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(54) **BIS(PHOSPHINE) BORONIUM SALT, PRODUCTION METHOD OF BIS(PHOSPHINE) BORONIUM SALT, AND BIS(PHOSPHINE) BORONIUM SALT PRODUCED BY THE PRODUCTION METHOD**

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C07F 5/02 (2006.01)

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(58) **Field of Classification Search** None
See application file for complete search history.

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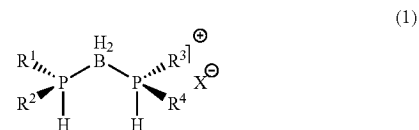
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(57) **ABSTRACT**

The present invention aims to provide a bis(phosphine) boronium salt to be used as a building block or the like for producing a diphosphine compound effective mainly for various kinds of transition metal-catalyzed reactions and further to provide a preferable production method of the bis(phosphine) boronium salt and a bis(phosphine) boronium salt to be produced by the production method.

The present invention provides a bis(phosphine) boronium salt having the structure defined by the following formula (1).



7 Claims, 1 Drawing Sheet

Fig. 1

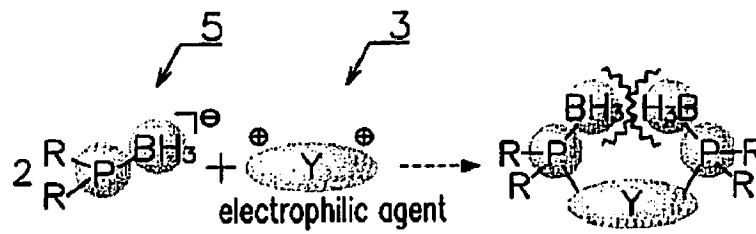
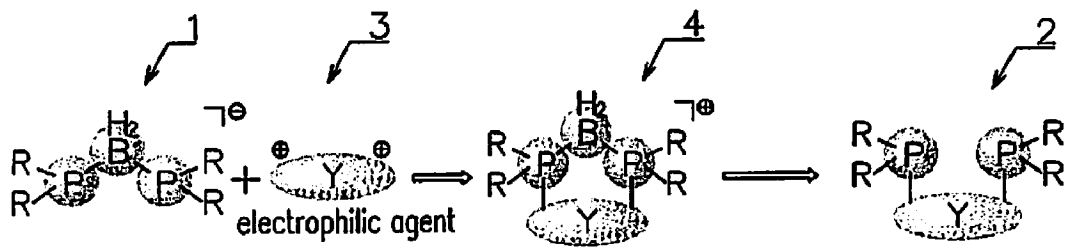


Fig.2



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**BIS(PHOSPHINE) BORONIUM SALT,
PRODUCTION METHOD OF
BIS(PHOSPHINE) BORONIUM SALT, AND
BIS(PHOSPHINE) BORONIUM SALT
PRODUCED BY THE PRODUCTION
METHOD**

TECHNICAL FIELD

The present invention relates to a bis(phosphine) boronium salt to be used as a building block or the like for producing a diphosphine compound effective mainly for various transition metal-catalyzed reactions, a production method of a bis(phosphine) boronium salt, and a bis(phosphine) boronium salt produced by the production method.

BACKGROUND ART

It is reported that a phosphine ligand having a sterically complicated structure is useful for various transition metal-catalyzed reactions such as cross-coupling and investigations on synthesis of phosphine compounds have been addressed as a most important issue at home and abroad (for example, Patent Documents 1 to 3) and have drawn attention as an important approach to develop a highly practical catalytic organic transformation process.

Patent Document 1: JP 2001-253889 A

Patent Document 2: JP 2003-292498 A

Patent Document 3: JP 2003-313194 A

DISCLOSURE OF THE INVENTION

Problems to be Solved by the Invention

However, with respect to synthesis of a diphosphine compound having two phosphino groups at close positions (particularly neighboring positions) and a further sterically complicated structure, almost no effective synthesis process has been proposed so far.

