

## A PROCESS FOR THE PREPARATION OF OPTICALLY ACTIVE CYCLOPROPYLAMINES.

### The Field of the Technology

This invention relates to a process for the production of optically active 2-(disubstituted  
5 aryl) cyclopropylamine derivatives and optically active 2-(disubstituted aryl) cyclopropane  
carboxamide derivative which are useful intermediates for the preparation of  
pharmaceutical agents, and in particular the compound [1S-(1 $\alpha$ , 2 $\alpha$ , 3 $\beta$  (1S\*,2R\*),5 $\beta$ )]-3-  
[7-[2-(3,4-difluorophenyl)-cyclopropyl]amino]-5-(propylthio)-3H-1,2,3-triazolo[4,5-  
d]pyrimidin-3-yl)-5-(2-hydroxyethoxy)-cyclopentane-1,2-diol. This compound, and  
10 similar such compounds, are disclosed in WO 00/34283 and WO 99/05143. These  
compounds are disclosed as P<sub>2T</sub> (which is now usually referred to as P<sub>2Y12</sub>) receptor  
antagonists. Such antagonists can be used as, *inter alia*, inhibitors of platelet activation,  
aggregation or degranulation.

### Background Technology

Some processes are known for the production of optically active 2-cyclopropane  
carboxamide derivatives, optically active 2-aryl cyclopropylamine derivatives and  
optically active 2-arylcyclopropane-1-carboxylate ester derivatives.

20 Examples of processes for the production of optically active 2-arylcyclopropane  
carboxamide derivatives, are

- (i) A process wherein excess thionyl chloride is reacted with optically active 2-  
phenylcyclopropane carboxylic acid in benzene solvent to form corresponding acid  
chloride, and after concentrating down excess thionyl chloride and benzene under reduced  
25 pressure, the acid chloride is isolated and purified by distillation, and, by causing ammonia  
water to act on this, 2-phenylcyclopropane carboxamide is obtained (J. Am. Chem. Soc.  
Vol.109, p.2311 (1987), Journal of Medicinal Chemistry Vol.20, p.771 (1977));
- (ii) A Process to obtain optically active 3-aryl-2-dimethylcyclopropane-1-carboxamide by  
causing ammonia water to act on the corresponding acid chloride formed by reacting  
30 thionyl chloride with optically active 3-aryl-2-dimethylcyclopropane-1-carboxylic acid (J.  
Org. Chem. Vol.68, p.621 (2003));

Examples of processes for the production of optically active 2-aryl cyclopropylamine derivatives, are:

- (iii) A process wherein chlorocarbonic acid ethyl ester is reacted with 2-aryl cyclopropane carboxylic acid to form mixed acid anhydride, and by causing to act sodium azide on this, corresponding acid azide is formed, and 2-aryl cyclopropylamine is obtained by Curtius rearrangement with this (Journal of Medicinal Chemistry Vol.20, p.771 (1977), WO01/92263);
- (iv) A process to obtain corresponding 2,2-dimethyl cyclopropylamine by causing chlorine, bromine or sodium hypochlorite to act on the optically active 2,2-dimethylcyclopropane-1-carboxamide in the presence of base (Kokoku 5-3865);

Examples of a process for the production of optically active 2-arylcyclopropane carboxylate ester derivatives, for example:

- (v) A process to obtain optically active cyclopropanecarboxylic acid derivative by cyclopropanation after deriving into optically active ester or amide via several steps using benzaldehyde derivative as the starting material (WO01 /92263);
- (vi) A process to obtain optically active 2-dihydrofuranyl cyclopropanecarboxylate derivative by reacting phosphonoacetic acid ester derivative with optically active dihydrobenzofuranyl ethylene oxide derivative in the presence of base (Organic Process Research & Development, vol 6, p.618 (2002));

Examples of a process to produce optically active 2-aryl cyclopropylamine derivatives from optically active 2-aryl cyclopropanecarboxylic acid, are:

- (vii) A process wherein benzaldehydes is used as the starting material and derived into optically active ester or amide via several steps, and thereafter optically active 2-aryl cyclopropane carboxylate ester is obtained by cyclopropanation. This optically active carboxylic acid derivative is formed into acid azide, and optically active 2-aryl cyclopropylamine derivative is produced by Curtius rearrangement (WO01/92263).

### Disclosure of the Invention

In the process for the production of optically active 2-arylcyclopropane carboxamides referred to in (i) above, only the process to produce 2-phenylcyclopropane carboxamide  
5 from 2-phenylcyclopropane carboxylic acid is described and a process for production for 2-(disubstituted aryl) cyclopropane carboxamide derivative is not disclosed. Moreover, in the process (ii), there is mentioned the process for production only of 2,2-dimethyl-3-phenylcyclopropane carboxamide and 2,2-dimethyl-3-isopropylidene cyclopropane carboxamide, and a process for production of 2-(disubstituted aryl) cyclopropane  
10 carboxamide derivative is not disclosed.

Secondly, in a process for the production of optically active 2-aryl cyclopropylamine derivative, optically active 2-aryl cyclopropylamine derivative is produced by Curtius rearrangement from optically active 2-arylcyclopropane carboxylic acid in the aforesaid  
15 process (iii), however, it is not suitable for a commercial preparation method from the viewpoint of safety because it is via an acid azide intermediate having explosive properties. Moreover, in the process (iv), optically active amine is produced from the optically active carboxamide by a Hofmann rearrangement. However, it is not suitable for a commercial preparation method from the viewpoint of economy because yield is low  
20 when the reaction is carried out using the sodium hypochlorite. Moreover, as for the aforesaid process (iv), only the process to produce optically active 2,2-dimethyl cyclopropylamine from optically active 2,2-dimethylcyclopropane carboxamide is mentioned, and a process for production of 2-(disubstituted aryl) cyclopropane carboxamide derivative is not disclosed.

25 Thirdly, in a process for the production of optically active 2-arylcyclopropane carboxylate ester derivative, in the aforesaid process (v), optically active 3,4-difluorophenyl cyclopropanecarboxylic derivative is obtained by cyclopropanation after converting 3,4-difluoro benzaldehyde starting material into optically active ester or amide via several  
30 steps. However, it is not commercially suitable from the viewpoint of productivity and economy. For example, the starting material is expensive, the stereoselectivity is insufficient in the cyclopropanation and also there are many numbers of steps. Moreover,

in process (vi), only an example of preparing optically active dihydrofuranyl cyclopropanecarboxylate ester from optically active dihydrobenzofuranyl ethylene oxide is mentioned. It is not a process for the production of general optical activity 2-arylcyclopropane carboxylate ester.

