivity. In addition, sulfide-based solid electrolytes are conventionally used for the solid electrolyte used in all-solid-state lithium batteries. Moreover, examples of processes used to produce sulfide-based solid

electrolytes include mechanical milling and rapid melt cooling.

[0005] Japanese Patent Application Publication No. 11-134937 (JP-A-11-134937), for example, discloses a process for producing ion-conducting sulfide glass by vitrifying a raw material of ion-conducting sulfide glass by mechanical milling.

5 Moreover, Japanese Patent Application Publication No. 2004-265685 (JP-A-2004-265685) and Japanese Patent Application Publication No. 2003-208919 (JP-A-2003-208919) disclose a process for producing lithium ion-conducting sulfide glass using mechanical milling. On the other hand, a process for producing a lithium ion-conducting solid electrolyte using rapid melt cooling is described in Japanese Patent
10 Application Publication No. 9-283156 (JP-A-9-283156).

[0006] In general, mechanical milling offers the advantage of allowing a desired sulfide glass to be more easily obtained in comparison with rapid melt cooling since it enables treatment to be carried out at normal temperatures. However, raw material compositions of sulfide-based glass are comparatively soft due to the presence of sulfur
15 (S) contained therein, and there was also the problem of adhered substances, including unreacted raw material composition, being formed on the inner surface of the milling pot during mechanical milling. Consequently, it was necessary to periodically remove adhered substances formed on the inner surface of the pot, thereby resulting in the problem of poor production efficiency.

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SUMMARY OF THE INVENTION

[0007] The invention provides a process for producing a sulfide-based solid electrolyte having superior production efficiency by eliminating the need to remove adhered substances formed on the inner surface of the milling pot during mechanical
25 milling.

[0008] In one aspect thereof, the invention relates to a process for producing a sulfide-based solid electrolyte including: a step of preparing a raw material composition containing at least sulfur (S); a step of adding to the raw material composition an adhesion inhibitor that inhibits formation of adhered substances, including the unreacted

raw material composition, on the inner surface of a milling pot during mechanical milling; and a step of synthesizing sulfide-based glass by carrying out mechanical milling on the raw material composition to which the adhesion inhibitor has been added.

[0009] According to this invention, the formation of adhered substances on the inner surface of the milling pot can be inhibited by adding an adhesion inhibitor of the raw material composition and carrying out mechanical milling. As a result, conventionally performed removal of adhered substances is no longer required, thereby making it possible to improve work efficiency. Moreover, the synthesis time of the sulfide-based glass can be shortened considerably since mechanical milling is carried out uniformly on the raw material composition. Moreover, sulfide-based glass having a homogeneous composition can be synthesized since mechanical milling is carried out uniformly.

[0010] The adhesion inhibitor is preferably a liquid having the property of not generating hydrogen sulfide in a reaction with the raw material composition. The use of such a liquid makes it possible to improve the dispersibility of the raw material composition in the milling pot, thereby enabling the formation of adhered substances to be inhibited.

[0011] The liquid is preferably an aprotic liquid. This is because an aprotic liquid can prevent the formation of hydrogen sulfide.

[0012] The aprotic liquid is preferably a non-polar aprotic liquid, and the non-polar aprotic liquid is more preferably an alkane that is a liquid at normal temperature (25°C). This is because such as alkane is able to effectively prevent formation of hydrogen sulfide.

[0013] The alkane that is a liquid at normal temperature (25°C) is preferably n-heptane. Because the SP value of the n-heptane is small (the polarity of the n-heptane is small), it is possible to effectively suppress a reaction between the n-heptane and a sulfide-based solid electrolyte (including the unreacted raw material composition), thereby suppressing deterioration of the sulfide-based solid electrolyte. Thus, it is possible to improve Li ion conductivity.

[0014] The adhesion inhibitor may also be a gel having the property of not generating hydrogen sulfide in a reaction with the raw material composition. The use of such a gel makes it possible to improve dispersibility of the raw material composition in the milling pot, thereby enabling the formation of adhered substances to be inhibited.

5 [0015] The raw material composition preferably further contains Li. This is because lithium allows the obtaining of a sulfide-based solid electrolyte useful as a solid electrolyte for an all-solid-state lithium battery, for example.

[0016] The raw material composition preferably contains at least Li_2S and P_2S_5 . This is because these compounds allow the obtaining of a sulfide-based solid electrolyte having superior Li ion conductivity.

10 [0017] The Li_2S and P_2S_5 preferably satisfy the relationship of $\text{Li}_2\text{S}:\text{P}_2\text{S}_5 = 70:30$ (molar ratio). This allows the obtaining of a sulfide-based solid electrolyte having even more superior Li ion conductivity.

[0018] The mechanical milling is preferably milling that uses a planetary ball mill. This is because the use of a planetary ball mill enables the raw material composition to be vitrified efficiently.

[0019] According to the invention, effects are demonstrated such as eliminating the need for removal of adhered substances, considerably shortening the synthesis time of sulfide-based glass, and allowing the obtaining of sulfide-based glass having a homogeneous composition.

BRIEF DESCRIPTION OF THE DRAWINGS

[0020] The features, advantages, and technical and industrial significance of this invention will be described in the following detailed description of example embodiments of the invention with reference to the accompanying drawings, in which like numerals denote like elements, and wherein:

FIG. 1 is a flow chart showing an example of the process for producing a sulfide-based solid electrolyte of the invention;

FIG. 2 is a schematic cross-sectional view for explaining adhered substances formed

during mechanical milling;

FIG. 3 shows measurement results of X-ray diffraction of glass powder obtained in Example 1;

FIG. 4 shows measurement results of X-ray diffraction of glass powder obtained in Comparative Example 1;

FIG. 5 shows measurement results of DSC on glass powder obtained in Example 1 and glass powder obtained in Comparative Example 1; and

FIG. 6 shows measurement results of Li ion conductivity of glass powder obtained in Examples 3 to 6.