The reason for that is attributed to the complicated steric structure of the diphosphine compound: that is, if two phosphino groups are introduced step by step into close positions, steric hindrance is caused in the introduction of the phosphino group in the second stage and accordingly the reaction is considerably inhibited.

Therefore, the present inventors have made intensive investigations to solve such a problem and consequently have developed a bis(phosphine) boronium salt having a structure comprising two phosphino groups bonded by boronium crosslinking, and have found that if a bis(phosphine) boronium salt is used as a building block for producing a diphosphine compound, the phosphine introduction reaction in the second stage becomes an inter-molecular reaction and the reaction can be relatively smoothly promoted.

Further, the present inventors have also found that the boronium crosslinking portion in the intermediate product after the coupling reaction also works as a protection group for the phosphino group which is relatively easily oxidized, and can be removed easily by a simple operation.

Further, the present inventors have also found that it is possible to easily obtain diphosphine compounds in various structures by combining bis(phosphine) boronium salts in various structures with electrophilic agents.

The present invention has been accomplished based on the above-mentioned findings and aims to provide a bis(phosphine) boronium salt to be used as a building block or the like for producing a diphosphine compound effective for various

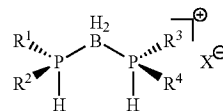
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transition metal-catalyzed reactions as well as a preferable production method of such bis(phosphine) boronium salt and a bis(phosphine) boronium salt produced by the production method.

Means for Solving the Problems

The bis(phosphine) boronium salt of the present invention is a compound having the structure defined by the following formula (1):

[Chem. 2]

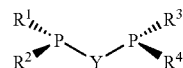


(1)

that is, a compound having a structure in which two phosphino groups are crosslinked by the boronium portion and is useful as a building block or the like for producing a diphosphine compound effective mainly for various transition metal-catalyzed reactions.

The diphosphine compound to be produced using the bis(phosphine) boronium salt of the present invention as a building block may be a compound having the structure defined by the following formula (2):

[Chem. 3]



(2)

that is, a compound having a structure comprising two phosphino groups introduced into a backbone portion Y.

That is, in the case two phosphino groups are to be introduced into a certain backbone portion in a nucleophilic manner, if phosphineborane or phosphine oxide having large substituent groups as a nucleophilic agent, significant steric hindrance is caused to make promotion of the reaction difficult and particularly, in the case of introducing an alkyl-substituted bulky phosphino group such as dialkylphosphine, the reaction is not at all promoted and the synthesis is regarded to be extremely difficult.

In this respect, if the bis(phosphine) boronium salt of the present invention is used as a building block for synthesizing a diphosphine compound, the problem of steric hindrance can be avoided and therefore, synthesis of a large amount of diphosphine, which has been conventionally impossible to be carried out, can be accomplished.

The synthesis of a diphosphine compound using the bis(phosphine) boronium salt of the present invention as a building block involves synthesizing, preferably under a basic condition, an intermediate product by a coupling reaction of the bis(phosphine) boronium salt of the present invention and an electrophilic agent having at least two electrophilic portions and then subjecting the intermediate product to treatment with a de-boronium agent (see the following formula (3)).

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m-tolyl group, a p-tolyl group, a 3,5-xylyl group, a 3,5-diisopropylphenyl group, a p-anisyl group, a p-trifluoromethylphenyl group, an α -naphthyl group, a β -naphthyl group and the like, and the alkoxy group may be selected properly from a methoxy group, an ethoxy group, an isopropoxy group, a tert-butoxy group, a phenoxy group and the like.

Since it is confirmed that high reactivity and stereoselectivity can be obtained in rhodium-catalyzed asymmetric hydrogenation or a palladium-catalyzed asymmetric carbon-carbon bond formation reaction using P-chiral phosphines having the asymmetric center on the phosphorus atom, it is preferable for the diphosphine compound produced using the bis(phosphine) boronium salt of the present invention as a building block that the phosphorus atom forms the asymmetric center.