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Fourthly, a process to produce optically active 2-aryl cyclopropylamine derivative from optically active 2-arylcyclopropane carboxylate ester derivative using (vii) is not commercially viable from a safety standpoint because the acid azide intermediate has expulsion properties. Also, purification is essential due to insufficient stereoselectivity during the cyclopropanation, making this process unsuitable for commercial preparation because of poor productivity.

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Thus, the processes outlined are unsuitable for commercial production. There is a need for a commercial process which addresses areas such as safety, economy, productivity and the like.

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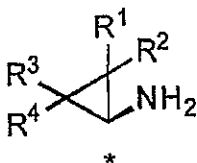
An efficient process has now been discovered for the production of optically active 2-aryl cyclopropylamine derivatives or salts thereof. The process affords high optical purity by using a readily available optically active styrene oxide derivative as the starting material.

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An efficient process for the production of optically active cyclopropylamine derivative by a Hofmann rearrangement using sodium hypochlorite has been discovered. This process can be used safely and inexpensively as a commercial preparation method.

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Thus, according to the present invention there is provided a process for the production of optically active cyclopropylamine derivatives represented by general formula (2) or salts thereof



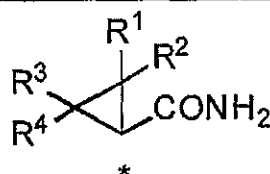
( 2 )

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(wherein  $R^1$ ,  $R^2$ ,  $R^3$  or  $R^4$  denote a hydrogen atom, optionally substituted 1-10C cyclic or acyclic alkyl group, optionally substituted 6-10C aryl group or optionally substituted 7-10C aralkyl group, and \* denotes an asymmetric carbon centre)

characterised by reacting optically active cyclopropane carboxamide derivative

5 represented by general formula (1)



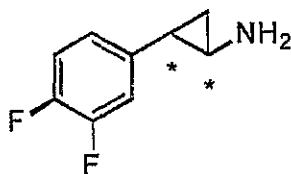
( 1 )

(wherein  $R^1$ ,  $R^2$ ,  $R^3$ ,  $R^4$  and \* have the same said definitions) with hypochlorite in water in the presence of alkali metal hydroxide of 5-30 equivalent.

10 Suitably, the hypochlorite is sodium hypochlorite; and in particular the quantity used of the hypochlorite is 1-5 mole equivalent with respect to compound of the formula (1). In a particular embodiment there is provided a process for the production of optically active cyclopropylamine derivatives or salts thereof wherein  $R^1$ ,  $R^2$ ,  $R^3$  is hydrogen atom and  $R^4$  is 3,4-difluorophenyl group.

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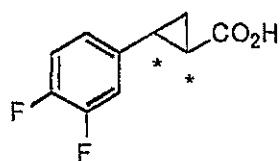
In a further embodiment, there is provided is a process for the production of an optically active 2-aryl cyclopropylamine derivative represented by general formula (9) or a salt thereof,



( 9 )

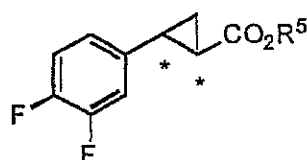
20 (wherein \* denotes an asymmetric carbon centre), wherein an optically active 2-aryl cyclopropanecarboxylic acid derivative represented by general formula (7)

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( 7 )

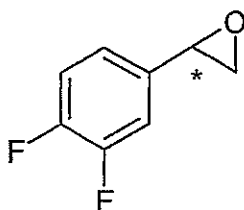
(wherein \* denotes an asymmetric carbon centre) is obtained by de-esterifying the optically active 2-arylcyclopropane carboxylate ester derivative represented by general formula (6)



( 6 )

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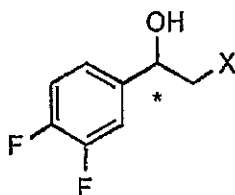
(wherein, R<sup>5</sup> denotes optionally substituted 1-10C cyclic or acyclic alkyl group, optionally substituted 6-10C aryl group or optionally substituted 7-10C aralkyl group, and \* denotes an asymmetric carbon centre) which is obtained by reacting the optically active styrene oxide derivative represented by general formula (3)



( 3 )

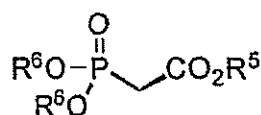
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(wherein \* denotes an asymmetric carbon centre) or optically active halohydrin derivative represented by or general formula (4)



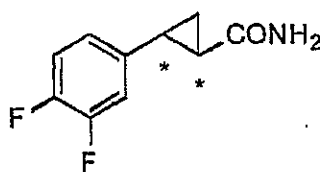
( 4 )

(wherein X denotes a halogen atom, and \* denotes an asymmetric carbon centre) with phosphonoacetic acid ester derivative represented by general formula (5)



( 5 )

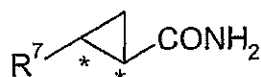
(wherein R<sup>5</sup> or R<sup>6</sup> denote optionally substituted 1-10C cyclic or acyclic alkyl group, optionally substituted 6-10C aryl group or optionally substituted 7-10C aralkyl group) in the presence of base, and optically active 2-aryl cyclopropane carboxamide derivative represented by obtained general formula (8)



( 8 )

(wherein \* denotes an asymmetric carbon centre) which is obtained by reacting the obtained aforesaid 2-aryl cyclopropanecarboxylic acid derivative with ammonia after being activated with carboxylic acid activator is reacted with oxidant.

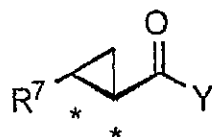
There is also provided a process for the production of optically an active 2-aryl cyclopropane carboxamide derivative represented by general formula (12)



( 12 )

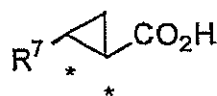
(wherein R<sup>7</sup> denotes an aryl group substituted by 2 or more halogen atoms, and \* denotes an asymmetric carbon centre) characterized by reacting with ammonia, optically active 2-aryl cyclopropanecarboxylic acid derivative represented by general formula (11)

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( 1 1 )

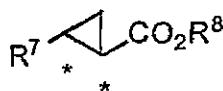
(wherein,  $R^7$  denotes an aryl group substituted by 2 or more halogen atoms, Y denotes carbonyl group activated group, and \* denotes an asymmetric carbon centre) which is obtained from an optically active 2-aryl cyclopropanecarboxylic acid derivative  
 5 represented by general formula (10)



( 1 0 )

(wherein  $R^7$  denotes an aryl group substituted by 2 or more halogen atoms, and \* denotes an asymmetric carbon centre) by reacting with a carboxylic acid activator.