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DETAILED DESCRIPTION OF EMBODIMENTS

[0021] FIG. 1 is a flow chart showing an example (embodiment) of a process for producing a sulfide-based solid electrolyte of the invention. In the process for producing a sulfide-based solid electrolyte shown in FIG. 1, lithium sulfide (Li_2S) and phosphorous pentasulfide (P_2S_5) are first prepared as raw materials followed by mixing at a prescribed ratio to prepare a raw material composition (raw material composition preparation step). Next, the raw material composition and grinding balls are placed in a mechanical milling pot followed by the further addition of dehydrated heptane (adhesion inhibitor) to the pot and sealing (adhesion inhibitor addition step). Next, by attaching the milling pot to a planetary ball mill and carrying out mechanical milling, the raw material composition is converted to an amorphous form to synthesize powdered sulfide-based glass (vitrification step). Next, the powdered sulfide-based glass is baked to synthesize powdered sulfide-based glass ceramic (baking step). This sulfide-based glass ceramic is the target sulfide-based solid electrolyte in FIG. 1.

[0022] Here, the sulfide-based solid electrolyte obtained according to this embodiment refers to a concept that includes both the sulfide-based glass ceramic obtained in the baking step described above, as well as the sulfide-based glass obtained in the vitrification step. Namely, the sulfide-based solid electrolyte obtained according to this embodiment may be the sulfide-based glass obtained in the vitrification step or the

sulfide-based glass ceramic obtained by baking the sulfide-based glass.

[0023] According to this embodiment, adhered substances can be inhibited from being formed on the inner surface of the milling pot by carrying out mechanical milling after adding an adhesion inhibitor to the raw material composition. As a result, the conventionally carried out removal of adhered substances is no longer necessary, thereby making it possible to improve work efficiency. Moreover, since mechanical milling is carried out uniformly on the raw material composition, time required to synthesize the sulfide-based glass can be shortened considerably. Moreover, since uniform mechanical milling is carried out, sulfide-based glass can be synthesized having a homogeneous composition.

[0024] FIG. 2 is a schematic cross-sectional view for explaining adhered substances formed during conventional mechanical milling. As shown in FIG. 2, since the raw material composition contains sulfur (S) and is comparatively soft, in the case of conventional mechanical milling, an adhered substance 2 containing unreacted raw material composition ends up forming on the inner surface of a milling pot 1. Consequently, in order to obtain sulfide-based glass having a homogeneous composition, it was necessary to scrape off the adhered substance 2 formed on the inner surface of the milling pot 1 resulting in the problem of poor production efficiency. This problem caused by adhered substances is unique to the case of using a soft raw material composition containing sulfur (S). In contrast, in this embodiment, the formation of adhered substances can be inhibited by adding an adhesion inhibitor. As a result, it is no longer necessary to remove the adhered substances, the time required to synthesize the sulfide-based glass can be shortened considerably, and sulfide-based glass can be obtained having a homogeneous composition as previously described. The following provides a detailed explanation of each step of the process for producing a sulfide-based solid electrolyte of this embodiment. Furthermore, each of the steps described to follow are normally carried out in an inert gas atmosphere (such as in an Ar gas atmosphere).

[0025] 1. Raw Material Composition Preparation Step

First, an explanation is provided of the raw material composition preparation step in

this embodiment. The raw material composition preparation step in this embodiment is a step in which a raw material composition is prepared that at least contains sulfur (S).

[0026] There are no particular limitations on the composition of the raw material composition provided it at least contains sulfur (S), and the composition thereof is preferably suitably selected depending on the intended use of the target sulfide-based solid electrolyte. For example, in the case of producing a sulfide-based solid electrolyte to be used in an all-solid-state lithium battery, the raw material composition preferably further contains Li in addition to S. This is done to obtain a sulfide-based solid electrolyte having superior Li ion conductivity. Moreover, the raw material composition preferably further contains a third component A in addition to the S and Li. Examples of the third component A include at least one type selected from the group consisting of P, Ge, B, Si, I, Al, Ga and As, with P being particularly preferable.

[0027] There are no particular limitations on the raw material compounds used in the raw material composition provided a raw material composition having a composition as described above can be obtained, and examples of such compounds include Li_2S , P_2S_5 , P_2S_3 , SiS_2 , AlS_2 , B_2S_2 , GeS_2 , Li, S, P, Ga_2S , As_2S_3 and Sb_2S_3 . In particular, the raw material composition preferably contains at least Li_2S and P_2S_5 . This is because a sulfide-based solid electrolyte can be obtained having superior Li ion conductivity. In this case, the raw material composition may contain only Li_2S and P_2S_5 , or it may contain other compounds in addition to the Li_2S and P_2S_5 .

[0028] There are no particular limitations on the contents of the compounds in the raw material composition, and the contents are preferably suitably selected according to the target composition of the raw material composition. For example, in the case the raw material composition contains Li_2S and P_2S_5 , the molar ratio of Li_2S to P_2S_5 is preferably 68:32 to 74:26 (molar ratio), and the ratio of $\text{Li}_2\text{S}:\text{P}_2\text{S}_5$ is particularly preferably 70:30 (molar ratio). This is because a sulfide-based solid electrolyte can be obtained having even more superior Li ion conductivity.

[0029] Furthermore, examples of compositions of the target sulfide-based solid electrolyte include $70\text{Li}_2\text{S}-30\text{P}_2\text{S}_5$, $80\text{Li}_2\text{S}-20\text{P}_2\text{S}_5$, $75\text{Li}_2\text{S}-25\text{P}_2\text{S}_5$, $\text{Li}_2\text{S}-\text{P}_2\text{S}_5$, $\text{Li}_2\text{S}-\text{SiS}_2$

and $\text{LiGe}_{0.25}\text{P}_{0.75}\text{S}_4$.

[0030] 2. Adhesion Inhibitor Addition Step

Next, an explanation is provided of the adhesion inhibitor addition step in this embodiment. The adhesion inhibitor addition step is a step in which an adhesion
5 inhibitor is added to the above-mentioned raw material composition for inhibiting the formation of adhered substances containing the unreacted raw material composition on the inner surfaces of a milling pot during mechanical milling.