In this respect, by using a compound containing a phosphorus atom forming the asymmetric center as a phosphine compound, a starting material, the method of the present invention makes it possible to produce a bis(phosphine) boronium salt having the asymmetric center on at least one of phosphorus atoms and use of the bis(phosphine) boronium salt as a building block makes it possible to produce a diphosphine compound having the asymmetric center on the phosphorus atom.

Further, in coupling reactions using various kinds of transition metal catalysts having monophosphine as a ligand, it is supposed that the reaction is smoothly promoted owing to prevention of stable dimer formation during the reaction by the catalytic active species because of the bulky ligand, that is, the steric hindrance, and accordingly, also with respect to the diphosphine compound to be produced using the bis(phosphine) boronium salt of the present invention as a building block, it is preferable to select a substituent group having a bulky structure for at least one of substituent groups R^1 to R^4 , for example, an alkyl group having a branched structure such as a tert-butyl group; an alkyl group having a cyclic structure such as a cyclohexyl group and an adamantyl group; and an aryl group.

In this respect, in the method of the present invention, it is made possible to produce a bis(phosphine) boronium salt having a bulky structure for at least one of the substituent groups R^1 to R^4 on the phosphorus atom by selecting a compound having a substituent group with a bulky structure, for example, an alkyl group having a branched structure such as a tert-butyl group; an alkyl group having a cyclic structure such as an adamantyl group and a cyclohexyl group; and an aryl group, for at least one of substituent groups on the phosphorus atom of the phosphine compound as a starting material, and by using the bis(phosphine) boronium salt as a building block, the diphosphine compound having a bulky structure for at least one of substituent groups of the phosphorus atom can be produced.

Use of the bis(phosphine) boronium salt of the present invention produced by the production method of the present invention mainly as a building block for synthesizing a diphosphine compound makes it possible to avoid the problem of the steric hindrance and to accomplish synthesis of a large amount of diphosphine which could not be conventionally synthesized.

EFFECTS OF THE INVENTION

The bis(phosphine) boronium salt of the present invention has the above-mentioned structure and is a novel compound capable of giving diphosphine compounds having various structures by a simple synthetic process.

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That is, if the bis(phosphine) boronium salt of the present invention is used as a building block for producing a diphosphine compound, the phosphine introduction reaction in the second stage becomes the intramolecular reaction and is relatively smoothly promoted.

The boronium crosslinking portion in an intermediate product after the coupling reaction works as a protection group for the phosphino groups, which tend to be oxidized relatively easily, and further it is easy to remove the portion by treatment with various kinds of de-boronium agents such as fluoride anions.

Further, combination of the bis(phosphine) boronium salt of the present invention which has various structures with an electrophilic agent can easily give diphosphine compounds having various structures.

The method of the present invention is a method preferable for producing the above-mentioned bis(phosphine) boronium salt of the present invention and includes two synthesis routes: a symmetric synthesis route and an asymmetric synthetic route.

The former symmetric synthetic route is characterized in that a phosphine compound is treated with mono-substituted borane having a substituent group for a hydrogen group on the boron atom and use of phosphine compounds having various steric and electronic characteristics makes it possible to give various kinds of symmetric bis(phosphine) boronium salts.

On the other hand, the latter asymmetric synthetic route is characterized in that phosphine compounds are treated with phosphineborane of which hydrogen group on the boron atom is mono-substituted, and it is possible to synthesize asymmetric bis(phosphine) boronium salts having various substitution patterns on two phosphorus atoms.

Accordingly, if the bis(phosphine) boronium salt of the present invention produced by the method of the present invention is used mainly as a building block for synthesizing a diphosphine compound, the problem of steric hindrance can be avoided and therefore, synthesis of a large amount of diphosphine, which has been conventionally impossible to be carried out, can be accomplished.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a schematic view showing a state of introducing two phosphino groups into a certain backbone portion in a nucleophilic manner using phosphineborane.