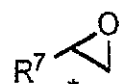
- 10 There is also provided a process for the production of an optically active 2-aryl cyclopropane carboxamide derivative, wherein the reaction is carried out by using the compound of formula (10) obtained by de-esterifying an optically active 2-aryl cyclopropane carboxylate ester derivative represented by general formula (13)



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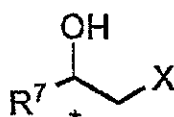
- 15 (wherein  $R^8$  denotes optionally substituted 1-10C cyclic or acyclic alkyl group, optionally substituted 6-10C aryl group or optionally substituted 7-10C aralkyl group, and  $R^7$  and \* have the same said definitions).

There is also provided a process for the production of an optically active 2-aryl cyclopropane carboxamide derivative, wherein the reaction is carried out by using the compound of formula (13) obtained by reacting the optically active styrene oxide derivative represented by general formula (14)



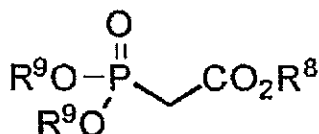
( 1 4 )

(wherein R<sup>7</sup> and \* have the same said definitions) or optically active halohydrin derivative represented by general formula (15)



( 1 5 )

(wherein R<sup>7</sup> and \* have the same said definitions) with phosphonoacetic acid ester derivative represented by general formula (16)



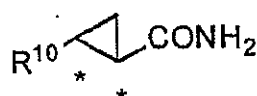
( 1 6 )

(wherein R<sup>9</sup> denotes optionally substituted 1-10C cyclic or acyclic alkyl group, optionally substituted 6-10C aryl group or optionally substituted 7-10C aralkyl group, and R<sup>8</sup> and \* have the same said definitions) in the presence of base. There is also provided a process for the production of an optically active 2-aryl cyclopropane carboxamide derivative to obtain (1R,2R)-2-aryl cyclopropane carboxamide derivative of formula (12) using a (1R,2R)-2-aryl cyclopropanecarboxylic acid derivative of formula (10). The present invention also provides a process for the production of an optically active 2-aryl cyclopropane carboxamide derivative to obtain a (1R,2R)-2-aryl cyclopropane carboxylic

acid derivative formula (10) using a (1R,2R)-2-aryl cyclopropane carboxylate ester derivative of formula (13).

There is also provided a process for the production of an optically active 2-aryl cyclopropane carboxamide derivative to obtain a (1R,2R)-2-aryl cyclopropane carboxylate ester derivative formula (13) using (S)-styrene oxide derivative formula (14) and (S)-halohydrin derivative formula (15). In particular there is provided a process for the production of an optically active 2-aryl cyclopropane carboxamide derivative, wherein R<sup>7</sup> is 3,4-difluorophenyl group.

The present invention also provides an optically active 2-arylcyclopropane carboxamide derivative represented by general formula (17)



( 1 7 )

(wherein R<sup>10</sup> denotes an aryl group substituted by 2 or more halogen atoms, and \* denotes an asymmetric carbon centre).

In particular, in the optically active 2-arylcyclopropane carboxamide derivative of formula (17), R<sup>10</sup> is a 3,4-difluorophenyl group.

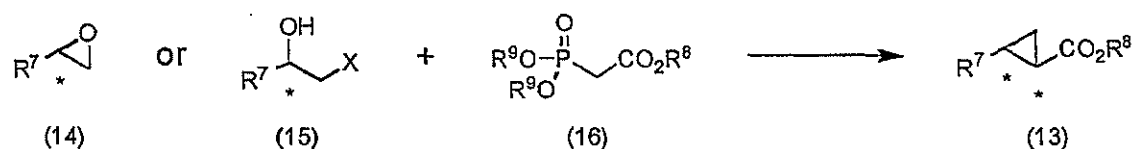
More particularly, the compound of formula (17) is a (1R,2R)-2-aryl cyclopropane carboxamide derivative.

The present invention provides a process for preparing an optically active aminocyclopropane derivative from inexpensive 3, 4-difluorobenzene using a Hoffmann re-arrangement. In general, the process is a safe and inexpensive way of preparing the optically active aminocyclopropane derivative which is useful as an intermediate in the manufacture of pharmaceuticals and pesticides.

The conversion of compounds of formula (14) to (2) comprises 4 steps, namely in total 1) cyclopropanation process, 2) deesterification process, 3) amidation process and 4) Hofmann rearrangement process. Hereinafter, the invention is described in detail for each process.

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Firstly, there will be described 1) cyclopropanation process.



#### Step 1. Cyclopropanation process

10 In compounds represented by formula (14),  $\text{R}^7$  denotes an aryl group substituted by 2 or more halogen atoms. Suitable values for  $\text{R}^7$  includes for example, a 2,3-difluorophenyl group, 3,4-difluorophenyl group, 2,4-difluorophenyl group, 2,3,4-trifluorophenyl group, 3,4,5-trifluorophenyl group, 2,3,4,5-tetrafluorophenyl group, 2,3,4,5,6-pentafluorophenyl group, 2,3-dichlorophenyl group, 3,4-dichlorophenyl group, 2,4-dichlorophenyl group, 2,3,4-trichlorophenyl group, 3,4,5-trichlorophenyl group, 2,3,4,5-tetrachlorophenyl group, 2,3,4,5,6-pentachlorophenyl group, 2,3-dibromophenyl group, 3,4-dibromophenyl group, 2,4-dibromophenyl group, 2,3,4-tribromophenyl group, 3,4,5-tribromophenyl group, 2,3,4,5-tetrabromophenyl group, 2,3,4,5,6-pentabromo phenyl group. 3,4-difluorophenyl group is preferred. Moreover, \* denotes an asymmetric carbon centre. In other words, a  
 15 styrene oxide derivative formula (14) contains an asymmetric carbon centre. This invention includes any optically active substance or racemic mixture of the compound of formula (14). Preferably it is optically active substance, and most preferably it is a  
 20 compound whose absolute configuration of asymmetric carbon centre is (S).

25 In the compound of formula (15),  $\text{R}^7$  denotes an aryl group substituted by 2 or more halogen atoms, and X denotes a halogen atom. Suitable values for  $\text{R}^7$  include for example, a 2,3-difluorophenyl group, 3,4-difluorophenyl group, 2,4-difluorophenyl group, 2,3,4-trifluorophenyl group, 3,4,5-trifluorophenyl group, 2,3,4,5-tetrafluorophenyl group, 2,3,4,5,6-pentafluorophenyl group, 2,3-dichlorophenyl group, 3,4-dichlorophenyl group,

2,4-dichlorophenyl group, 2,3,4-trichlorophenyl group, 3,4,5-trichlorophenyl group,  
2,3,4,5-tetrachlorophenyl group, 2,3,4,5,6-pentachlorophenyl group, 2,3-dibromophenyl  
group, 3,4-dibromophenyl group, 2,4-dibromophenyl group, 2,3,4-tribromophenyl group,  
3,4,5-tribromophenyl group, 2,3,4,5-tetrabromophenyl group, 2,3,4,5,6-pentabromo phenyl  
5 group. A 3,4-difluorophenyl group is preferred.