[0031] There are no particular limitations on the adhesion inhibitor used in this embodiment provided it has the property of not generating hydrogen sulfide in a reaction
10 with the raw material composition. The adhesion inhibitor normally has the property of not generating hydrogen sulfide not only in a reaction with the raw material composition, but also in reactions with sulfide-based glass and precursors thereof. Examples of forms of the adhesion inhibitor include liquids, gels and solids.

[0032] (1) Case of a Liquid Adhesion Inhibitor

15 In this embodiment, the adhesion inhibitor is preferably a liquid having the property of not generating hydrogen sulfide in a reaction with the raw material composition. The use of such a liquid makes it possible to improve the dispersibility of the raw material composition in the milling pot, thereby making it possible to inhibit the formation of adhered substances. Furthermore, in the case of using such a liquid, it is suitable to use
20 so-called wet mechanical milling. Since wet mechanical milling typically has lower shearing force than dry mechanical milling, it has conventionally not been used for the production of solid electrolytes requiring adequate vitrification. In addition, suitably selecting the type of liquid in consideration of the boiling point of the liquid offers the advantage of being able to easily remove the adhesion inhibitor by drying only.

25 **[0033]** There are no particular limitations on the above-mentioned liquid provided it has a desired fluidity and has the property of not generating hydrogen sulfide in a reaction with the raw material composition. Hydrogen sulfide is generally generated as a result of protons that have dissociated from molecules of the liquid reacting with the raw material composition and sulfide-based glass. Consequently, the

liquid preferably has aprotic properties to a degree that does not result in the formation of hydrogen sulfide. Namely, the liquid is preferably an aprotic liquid. This is because an aprotic liquid is able to prevent the formation of hydrogen sulfide. In addition, aprotic liquids used in this embodiment can normally be divided into polar aprotic liquids
5 and non-polar aprotic liquids.

[0034] There are no particular limitations on the polar aprotic liquid, and examples include ketones such as acetone, nitriles such as acetonitrile, amides such as N,N-dimethylformamide (DMF) and sulfoxides such as dimethylsulfoxide (DMSO).

[0035] One example of a non-polar aprotic liquid is an alkane that is a liquid at
10 normal temperature (25°C). The alkane may be a chain alkane or a cyclic alkane. The number of carbon atoms of the alkane is preferably, for example, 5 or more. On the other hand, there are no particular limitations on the upper limit on the number of carbon atoms of the alkane provided it is a liquid at normal temperature. Specific examples of the alkane include pentane, hexane, heptane, octane, nonane, decane, undecane, dodecane
15 and paraffin. Furthermore, the chain alkane may have a branch. The chain alkane is preferably a straight-chain alkane. Specific examples of the straight-chain alkane include n-heptane and n-dacane, and the n-heptane is more preferable. Because the SP value of the n-heptane is small, it is possible to effectively suppress a reaction with between the n-haptane and a sulfide-based solid electrolyte (including the unreacted raw
20 material composition), thereby suppressing deterioration of the sulfide-based solid electrolyte. Thus, it is possible to improve Li ion conductivity. In addition, the number of carbon atoms of the cyclic alkane is preferably, for example, 5 or more and particularly preferably 6 or more. On the other hand, there are no particular limitations on the upper limit of the number of carbon atoms of the cyclic alkane provided it is a
25 liquid at normal temperature. Specific examples of the cyclic alkane include cyclopentane, cyclohexane, cycloheptane, cyclooctane and cycloparaffin.

[0036] In addition, other examples of non-polar aprotic liquids include aromatic hydrocarbons such as benzene, toluene or xylene, chain ethers such as diethyl ether or dimethyl ether, cyclic ethers such as tetrahydrofuran, alkyl halides such as chloroform,

methyl chloride or methylene chloride, esters such as ethyl acetate, and fluorine compounds such as fluorobenzene, fluoroheptane, 2,3-dihydroperfluoropentane or 1,1,2,2,3,3,4-heptafluorocyclopentane.

[0037] The polarity of the aprotic liquid can be evaluated using a solution
5 parameter (SP value). The aprotic liquid with lower polarity is more preferable. In
other words, the aprotic liquid with a smaller SP value (MJ/cm^3) is more preferable.
More specifically, the SP value of the aprotic liquid is preferably equal to or smaller than
18.5, and is more preferably equal to or smaller than 16. By using the aprotic liquid
with a low SP value, it is possible to effectively suppress a reaction between the aprotic
10 liquid and the sulfide-based solid electrolyte (including the unreacted raw material
composition), thereby suppressing deterioration of the sulfide-based solid electrolyte.
Thus, it is possible to improve Li ion conductivity.

[0038] The above-mentioned liquid is normally a liquid at normal temperature
(25°C). The boiling point of the liquid is preferably within the range of, for example, 60
15 to 300°C , and more preferably within the range of 80 to 200°C . If the boiling point of
the liquid is excessively low, it ends up vaporizing within the milling pot and may not be
able to inhibit the formation of adhered substances, while if the boiling point is
excessively high, it may be difficult to remove the liquid.

[0039] In addition, the above-mentioned liquid used in this embodiment
20 preferably has a low water content. This is because a low water content makes it
possible to inhibit the formation of hydrogen sulfide. The amount of water contained in
the liquid is preferably, for example, 100 ppm or less, and is more preferably 50 ppm or
less. An example of a method for lowering the water content is distillation. Namely,
the liquid is preferably that which has undergone distillation.

[0040] In this embodiment, there are no particular limitations on the amount of
25 the liquid added to the raw material composition provided the formation of adhered
substances is able to be inhibited. The amount of the liquid is preferably, for example,
50 parts by weight or more, more preferably 100 parts by weight or more, and even more
preferably 200 parts by weight or more, with respect to 100 parts by weight of the raw

material composition. If the amount of the liquid added is excessively low, there is the possibility of the formation of adhered substances not being inhibited. On the other hand, the amount of the liquid is preferably, for example, 1000 parts by weight or less, and more preferably 500 parts by weight or less, with respect to 100 parts by weight of the raw material composition. If the amount of the liquid added is excessively high, there is the possibility of synthesis of sulfide-based glass requiring an excessively long period of time.