FIG. 2 is a schematic view showing a state of introducing two phosphino groups into a certain backbone portion in a nucleophilic manner using a bis(phosphine)boronium salt of the present invention.

EXPLANATION OF SYMBOLS

- 1 deprotonated bis(phosphine)boronium salt
- 2 diphosphine compound
- 3 electrophilic agent
- 4 intermediate product
- 5 deprotonated phosphineborane

BEST MODES FOR CARRYING OUT THE INVENTION

Hereinafter, Examples of the present invention will be described, however it is not intended that the present invention be limited to the described Examples.

FIG. 1 illustrates a conventional method for introducing two phosphino groups into an electrophilic agent 3 having a certain backbone portion in a nucleophilic manner using

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phosphineborane (deprotonated form) **5**: that is, if the phosphineborane **5** is used as a nucleophilic agent, significant steric hindrance is caused at the time of introduction of the phosphino group in the second stage and accordingly the reaction is hardly promoted and particularly, the reaction of introduction of an alkyl-substituted bulky phosphino group such as dialkylphosphine cannot be promoted at all.

On the other hand, FIG. 2 illustrates a method for introducing two phosphino groups into an electrophilic agent **3** having a certain backbone portion in a nucleophilic manner using a bis(phosphine) boronium salt (deprotonated form) **1** of the present invention: that is, since the bis(phosphine) boronium

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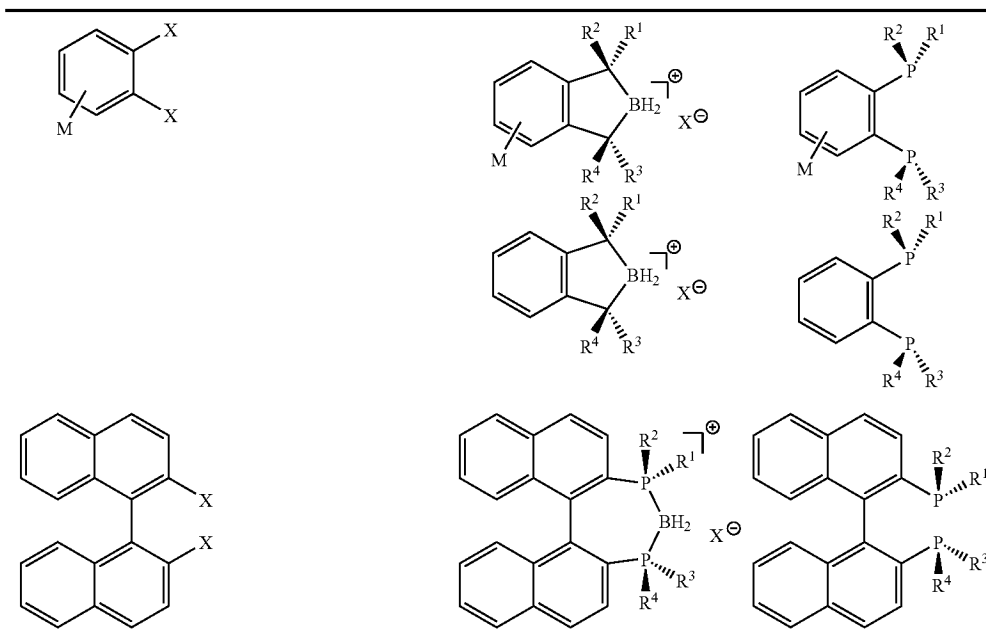
salt **1** of the present invention is used as a building block for the synthesis of a diphosphine compound **2**, the phosphine introduction reaction in the second stage becomes an intramolecular reaction to avoid the problem of the steric hindrance and therefore, synthesis of a large amount of diphosphine compounds **2**, which cannot be conventionally synthesized, can be accomplished.