Moreover, \* denotes an asymmetric carbon centre. In other words, the halohydrin  
derivative represented by general formula (15) contains asymmetric carbon centre. The  
invention includes any optically active substance or racemic mixture of the compound of  
10 formula (15). Preferably it is optically active substance, and most preferably it is a  
compound whose absolute configuration of asymmetric carbon centre is (S).

In the compound of formula (16),  $R^8$  denotes an optionally substituted 1-10C cyclic or  
acyclic alkyl group, optionally substituted 6-10C aryl group or optionally substituted 7-  
15 10C aralkyl group, and  $R^9$  denotes an optionally substituted 1-10C cyclic or acyclic alkyl  
group, optionally substituted 6-10C aryl group or optionally substituted 7-10C aralkyl  
group. Suitable values for a 1-10C cyclic or acyclic alkyl group include for example, a  
methyl group, ethyl group, n-propyl group, i-propyl group, cyclopropyl group, n-butyl  
group, s-butyl group, i-butyl group, t-butyl group, cyclobutyl group, n-pentyl group,  
20 neopentyl group, cyclopentyl group, n-hexyl group, cyclohexyl group, n-heptyl group,  
cyclohexylmethyl group, n-octyl group, n-decyl group. Suitable values for an optionally  
substituted 6-10C aryl group include for example phenyl group, o-methoxyphenyl group,  
m-methoxyphenyl group, p-methoxy phenyl group, o-nitrophenyl group, m-nitrophenyl  
group, p-nitrophenyl group, o-chlorophenyl group, m-chlorophenyl group, p-chlorophenyl  
25 group, o-methylphenyl group, m-methylphenyl group, p-methylphenyl group. Suitable  
values for an optionally substituted 7-10C aralkyl group include for example a, benzyl  
group, o-methoxybenzyl group, m-methoxybenzyl group, p-methoxybenzyl group, o-  
nitrobenzyl, m-nitrobenzyl, p-nitrobenzyl, o-chlorobenzyl group, m-chlorobenzyl group, p-  
chlorobenzyl group, o-methylbenzyl group, m-methylbenzyl group, p-methylbenzyl group.

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In particular one or both of  $R^8$  and  $R^9$  are methyl group or ethyl group, and preferably  
both of  $R^8$  and  $R^9$  are methyl group or ethyl group.

In the compound of formula (13), values of substituents  $R^7$ ,  $R^8$  originate from respective values in the styrene oxide derivative of formula (14) or a halohydrin derivative represented by the formula (15) and carboxylate ester derivative represented by general formula (16). In other words,  $R^7$  denotes an aryl group substituted by 2 or more halogen atoms, and  $R^8$  denotes optionally substituted 1-10C cyclic or acyclic alkyl group, optionally substituted 6-10C aryl group or optionally substituted 7-10C aralkyl group, and  $R^9$  denotes optionally substituted 1-10C cyclic or acyclic alkyl group, optionally substituted 6-10C aryl group or optionally substituted 7-10C aralkyl group. Suitable values for an aryl group substituted by 2 or more halogen atoms, include, for example, a 2,3-difluorophenyl group, 3,4-difluorophenyl group, 2,4-difluorophenyl group, 2,3,4-trifluorophenyl group, 3,4,5-trifluorophenyl group, 2,3,4,5-tetrafluorophenyl group, 2,3,4,5,6-pentafluorophenyl group, 2,3-dichlorophenyl group, 3,4-dichlorophenyl group, 2,4-dichlorophenyl group, 2,3,4-trichlorophenyl group, 3,4,5-trichlorophenyl group, 2,3,4,5-tetrachlorophenyl group, 2,3,4,5,6-pentachlorophenyl group, 2,3-dibromophenyl group, 3,4-dibromophenyl group, 2,4-dibromophenyl group, 2,3,4-tribromophenyl group, 3,4,5-tribromophenyl group, 2,3,4,5-tetrabromophenyl group or 2,3,4,5,6-pentabromophenyl group. Suitable values for a 1-10C cyclic or acyclic alkyl group, include, for example, a methyl group, ethyl group, n-propyl group, i-propyl group, cyclopropyl group, n-butyl group, s-butyl group, i-butyl group, t-butyl group, cyclobutyl group, n-pentyl group, neopentyl group, cyclopentyl group, n-hexyl group, cyclohexyl group, n-heptyl group, cyclohexylmethyl group, n-octyl group or n-decyl group. Suitable values for an optionally substituted 6-10C aryl group include, for example, a phenyl group, o-methoxyphenyl group, m-methoxyphenyl group, p-methoxyphenyl group, o-nitrophenyl group, m-nitrophenyl group, p-nitrophenyl group, o-chlorophenyl group, m-chlorophenyl group, p-chlorophenyl group, o-methylphenyl group, m-methylphenyl group or p-methylphenyl group. Suitable values for an optionally substituted 7-10C aralkyl group include, for example, a benzyl group, o-methoxybenzyl group, m-methoxybenzyl group, p-methoxybenzyl group, o-nitrobenzyl, m-nitrobenzyl, p-nitrobenzyl, o-chlorobenzyl group, m-chlorobenzyl group, p-chlorobenzyl group, o-methylbenzyl group, m-methylbenzyl group or p-methylbenzyl group. It is generally preferred that  $R^7$  is a 3,4-difluorophenyl group and  $R^8$  is an ethyl group.

Moreover, \* denotes an asymmetric carbon centre. In other words, an ester derivative represented by the formula (13) contains asymmetric carbon centres. The invention includes any optically active substance or racemic mixture of the compound of formula (13). Preferably it is an optically active substance, and most preferably it is a compound whose absolute configuration of asymmetric carbon centre is (1R, 2R).

The optically active halohydrin derivative represented by the formula (15) which is a starting material of this invention can be readily obtained, for example, by enantioselectively reacting a  $\alpha$ -halomethyl arylketone derivative obtained by reacting a benzene derivative with  $\alpha$ -halo acetic acid chloride in the presence of aluminum chloride. The optically active styrene oxide derivative of formula (14) can be readily obtained by epoxidation of an optically active  $\alpha$ -halohydrin derivative of formula (15).

A compound of formula (14) or of formula (15) is reacted with a compound of formula (16) in the presence of base and thereby converted to compound of formula (13).