[0041] (2) Case of a Gelatinous Adhesion Inhibitor

In this embodiment, the adhesion inhibitor may also be a gel having the property of not generating hydrogen sulfide in a reaction with the raw material composition. The use of such a gel makes it possible to improve dispersibility of the raw material composition in the milling pot, thereby enabling inhibition of the formation of adhered substances.

[0042] There are no particular limitations on the gel provided it has the property of not generating hydrogen sulfide in a reaction with the raw material composition. In addition, the amount of the gel added to the raw material composition is preferably suitably selected according to the amount and type of the raw material composition.

[0043] (3) Case of a Solid Adhesion Inhibitor

In this embodiment, the adhesion inhibitor may also be a solid having the property of not generating hydrogen sulfide in a reaction with the raw material composition. The use of such a solid makes it possible to improve dispersibility of the raw material composition in the milling pot, thereby enabling inhibition of the formation of adhered substances.

[0044] There are no particular limitations on the solid provided it has the property of not generating hydrogen sulfide in a reaction with the raw material composition. An example of the solid is a ball (adhesion inhibition ball) smaller than crushing balls used in mechanical milling. The addition of an adhesion inhibition ball makes it possible to reduced adhesion between the inner surface of the milling pot and adhered substances. An example of the material of such an adhesion inhibition ball is

the same material as that of crushing balls used in mechanical milling to be described later. In addition, although there are no particular limitations on the diameter of the adhesion inhibition ball provided it is smaller than the crushing balls, it is preferably, for example, 3 mm or less. In addition, the amount of the solid added to the raw material composition is preferably suitably selected according to the amount and type of raw material composition.

[0045] 3. Vitrification Step

Next, an explanation is provided of the vitrification step in this embodiment. The vitrification step is a step for synthesizing sulfide-based glass by carrying out mechanical milling on the raw material composition to which the adhesion inhibitor has been added as previously described. As was previously described, the sulfide-based glass obtained in the vitrification step may be used as a sulfide-based solid electrolyte.

[0046] There are no particular limitations on the type of mechanical milling apparatus used in this embodiment provided it causes vitrification of the raw material composition, and examples include ball mill apparatuses such as a planetary ball mill, rotary ball mill, agitating ball mill or vibration ball mill, and crushing machines using a ring. A planetary ball mill is particularly preferable. This is because a planetary ball mill is able to efficiently vitrify the raw material composition.

[0047] Various conditions for the above-mentioned mechanical milling are preferably set to a degree that allows the obtaining of a desired sulfide-based glass, and are preferably suitably selected according to the type of mechanical milling. For example, in the case of synthesizing sulfide-based glass with a planetary ball mill, normally the raw material composition and grinding balls are added to the milling pot followed by carrying out treatment for a prescribed rotational speed and time. In general, the formation rate of sulfide-based glass increases the higher the rotational speed, while the conversion rate of the raw materials to sulfide-based glass increases as the duration of treatment time becomes longer. The rotational speed of the planetary ball mill is, for example, within the range of 100 to 500 rpm, and preferably within the range of 200 to 400 rpm. In addition, the treatment time when using the planetary ball mill is

preferably an amount of time of a degree to which vitrification of the raw material composition is able to proceed adequately.

[0048] 4. Baking Step

Next, an explanation is provided of the baking step in this embodiment. The
5 baking step is a step for obtaining sulfide-based glass ceramic by carrying out baking treatment for improving crystallinity of sulfide-based glass. As was previously described, the sulfide-based glass ceramic obtained in the baking step may be used as a sulfide-based solid electrolyte.

[0049] There are no particular limitations on the temperature of baking
10 treatment provided it is a temperature that allows the obtaining of a desired sulfide-based glass ceramic, and for example, is within the range of 150 to 360°C and preferably within the range of 200 to 350°C. If the temperature of baking treatment is too low, the glass transition temperature of the sulfide-based glass may not be reached thereby preventing crystallization from proceeding, while if the temperature of baking treatment is too high,
15 a desired crystal structure may not be able to be formed. In addition, the duration of baking treatment is, for example, within the range of 1 minute to 10 hours, and preferably within the range of 0.5 to 3 hours.

[0050] 5. Other

The sulfide-based solid electrolyte obtained according to this embodiment is useful,
20 for example, as a solid electrolyte for an all-solid-state battery (and particularly an all-solid-state lithium battery). For example, a powder of the sulfide-based solid electrolyte can be compression molded for use as a solid electrolyte film. In addition, a method for producing an all-solid-state battery can also be provided that uses a sulfide-based solid electrolyte obtained according to the process for producing a
25 sulfide-based solid electrolyte described above.

[0051] This invention is not limited to the embodiments thus described. These embodiments are intended to exhibit examples. Anything that embodies configurations substantially equivalent to the technical ideas disclosed in the claims of this invention and provides operational advantage identical to this invention is within the technical scope of

the claims of this invention.

[0052] The following provides an explanation of specific examples.

[Example 1]

[0053] Lithium sulfide (Li_2S) crystals and phosphorous pentasulfide (P_2S_5) were
5 used as starting materials. 2 g of these powders were weighed at a molar ratio of 70/30
($\text{Li}_2\text{S}/\text{P}_2\text{S}_5$) in a dry box in the presence of an argon atmosphere and then placed in a 45
ml zirconia milling pot. Next, 4 g of dehydrated heptane were weighed and placed in
the milling pot followed by further placing zirconia grinding balls (diameter: 10 mm, 29
balls) in the milling pot and completely sealing the pot.

10 [0054] The milling pot was then attached to a planetary ball mill followed
initially by carrying out milling for several minutes at low speed (rotational speed: 60
rpm) to sufficiently mix the materials. Subsequently, the rotational speed was gradually
increased to carry out mechanical milling at 370 rpm for 5 hours, 9 hours and 15 hours,
respectively. After drying was performed, glass powder was obtained.