Table 1 shows examples of various kinds of structures of diphosphine compounds produced by a coupling reaction of the bis(phosphine) boronium salt of the present invention with electrophilic agents having various structures.

TABLE 1

Electrophilic agent	Bis(phosphine) boronium salt	Intermediate product	Diphosphine compound

TABLE 1-continued



That is, combinations of the bis(phosphine) boronium salt of the present invention with electrophilic agents having various structures easily gives diphosphine compounds in various structures and a library of diphosphine compounds can be built according to the combinations.

Example 1

Preparation of (S)-(tert-butyl(methyl)phosphine)(di-tert-butylphosphine) boronium iodide

A 30 mL two-necked eggplant flask was charged with 133 mg of lithium aluminum hydride and a magnetic stirrer and purged with nitrogen and 5 mL of diethyl ether was added and the contents were cooled to 0° C. while being stirred.

Further, 627 μ L of di-tert-butylchlorophosphine was added and after the resulting mixture was stirred at room temperature for 1 hour, the resulting reaction solution was filtered by passing the solution through about 3 mg of basic alumina under nitrogen gas flow and eluted with about 15 mL of diethyl ether.

The filtrate was concentrated under reduced pressure and the obtained residue was dissolved in 9 mL of dichloromethane after reintroduction of nitrogen.

Meanwhile, a 30 mL two-necked eggplant flask was charged with 300 mg of (S)-tert-butyl(methyl)phosphineborane, 323 mg of iodine, and a magnetic stirrer and purged with nitrogen and 5 mL of dichloromethane was added for dissolution and the resulting solution was stirred at room temperature for 7 hours and mixed with the prepared dichloromethane solution of di-tert-butylphosphine at room temperature and stirred for 24 hours.

After 1 mL of a 1.2 M borane-tetrahydrofuran complex tetrahydrofuran solution was added and stirred at room temperature for 30 minutes, 5 mL of water and 7 mL of dichloromethane were added and after the resulting mixture was

separated, the water layer was extracted with 7 mL of dichloromethane two times and sodium sulfate was added to dehydrate the organic layer.

The resulting product was filtered and the filtrate was concentrated by a rotary evaporator to obtain a residue which was purified by silica gel column chromatography (chloroform/methanol=20/1 \rightarrow 10/1) to obtain 618 mg of (S)-(tert-butyl(methyl)phosphine)(di-tert-butylphosphine) boronium iodide as a white solid (yield 62%).

Example 2

Preparation of bis(di-tert-butylphosphine)boronium bromide

A 30 mL two-necked eggplant flask was charged with 229 mg of lithium aluminum hydride and a magnetic stirrer and purged with nitrogen and 5 mL of diethyl ether was added and the contents were cooled to 0° C. while being stirred.

Further, 1.08 mL of di-tert-butylchlorophosphine was added and stirred at room temperature for 1 hour, and thereafter, the resulting reaction solution was filtered by passing the solution through about 3 g of basic alumina under nitrogen gas flow and eluted with about 15 mL of diethyl ether.

The filtrate was concentrated under reduced pressure and after nitrogen reintroduction, the obtained residue was dissolved in 10 mL of dichloromethane and mixed with 2.7 mL of a 1.0 M monobromoborane-methyl sulfide complex dichloromethane solution at room temperature and stirred for 40 hours.

The resulting reaction solution was concentrated under reduced pressure and the obtained residue was purified by silica gel column chromatography (chloroform/methanol=20/1 \rightarrow 10/1) to obtain 743 mg of bis(di-tert-butylphosphine)boronium bromide as a white solid (yield 72%).

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Example 3

Preparation of meso- and (S,S)-bis(tert-butyl(methyl)phosphine)boronium iodide

A 50 mL two-necked eggplant flask was charged with 3.04 g of potassium carbonate, 1.02 g of tert-butyl(methyl)phosphonium iodide, and a magnetic stirrer and purged with nitrogen and 10 mL of dichloromethane was added.