Examples of suitable bases include, for example, an organolithium compound such as methyllithium, n-butyllithium, t-butyllithium, phenyl lithium or the like, a Grignard reagent such as n-butylmagnesiumchloride, methyl magnesium bromide or the like; an alkaline earth metal amide or alkali metal amide such as lithium amide, sodium amide, lithium diisopropyl amide, magnesium diisopropyl amide, lithium hexamethyl disilazide, sodium hexamethyl disilazide, potassium hexamethyl disilazide or the like; an alkali metal alkoxide such as sodium methoxide, sodium ethoxide, sodium-t-butoxide, lithium methoxide, lithium ethoxide, lithium-t-butoxide, potassium-t-butoxide or the like; an alkaline earth metal hydride or alkali metal hydride such as lithium hydride, sodium hydride, potassium hydride, calcium hydride or the like.

A base of an alkali metal-t-butoxide, alkali metal hydride or the like is generally preferred.

The quantity of base used differs depending on species of base used, species of solvent and reaction conditions. A particular quantity is a 1-5 fold molar ratio, preferably 1-3 fold molar ratio with respect to compound of formula (14) or (15).

The quantity of compound of formula (16) used differs depending on species of solvent and reaction conditions. A particular quantity is a 1-5 fold molar ratio, preferably 1-3 fold molar ratio with respect to compound of formula (14) or (15).

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In general a solvent is usually used in the reaction. Examples include, for example, dichloromethane, chloroform, dichloroethane, benzene, toluene, diethyl ether, ethylene glycol dimethylether, methyl-t-butyl ether, diisopropyl ether, tetrahydrofuran, 1,4-dioxane, N,N-dimethylformamide, N-methylpyrrolidone, 1,3-dimethyl imidazolidinone,  
10 dimethylsulfoxide, acetone, acetonitrile, ethyl acetate, isopropyl acetate ester, acetic acid-t-butyl, t-butanol and the like. The solvent may be used alone or as an admixture thereof, and in this case, the mixed proportions thereof are not restricted.

15

A solvent of toluene, ethylene glycol dimethylether, tetrahydrofuran or 1,4-dioxane is generally preferred.

Suitable values of the reaction temperature include values selected from the range of -30°C to boiling point of solvent used, and a temperature in the range of 20°C-90°C. Generally, the reaction time required is usually 30 mins to 24 hours.

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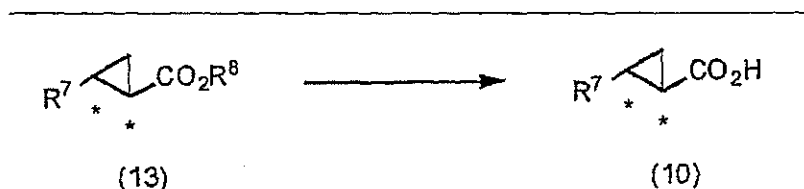
On completion of the reaction, solvent may be removed by distillation. The reaction mixture may then be added to water or water is added to it, and thereafter, it may be neutralized by addition of an appropriate quantity of acid. The compound of formula (13) may be obtained by using procedures such as extraction with an organic solvent such as  
25 toluene, ethyl acetate, isopropyl acetate, diethyl ether, dichloromethane, chloroform or the like, washing with water and concentration. The compound obtained may be purified further by column chromatography or distillation.

30

Examples of the acid used for neutralization after completion of the reaction include organic carboxylic acid such as formic acid, acetic acid, propionic acid, trifluoroacetic acid, chloroacetic acid, dichloroacetic acid, trichloroacetic acid, oxalic acid, benzoic acid, phthalic acid, fumaric acid, mandelic acid or the like; an optically active organic

carboxylic acid such as tartaric acid, lactic acid, ascorbic acid, amino acid or the like; an organic sulfonic acid such as methanesulfonic acid, trifluoromethanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, camphor sulfonic acid or the like; an inorganic acid such as hydrochloric acid, sulphuric acid, nitric acid, phosphoric acid,  
 5 carbonic acid or the like. Hydrochloric acid or sulfuric acid are generally preferred.

Next, there will be described 2) deesterification process.



## 10 Step 2. Deesterification process

The values of  $\text{R}^7$ ,  $\text{R}^8$  and \* in the compound of formula (13), including the suitable and preferred values, are the same as those mentioned above in 1) cyclopropanation process. In the compound of formula (10), the values of substituent  $\text{R}^7$  including the suitable and  
 15 preferred values, originate from the ester derivative of formula (13). In other words,  $\text{R}^7$  denotes an aryl group substituted by 2 or more halogen atoms. Suitable values of an aryl group substituted by 2 or more halogen atoms include, for example, a 2,3-difluorophenyl group, 3,4-difluorophenyl group, 2,4-difluorophenyl group, 2,3,4-trifluorophenyl group, 3,4,5-trifluorophenyl group, 2,3,4,5-tetrafluorophenyl group, 2,3,4,5,6-pentafluorophenyl  
 20 group, 2,3-dichlorophenyl group, 3,4-dichlorophenyl group, 2,4-dichlorophenyl group, 2,3,4-trichlorophenyl group, 3,4,5-trichlorophenyl group, 2,3,4,5-tetrachlorophenyl group, 2,3,4,5,6-pentachlorophenyl group, 2,3-dibromophenyl group, 3,4-dibromophenyl group, 2,4-dibromophenyl group, 2,3,4-tribromophenyl group, 3,4,5-tribromophenyl group, 2,3,4,5-tetrabromophenyl group or 2,3,4,5,6-pentabromophenyl group. A 3,4-  
 25 difluorophenyl group is generally preferred.

Moreover, \* denotes an asymmetric carbon centre. In other words the carboxylic acid derivative of formulae (10) contains asymmetric carbon centres. The invention includes any optically active substance or racemic mixture of the compound of formula (10).

Preferably it is an optically active substance, and most preferably it is a compound whose absolute configuration of asymmetric carbon centre is (1R, 2R).

In this step, the compound of formula (13) is converted to the compound of formula (10) by deesterifying, and reaction conditions of deesterification of Compound (13) are not restricted. The reaction may be carried out using general deesterification conditions. Examples of conditions for deesterification include a process of oxidative elimination of p-methoxybenzyl ester using DDQ (2,3-dichloro-5,6-dicyanobenzoquinone) and CAN (cerium nitrate), a process to eliminate benzyl ester, t-butyl ester using iodotrimethylsilane, a process of reductive elimination of benzyl ester using palladium catalyst under a hydrogen atmosphere, a process to eliminate t-butyl ester using TFA (trifluoroacetic acid), a process to eliminate ester group by acid or alkali hydrolysis, or the like. From the point of inexpensiveness and the point that the process can be applied for most kinds of ester group, the process to eliminate ester group by acid or alkali hydrolysis is preferred, and the process to eliminate ester group by alkali hydrolysis is more preferred.