15 [Example 2]

[0055] Glass powder was obtained in the same manner as Example 1 with the
exception of changing the amount of raw material composition to 20 g, changing the
amount of dehydrated heptane to 40 g, using a 500 ml zirconia milling pot, and using
zirconia grinding balls (diameter: 5 mm, 600 g).

20 [Comparative Example 1]

[0056] Glass powder was obtained in the same manner as Example 1 with the
exception of not adding dehydrated heptane, changing durations of mechanical milling at
370 rpm to 20, 25 and 30 hours, respectively, and scraping off unreacted raw material
composition adhered to the surface of the milling pot every 5 hours.

25 [Evaluation 1]

(1) X-ray Diffraction Measurement

[0057] FIG. 3 shows measurement results of X-ray diffraction of the glass
powder obtained in Example 1. As shown in FIG. 3, although a peak was observed for
lithium sulfide (Li_2S) of the raw material in glass powder mechanically milled for 5 hours,

this peak corresponding to (Li_2S) of the raw material was no longer observed in glass powders mechanically milled for 9 and 15 hours, thereby confirming the adequate progression of vitrification. Similarly, vitrification was also confirmed to have adequately progressed at 9 hours for the glass powder obtained in Example 2.

5 **[0058]** On the other hand, FIG. 4 shows measurement results of X-ray diffraction of the glass powder obtained in Comparative Example 1. As shown in FIG. 4, peaks corresponding to (Li_2S) of the raw material were observed in the glass powder following mechanical milling for 20 hours and 25 hours. In contrast, the peak corresponding to (Li_2S) of the raw material was no longer observed in the glass powder
10 mechanically milled for 30 hours, thereby confirming the adequate progression of vitrification.

[0059] Namely, in Comparative Example 1, although it was necessary to carry out mechanical milling for about 30 hours in order to synthesize sulfide-based glass, in Examples 1 and 2, sulfide-based glass was able to be synthesized in about 9 hours. This
15 shortening of synthesis time is thought to be the result of inhibition of adhesion of unreacted raw material to the surface of the milling pot brought about by the addition of an adhesion inhibitor (dehydrated heptane). Moreover, a desired sulfide-based glass was able to be synthesized in Examples 1 and 2 without having to carry out work for removing unreacted raw material every 5 hours as in Comparative Example 1. In
20 addition, although mechanical milling was carried in Example 2 on a scale ten times that of Example 1, a desired sulfide-based glass was confirmed to be able to be synthesized in a short period of time in this case as well.

(2) DSC Measurement

[0060] FIG. 5 shows measurement results of a differential scanning calorimeter
25 (DSC) for the glass powder obtained in Example 1 (following mechanical milling for 15 hours) and the glass powder obtained in Comparative Example 1 (following mechanical milling for 30 hours). As shown in FIG. 5, Example 1 demonstrated a narrower peak width than Comparative Example 1, thereby confirming that sulfide-based glass having a more homogeneous composition was synthesized in Example 1.

(3) Electrical Conductivity

[0061] The glass powder obtained in Example 1 (following mechanical milling for 15 hours) and the glass powder obtained in Comparative Example 1 (following mechanical milling for 30 hours) were respectively heated in an Ar atmosphere under conditions of 290°C for 2 hours to obtain sulfide-based glass ceramic powder. Next, the resulting sulfide-based glass ceramic powder was molded into pellets at a pressure of 5.1 t/cm². Next, the electrical conductivity (Li ion conductivity) of the resulting pellets was measured according to the alternating current 2-terminal method. As a result, electrical conductivity of the pellets using the glass powder obtained in Example 1 (room temperature (25°C)) was 1.7×10^{-3} S/cm, while the electrical conductivity of pellets using the glass powder obtained in Comparative Example 1 (room temperature (25°C)) was 1.5×10^{-3} S/cm, thereby confirming that both types of pellets have about the same degree of electrical conductivity.

[Example 3]

[0062] Lithium sulfide (Li₂S) crystals and phosphorous pentasulfide (P₂S₅) were used as starting materials. 2 g of these powders were weighed at a molar ratio of 75/25 (Li₂S/ P₂S₅) in a dry box in the presence of an argon atmosphere and then placed in a 45 ml zirconia milling pot. Next, 4 g of dehydrated heptane (dehydrated n-heptane) were weighed and placed in the milling pot followed by further placing zirconia grinding balls (diameter: 10 mm, 29 balls) in the milling pot and completely sealing the pot.

[0063] The milling pot was then attached to a planetary ball mill followed initially by carrying out milling for several minutes at low speed (rotational speed: 60 rpm) to sufficiently mix the materials. Subsequently, the rotational speed was gradually increased to carry out mechanical milling at 370 rpm for 9 hours. After drying was performed, glass powder was obtained.

[Example 4]

[0064] Glass powder was obtained in the same manner as Example 3 with the exception of using dehydrated n-decane, instead of dehydrated n-heptane.

[Example 5]

[0065] Glass powder was obtained in the same manner as Example 3 with the exception of using dehydrated toluene, instead of dehydrated heptane.

[Example 6]

[0066] Glass powder was obtained in the same manner as Example 3 with the exception of using dehydrated xylene, instead of dehydrated heptane.

[Evaluation 2]

[0067] X-ray diffraction was performed on the glass powder obtained in Examples 3 to 6 in the same manner as the manner described above. As a result, it was confirmed that sulfide-based glass was synthesized in each of the Examples 3 to 6. Next, the glass powder obtained in each of the Examples 3 to 6 was molded into pellets at a pressure of 5.1 t/cm². Next, the electrical conductivity (Li ion conductivity) of the resulting pellets (at room temperature (25°C)) was measured according to the alternating current 2-terminal method. Table 1 and FIG. 6 show measurement results. Table 1 also shows the SP value of the aprotic liquid and the amount of water contained in the aprotic liquid in each of Examples 3 to 6.