After the contents were stirred at room temperature for 1 hour, the resulting reaction solution was filtered by passing the solution through about 3 g of basic alumina under nitrogen gas flow and eluted with about 15 mL of dichloromethane.

Meanwhile, a 100 mL two-mouth eggplant flask was loaded with 400 mg of (S)-tert-butyl(methyl)phosphineborane, 430 mg of iodine, and a magnetic stirrer and purged with nitrogen and 5 mL of dichloromethane was added for dissolution and the resulting solution was stirred at room temperature for 7 hours and mixed with the prepared tert-butyl(methyl)phosphine dichloromethane solution at room temperature and stirred for 24 hours.

After 1.4 mL of a 1.2 M borane-tetrahydrofuran complex tetrahydrofuran solution was added and stirred at room temperature for 30 minutes, 5 mL of water and 7 mL of dichloromethane were added and after the resulting mixture was fiercely stirred and an organic layer and a water layer were separated, the water layer was extracted with 7 mL of dichloromethane two times and sodium sulfate was added to dehydrate the organic layer.

The resulting product was filtered and the filtrate was concentrated by a rotary evaporator to obtain a residue which was refined by silica gel column chromatography (chloroform/methanol=20/1→10/1) to obtain 697 mg of bis(tert-butyl(methyl)phosphine)boronium iodide as a mixture of meso and (S,S) isomers as a white solid (yield 59%).

The obtained product was dissolved in 6 mL of tetrahydrofuran and recrystallized to obtain 208 mg of meso isomer as a plate-like crystal and 151 mg of (S,S) isomer as a needle-like crystal.

Example 4

Preparation of bis(diphenylphosphine)boronium bromide

A 30 mL two-necked eggplant flask was charged with 20 mL of a hexane solution containing 10% by weight of diphenylphosphine and a magnetic stirrer and the solution was concentrated under reduced pressure and after nitrogen introduction, the obtained residue was dissolved in 5 mL of dichloromethane.

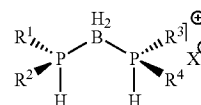
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Further, 3.5 mL of a 1.0 M monobromoborane-methyl sulfide complex dichloromethane solution was added to the resulting solution at room temperature and stirred for 35 hours.

Further, the resulting reaction solution was concentrated under reduced pressure and the obtained residue was washed with tetrahydrofuran to obtain 1.0 g of bis(diphenylphosphine)boronium bromide as a white solid (yield 65%).

What is claimed is:

1. A bis(phosphine) boronium salt having the structure defined by the following formula:



(1)

wherein R1, R2, R3, and R4 are independently selected from the group consisting of an alkyl group, an aryl group, a hydroxyl group, and an alkoxy group, and wherein X is selected from the group consisting of a halide ion, CF₃SO₃—, BF₄—, PF₆— and SbF₆—.

2. A production method of the bis(phosphine) boronium salt according to claim 1, comprising treatment of a phosphine compound with mono-substituted borane.

3. A production method of the bis(phosphine) boronium salt according to claim 1, comprising treatment of a phosphine compound with mono-substituted phosphineborane having a substituent group for a hydrogen atom on the boron atom.

4. The production method of the bis(phosphine) boronium salt according to claim 2, wherein the phosphorus atom of the phosphine compound forms an asymmetric center.

5. The production method of the bis(phosphine) boronium salt according to claim 2, wherein at least one of substituent groups of the phosphine compound is an alkyl group having a branched structure, an alkyl group having a cyclic structure, or an aryl group.

6. A bis(phosphine) boronium salt produced by the production method of a bis(phosphine) boronium salt according to claim 2.

7. A bis(phosphine) boronium salt as described in claim 1, wherein the alkyl group is selected from the group consisting of a methyl group, an ethyl group, an isopropyl group, a cyclohexyl group, a tert-butyl group, an adamantyl group and a benzyl group.

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