Suitable alkalis include an alkali metal hydroxide such as lithium hydroxide, sodium hydroxide, potassium hydroxide, cesium hydroxide or the like; an alkaline earth metal hydroxide such as magnesium hydroxide, calcium hydroxide, barium hydroxide or the like; an alkali metal carbonate such as lithium carbonate, sodium carbonate, potassium carbonate, cesium carbonate or the like. An inorganic acid such as hydrochloric acid, sulphuric acid, nitric acid, phosphoric acid, perchloric acid or the like are generally preferred.

Suitable reaction solvents for deesterification include, for example, water, tetrahydrofuran, 1,4-dioxane, diethyl ether, methyl-t-butyl ether, toluene, benzene, N,N-dimethylformamide, dimethylsulfoxide, dichloromethane, chloroform, acetone, acetonitrile, butanol, propanol, ethanol, methanol, water and the like. The solvent may be used alone or as a mixture thereof, and in this case, the mixed proportions are not limited in particular.

In general a solvent of toluene, tetrahydrofuran, ethanol or methanol is preferred.

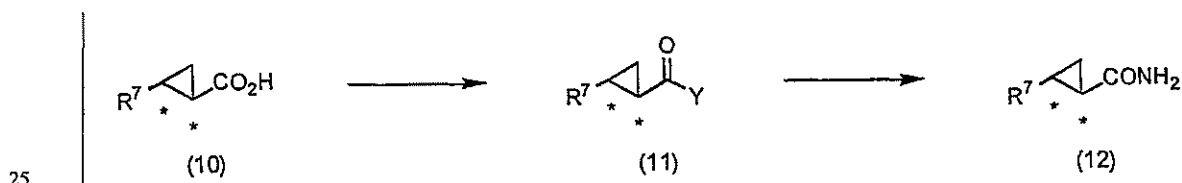
Suitable reaction temperatures, include those selected from the range of -30°C to boiling point of solvent used, and preferably it is 0°C-80°C. The reaction time is required usually to be 30 mins to 27 hours.

5

On completion of the reaction, the solvent may be removed by distillation, and thereafter the mixture added to water or water is added to it as required. The mixture is neutralized by addition of acid. The compound of formula (10) may be obtained by procedures such as extraction with an organic solvent such as toluene, ethyl acetate, isopropyl acetate,  
 10 diethyl ether, dichloromethane, chloroform or the like; washing with water, concentration and the like. The obtained compound may be further purified by column chromatography or crystallisation, or it may be used in the following step without treatment.

Suitable acids used for neutralization after completion of the reaction include, for example,  
 15 an organic carboxylic acid such as formic acid, acetic acid, propionic acid, trifluoroacetic acid, chloroacetic acid, dichloroacetic acid, trichloroacetic acid, oxalic acid, benzoic acid, phthalic acid, fumaric acid, mandelic acid or the like; an optically active organic carboxylic acid such as tartaric acid, lactic acid, ascorbic acid, amino acid or the like; an organic sulfonic acid such as methanesulfonic acid, trifluoromethanesulfonic acid,  
 20 benzenesulfonic acid, p-toluenesulfonic acid, camphor sulfonic acid or the like; an inorganic acid such as hydrochloric acid, sulphuric acid, nitric acid, phosphoric acid, carbonic acid or the like. Hydrochloric acid and sulfuric acid are generally preferred.

Next, a description will be given of 3) amidation process.



### Step 3. Amidation process

In the compound of formula (10), the values of substituent  $\text{R}^7$  and \* (including the suitable and preferred values) are the same as those mentioned above in 2) deesterification process.

In the compound of formula (11), values of substituent  $R^7$  originate from the ester derivative of formula (10). In other words,  $R^7$  may represent an aryl group substituted by 2 or more halogen atoms. Suitable values for an aryl group substituted by 2 or more halogen atoms include a 2,3-difluorophenyl group, 3,4-difluorophenyl group, 2,4-difluorophenyl group, 2,3,4-trifluorophenyl group, 3,4,5-trifluorophenyl group, 2,3,4,5-tetrafluorophenyl group, 2,3,4,5,6-pentafluorophenyl group, 2,3-dichlorophenyl group, 3,4-dichlorophenyl group, 2,4-dichlorophenyl group, 2,3,4-trichlorophenyl group, 3,4,5-trichlorophenyl group, 2,3,4,5-tetrachlorophenyl group, 2,3,4,5,6-pentachlorophenyl group, 2,3-dibromophenyl group, 3,4-dibromophenyl group, 2,4-dibromophenyl group, 2,3,4-tribromophenyl group, 3,4,5-tribromophenyl group, 2,3,4,5-tetrabromophenyl group, 2,3,4,5,6-pentabromo phenyl group or the like. A 3,4-difluorophenyl group is generally preferred. Moreover, Y denotes an activated carbonyl group activated group, and it is derived from the carboxylic acid activator described later.

Moreover, \* denotes an asymmetric carbon centre. In other words the carboxylic acid derivative of formula (11) contains asymmetric carbon centres. The invention includes any optically active substance or racemic mixture of the compound of formula (11). Preferably it is an optically active substance, and most preferably it is a compound whose absolute configuration of asymmetric carbon centre is (1R, 2R).

In the compound of formula (12), values of substituent  $R^7$  originate from the ester derivative of formula (10). In other words,  $R^7$  may denote an aryl group substituted by 2 or more halogen atoms. Suitable values for an aryl group substituted by 2 or more halogen atoms include a 2,3-difluorophenyl group, 3,4-difluorophenyl group, 2,4-difluorophenyl group, 2,3,4-trifluorophenyl group, 3,4,5-trifluorophenyl group, 2,3,4,5-tetrafluorophenyl group, 2,3,4,5,6-pentafluorophenyl group, 2,3-dichlorophenyl group, 3,4-dichlorophenyl group, 2,4-dichlorophenyl group, 2,3,4-trichlorophenyl group, 3,4,5-trichlorophenyl group, 2,3,4,5-tetrachlorophenyl group, 2,3,4,5,6-pentachlorophenyl group, 2,3-dibromophenyl group, 3,4-dibromophenyl group, 2,4-dibromophenyl group, 2,3,4-tribromophenyl group, 3,4,5-tribromophenyl group, 2,3,4,5-tetrabromophenyl group, 2,3,4,5,6-pentabromo phenyl group or the like. A 3,4-difluorophenyl group is generally preferred.

Moreover, \* denotes an asymmetric carbon centre. In other words, the carboxylic acid derivative of formula (12) contains asymmetric carbon centres. The invention includes any optically active substance or racemic mixture of the compound of formula (12).