TABLE 1

	Aprotic liquid	Chain/Cyclic	SP value (MJ/cm ³)	Amount of water (ppm)	Li ion Conductivity (S/cm)
Example 3	n-heptane	Chain (straight-chain)	14.4	34	6.4×10^{-4}
Example 4	n-decane	Chain (straight-chain)	15.6	28	5.8×10^{-4}
Example 5	toluene	Cyclic	18.2	30	4.0×10^{-4}
Example 6	xylene	Cyclic	18.0	35	4.3×10^{-4}

[0068] As shown in Table 1 and FIG. 6, it was confirmed that the glass powder obtained in each of Examples 3 to 6 had Li ion conductivity of 10^{-4} S/cm or higher. The glass powder obtained in each of Examples 3 and 4 had higher Li ion conductivity than that of the glass powder obtained in each of Examples 5 and 6. It is considered that because the aprotic liquid with a smaller SP value, that is, the aprotic liquid with lower polarity was used in each of Examples 3 and 4, the reaction between the aprotic liquid and the sulfide-based solid electrolyte (including the unreacted raw material composition)

was more effectively suppressed, deterioration of the sulfide-based solid electrolyte was more effectively suppressed, and thus, Li ion conductivity was improved in each of Examples 3 and 4.

5

CLAIMS

1. A process for producing a sulfide-based solid electrolyte, comprising:
 - preparing a raw material composition containing at least sulfur;
 - adding to the raw material composition an adhesion inhibitor that inhibits formation of adhered substances including the unreacted raw material composition, on an inner surface of a milling pot during mechanical milling; and
 - synthesizing sulfide-based glass by carrying out mechanical milling on the raw material composition to which the adhesion inhibitor has been added.

2. The production process according to claim 1, wherein the adhesion inhibitor is a liquid having the property of not generating hydrogen sulfide in a reaction with the raw material composition.

3. The production process according to claim 2, wherein the liquid is an aprotic liquid.

4. The production process according to claim 3, wherein the aprotic liquid is a non-polar aprotic liquid.

5. The production process according to claim 4, wherein the non-polar aprotic liquid is an alkane that is a liquid at normal temperature.

6. The production process according to claim 5, wherein the non-polar liquid is an alkane that is a liquid at 25°C.

7. The production process according to claim 5 or 6, wherein the number of carbon atoms of the alkane is 5 or more.

8. The production process according to claim 7, wherein the number of carbon atoms of the alkane is 6 or more.

9. The production process according to claim 8, wherein the alkane that is the liquid at normal temperature is n-heptane.
10. The production process according to any one of claims 5 to 9, wherein the boiling point of the alkane is 60 to 300°C.
11. The production process according to claim 10, wherein the boiling point of the alkane is 80 to 200°C.
12. The production process according to any one of claims 2 to 11, wherein the water concentration in the liquid is 100 ppm or less.
13. The production process according to any one of claims 2 to 12, wherein the amount of the liquid is 50 to 1000 parts by weight with respect to 100 parts by weight of the raw material composition.
14. The production process according to claim 1, wherein the adhesion inhibitor is a gel having the property of not generating hydrogen sulfide in a reaction with the raw material composition.
15. The production process according to any one of claims 1 to 14, wherein the raw material composition further contains Li.
16. The production process according to any one of claims 1 to 15, wherein the raw material composition contains at least Li_2S and P_2S_5 .
17. The production process according to claim 16, wherein the molar ratio of Li_2S to P_2S_5 is 68:32 to 74:26.

18. The production process according to claim 17, wherein the molar ratio of Li_2S to P_2S_5 is 70:30.

19. The production process according to any one of claims 1 to 18, wherein the mechanical milling is milling that uses a planetary ball mill.

FIG. 1

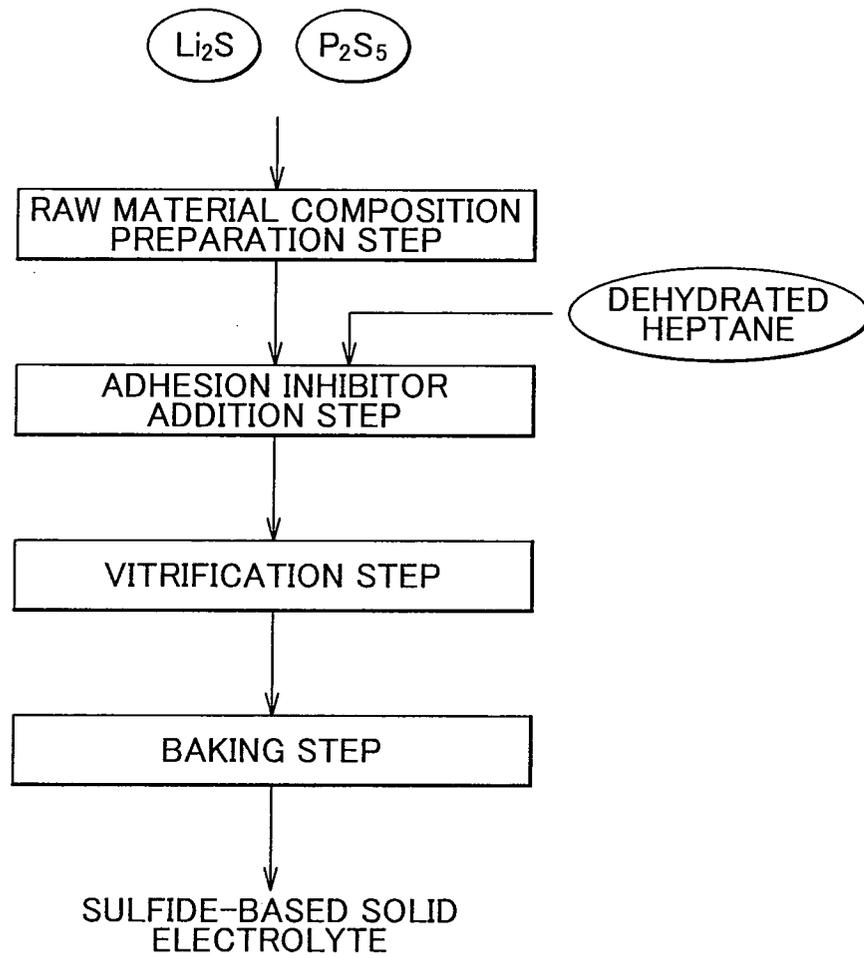


FIG. 2

RELATED ART

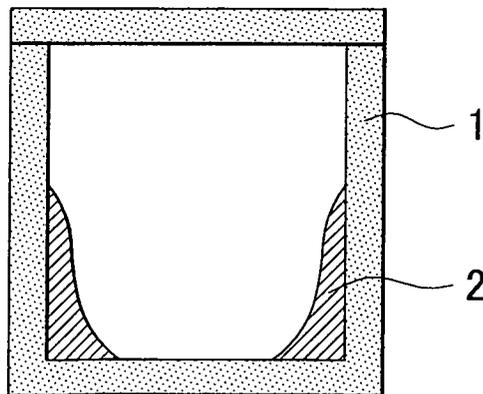


FIG. 3

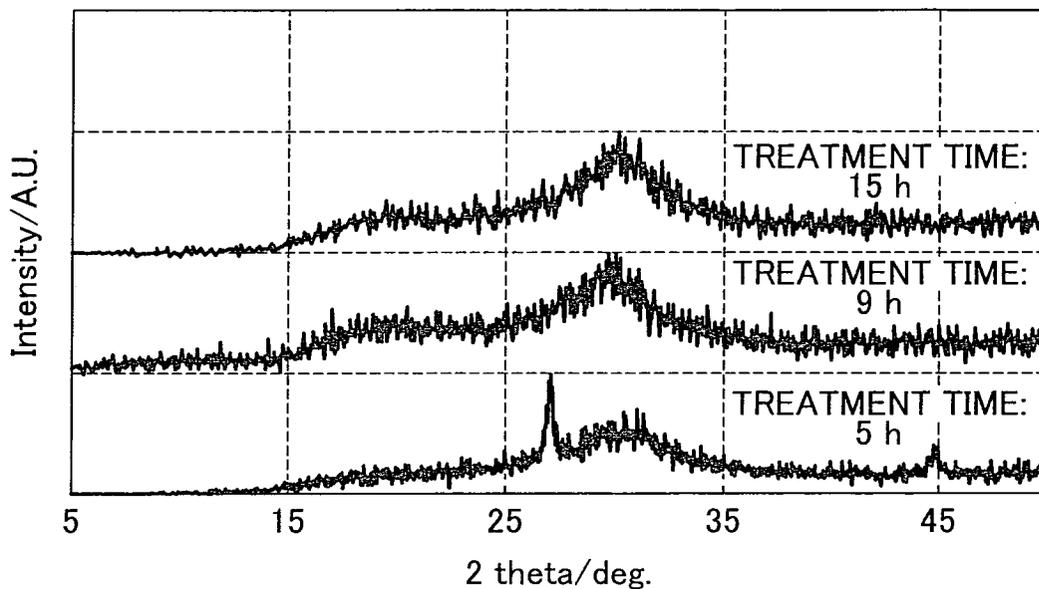


FIG. 4

COMPARATIVE EXAMPLE

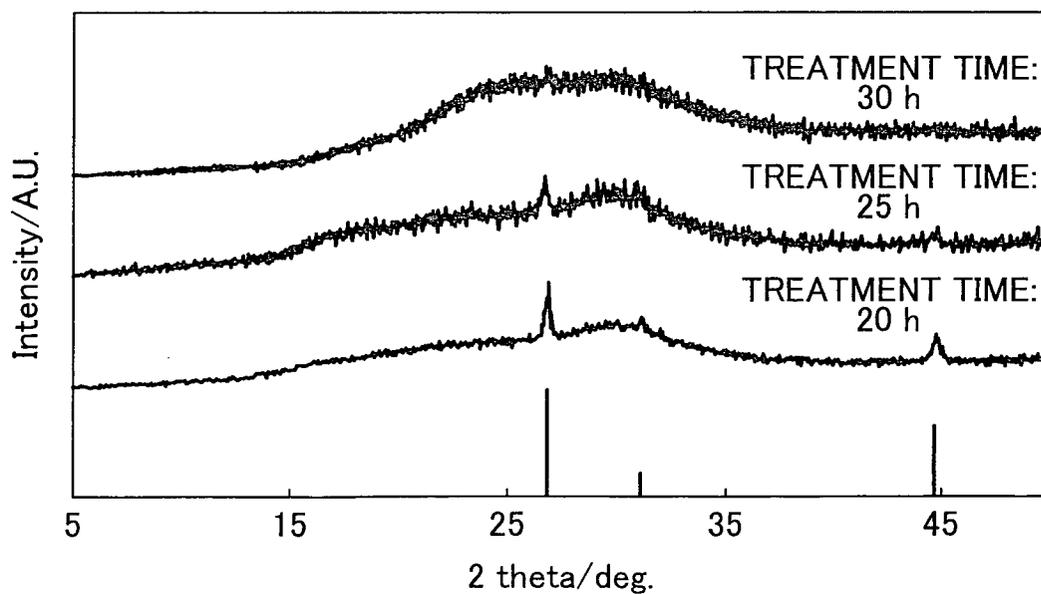


FIG. 5

EXAMPLE 1: WET MM, 15 h
COMPARATIVE EXAMPLE 1: DRY MM, 30 h

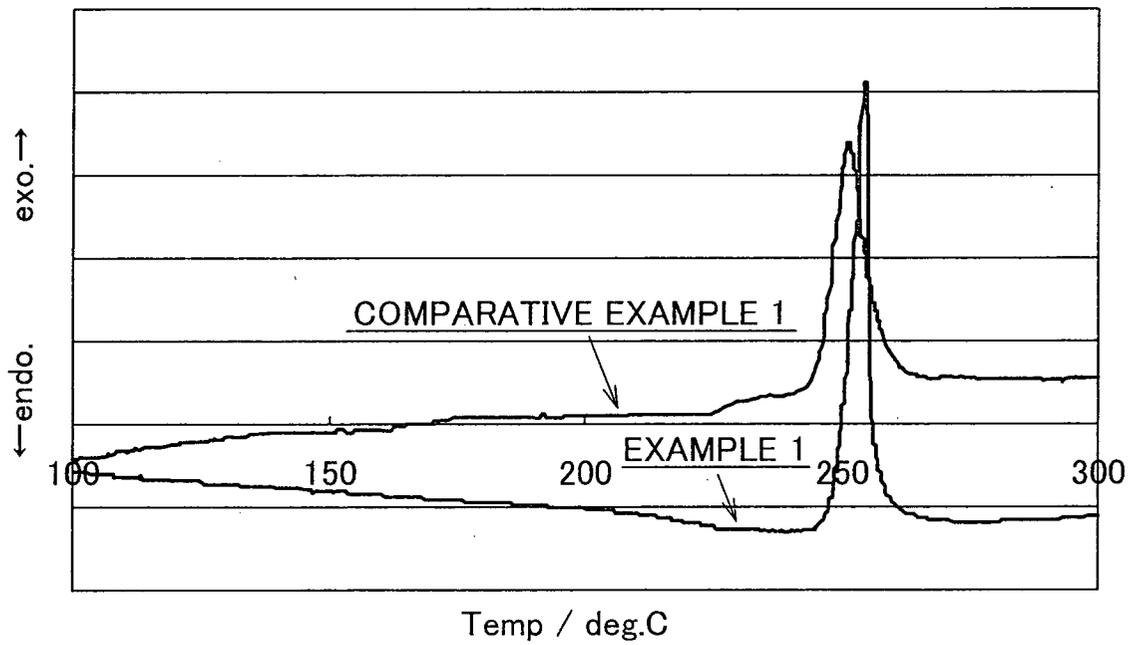
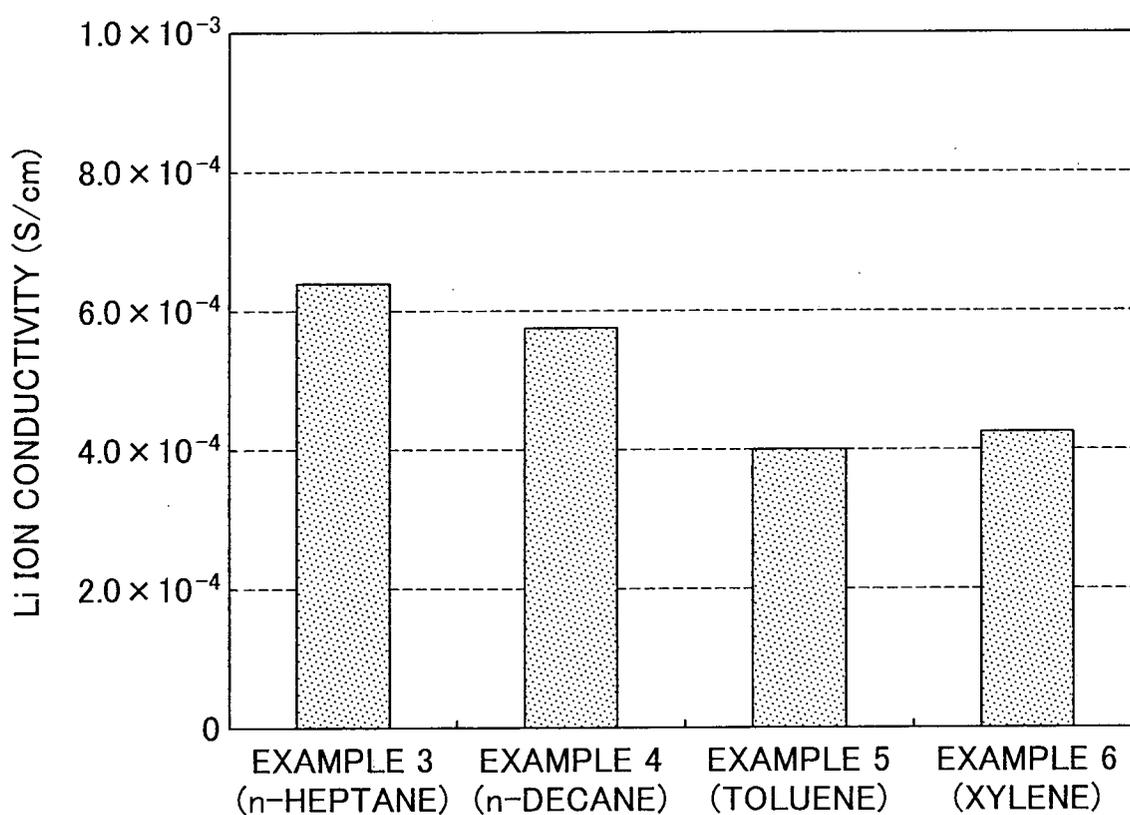


FIG. 6



INTERNATIONAL SEARCH REPORT

International application No
PCT/IB2009/006164

A. CLASSIFICATION OF SUBJECT MATTER INV. H01M10/36 H01M6/18		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols) H01M		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used) EPO-Internal, WPI Data		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 1 466 865 A (IDEMITSU PETROCHEMICAL CO [JP]; AKIBA IWAO [JP] IDEMITSU KOSAN CO [JP]) 13 October 2004 (2004-10-13) cited in the application * see claims * the whole document	1-19
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	-/--	
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input checked="" type="checkbox"/> See patent family annex.		
* Special categories of cited documents :		
A document defining the general state of the art which is not considered to be of particular relevance *E* earlier document but published on or after the international filing date *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) *O* document referring to an oral disclosure, use, exhibition or other means *P* document published prior to the international filing date but later than the priority date claimed		*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. *&* document member of the same patent family
Date of the actual completion of the international search <p align="center">15 October 2009</p>		Date of mailing of the international search report <p align="center">23/10/2009</p>
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016		Authorized officer <p align="center">Stellmach, Joachim</p>

INTERNATIONAL SEARCH REPORT

International application No

PCT/IB2009/006164

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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