5 Preferably it is an optically active substance, and most preferably it is a compound whose absolute configuration of asymmetric carbon centre is (1R, 2R).

The compound of formula (10) may be formed into the compound of formula (11) by reacting with a carboxylic acid activator to activate the carbonyl moiety. The activated  
10 compound is converted to the compound of formula (12) by reacting with ammonia. Suitable carboxylic acid activators include, for example, a dehydrocondensation agent such as dicyclohexylcarbodiimide (DCC) and carbonyldiimidazole; chlorocarbonic acid esters such as methyl chlorocarbonate ester, ethyl chlorocarbonate ester, propyl chlorocarbonate ester, isopropyl chlorocarbonate ester, chlorocarbonate butyl ester, t-butyl  
15 chlorocarbonate, benzyl chlorocarbonate or the like; an acid anhydride such as acetic anhydride, anhydrous trifluoroacetic acid, anhydrous methanesulfonic acid, anhydrous trifluoromethanesulfonic acid or the like; an carboxylic acid ester species such as carbonic acid di-t-butyl, dimethyl carbonate, diethyl carbonate or the like, acid chloride such as methanesulfonyl chloride, p-toluenesulphonyl chloride, phosphorus pentachloride,  
20 phosphorus trichloride, phosphorus oxychloride, acetyl chloride, propionyl chloride, pivaloyl chloride, benzoyl chloride, thionyl chloride, chlorosulfuric acid, oxalyl chloride; phosgene or the like, and a metal chloride such as titanium chloride, aluminum chloride, ferric chloride or the like may be proposed.

25 Particular carboxylic acid activators are chlorocarbonate ester, acid anhydride, carboxylic acid ester, acid chloride except phosgene. In general thionyl chloride is preferred particularly as it offers advantages from the point of handling and post-treatment after reaction.

30 The quantity used of carboxylic acid activator differs depending on species of base used and species of solvent and of reaction conditions. In particular a 1-3 fold molar ratio may

be used, and preferably a 1-1.5 fold molar ratio with respect to compound represented by the aforesaid formula (10).

When reacting the compound of formula (10) with the carboxylic acid activator, a base  
5 may be used in accordance with requirements. Suitable bases include, for example, an organolithium compound such as methyllithium, n-butyllithium, t-butyllithium, phenyl lithium or the like, a Grignard reagent such as n-butyl magnesium chloride, methyl magnesium bromide or the like, alkaline earth metal amide or alkali metal amide such as lithium amide, sodium amide, lithium diisopropyl amide, magnesium diisopropyl amide,  
10 lithium hexamethyl disilazide, sodium hexamethyl disilazide, potassium hexamethyl disilazide or the like, alkali metal alkoxide such as sodium methoxide, sodium ethoxide, sodium-t-butoxide, lithium methoxide, lithium ethoxide, lithium-t-butoxide, potassium-t-butoxide or the like, alkaline earth metal hydride or alkali metal hydride such as lithium hydride, sodium hydride, potassium hydride, calcium hydride or the like, alkaline earth  
15 metal hydroxide or alkali metal hydroxide such as lithium hydroxide, sodium hydroxide, potassium hydroxide, cesium hydroxide, magnesium hydroxide, calcium hydroxide or the like, alkali metal carbonate such as lithium carbonate, sodium carbonate, potassium carbonate or the like, alkali metal bicarbonate such as lithium bicarbonate, sodium bicarbonate, potassium bicarbonate or the like, organic tertiary amine such as  
20 triethylamine, diisopropyl ethylamine, DBU (1,8-diazabicyclo[5,4,0]undecene) or the like, basic organic solvent such as N,N-dimethylformamide or the like.

In particular the base may be an alkali metal alkoxide, alkaline earth metal hydride or alkali metal hydride, alkaline earth metal hydroxide or alkali metal hydroxide, alkaline  
25 earth carbonate or alkali metal carbonate, alkali metal bicarbonate, or organic tertiary amine. In general an alkaline earth metal hydroxide or alkali metal hydroxide, alkaline earth carbonate or alkali metal carbonate, alkali metal bicarbonate, organic tertiary amine or the like is preferred.

30 The quantity used of base differs depending on the species of base used and species of solvent and reaction conditions. In particular a 1-3 fold molar ratio may be used, and

preferably a 1-1.5 fold molar ratio with respect to compound represented by the aforesaid formula (10).

Suitable forms of the ammonia used include, for example, liquid ammonia, ammonia gas,  
5 ammonia solution in organic solvent and ammonia water. Particular examples are ammonia gas, ammonia in an organic solvent, ammonia water, and ammonia water is generally preferred.

When the form of ammonia is ammonia water the concentration of ammonia water used is  
10 not limited. In particular 5-30 wt % may be used, and 20-28 wt % is generally preferred.

The quantity of ammonia used differs depending on the form of used ammonia, species of solvent and reaction conditions. In particular a 1-6 fold molar ratio may be used, and preferably a 3-5 fold molar ratio with respect to compound represented by the aforesaid  
15 formula (10).

Generally a solvent is usually used in the reaction. Suitable solvents include for example dichloromethane, chloroform, dichloroethane, benzene, toluene, diethyl ether, methyl-t-butyl ether, diisopropyl ether, tetrahydrofuran, 1,4-dioxane, N,N-dimethylformamide, N-methylpyrrolidone, 1,3-dimethyl imidazolidinone, dimethylsulfoxide, acetone, acetonitrile,  
20 ethyl acetate, isopropyl acetate ester and acetic acid-t-butyl and the like.

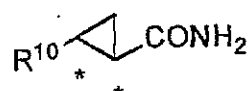
The solvent may be used alone or by mixing, and in this case, the mixing proportion is not limited. Generally a solvent of toluene, ethyl acetate and isopropyl acetate are preferred.  
25

Suitable reaction temperatures, include those selected from the range of -30°C to boiling point of solvent used and preferably it is selected from the range of 0°C-60°C. The reaction time required is usually 10 mins to 24 hours.

30 On completion of the reaction, the solvent is removed by distillation in accordance with requirements, and thereafter the reaction mixture is added to water or water is added to it. The compound of formula (12) is obtained using procedures such as extraction with an

organic solvent such as toluene, ethyl acetate, isopropyl acetate ester, diethyl ether, dichloromethane, chloroform or the like, washing with water, and concentration. The obtained compound may be further purified by column chromatography or crystallisation, or it may be used in the following step without treatment.

The compound of formula (17)

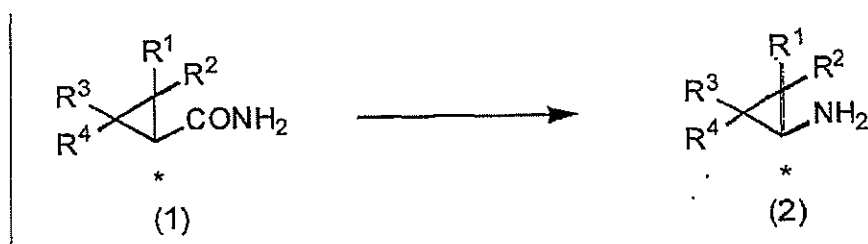


( 1 7 )

produced by the aforesaid process is a novel compound, and is therefore provided as a further feature of the present invention. In formula (17), R<sup>10</sup> denotes an aryl group substituted by 2 or more halogen atoms. Suitable values for the aryl group substituted by 2 or more halogen atoms include a 2,3-difluorophenyl group, 3,4-difluorophenyl group, 2,4-difluorophenyl group, 2,3,4-trifluorophenyl group, 3,4,5-trifluorophenyl group, 2,3,4,5-tetrafluorophenyl group, 2,3,4,5,6-pentafluorophenyl group, 2,3-dichlorophenyl group, 3,4-dichlorophenyl group, 2,4-dichlorophenyl group, 2,3,4-trichlorophenyl group, 3,4,5-trichlorophenyl group, 2,3,4,5-tetrachlorophenyl group, 2,3,4,5,6-pentachlorophenyl group, 2,3-dibromophenyl group, 3,4-dibromophenyl group, 2,4-dibromophenyl group, 2,3,4-tribromophenyl group, 3,4,5-tribromophenyl group, 2,3,4,5-tetrabromophenyl group, 2,3,4,5,6-pentabromo phenyl group or the like. A 3,4-difluorophenyl group is generally preferred.

Moreover, \* denotes an asymmetric carbon centre. In other words, the carboxamide derivative of formulae (17) contains asymmetric carbon centres. The invention includes any optically active substance or racemic mixture of the compound of formula (17). Preferably it is optically active substance, and most preferably it is a compound whose absolute configuration of asymmetric carbon centre is (1R, 2R).

Next, a description will be given of 4) Hofmann rearrangement process.



#### Step 4. Hofmann rearrangement step

In the compound of formula (1),  $\text{R}^1$ ,  $\text{R}^2$ ,  $\text{R}^3$  and  $\text{R}^4$  each independently denote hydrogen atom, optionally substituted 1-10C cyclic or acyclic alkyl group, optionally substituted 6-10C aryl group or optionally substituted 7-10C aralkyl group, and they may be the same or different to each other. Suitable values of an optionally substituted cyclic or acyclic alkyl group of carbon number 1-10 include a methyl group, ethyl group, n-propyl group, i-propyl group, cyclopropyl group, n-butyl group, s-butyl group, i-butyl group, t-butyl group, cyclobutyl group, n-pentyl group, neopentyl group, cyclopentyl group, n-hexyl group, cyclohexyl group, n-heptyl group, cyclohexylmethyl group, n-octyl group, n-decyl group and the like. Suitable values of an optionally substituted 6-10C aryl group include a phenyl group, o-methoxyphenyl group, m-methoxyphenyl group, p-methoxyphenyl group, o-nitrophenyl group, m-nitrophenyl group, p-nitrophenyl group, o-fluorophenyl group, m-fluorophenyl group, p-fluorophenyl group, o-chlorophenyl group, m-chlorophenyl group, p-chlorophenyl group, 2,3-difluorophenyl group, 3,4-difluorophenyl group, 2,4-difluorophenyl group, 2,3,4-trifluorophenyl group, 3,4,5-trifluorophenyl group, 2,3,4,5-tetrafluorophenyl group, 2,3,4,5,6-pentafluorophenyl group, 2,3-dichlorophenyl group, 3,4-dichlorophenyl group, 2,4-dichlorophenyl group, 2,3,4-trichlorophenyl group, 3,4,5-trichlorophenyl group, 2,3,4,5-tetrachlorophenyl group, 2,3,4,5,6-pentachlorophenyl group, 2,3-dibromophenyl group, 3,4-dibromophenyl group, 2,4-dibromophenyl group, 2,3,4-tribromophenyl group, 3,4,5-tribromophenyl group, 2,3,4,5-tetrabromophenyl group, 2,3,4,5,6-pentabromophenyl group, o-methylphenyl group, m-methylphenyl group, p-methylphenyl group and the like. Suitable values of an optionally substituted 7-10C aralkyl group include a benzyl group, o-methoxybenzyl group, m-methoxybenzyl group, p-methoxybenzyl group, o-nitrobenzyl group, m-nitrobenzyl group, p-nitrobenzyl group, o-chlorobenzyl group, m-chlorobenzyl group, p-chlorobenzyl group, o-methylbenzyl group, m-methylbenzyl group, p-methylbenzyl group and the like. Preferably any of  $\text{R}^1$ ,  $\text{R}^2$ ,  $\text{R}^3$

and R<sup>4</sup> is a 3,4-difluorophenyl group, and more preferably the substituent except 3,4-difluorophenyl group is a hydrogen atom.

Moreover, \* denotes an asymmetric carbon centre. In other words, the compound of formula (1) has asymmetric carbon centre. The invention includes any optically active substance or racemic mixture of the compound of formula (1). Preferably it is optically active substance, and most preferably it is a compound whose absolute configuration of asymmetric carbon centre is (1R, 2R).

In the compound of formula (2), values (including suitable and preferred) for R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> originate from the compound of formula (1). In other words, R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup> each independently denote a hydrogen atom, an optionally substituted 1-10C cyclic or an acyclic alkyl group, optionally substituted 6-10C aryl group or optionally substituted 7-10C aralkyl group, and they may be the same or different to each other. Suitable values for an optionally substituted 1-10C cyclic or acyclic alkyl group include a methyl group, ethyl group, n-propyl group, i-propyl group, cyclopropyl group, n-butyl group, s-butyl group, i-butyl group, t-butyl group, cyclobutyl group, n-pentyl group, neopentyl group, cyclopentyl group, n-hexyl group, cyclohexyl group, n-heptyl group, cyclohexylmethyl group, n-octyl group, n-decyl group and the like. Suitable values for an optionally substituted 6-10C aryl group include a phenyl group, o-methoxyphenyl group, m-methoxyphenyl group, p-methoxyphenyl group, o-nitrophenyl group, m-nitrophenyl group, p-nitrophenyl group, o-fluorophenyl group, m-fluorophenyl group, p-fluorophenyl group, o-chlorophenyl group, m-chlorophenyl group, p-chlorophenyl group, 2,3-difluorophenyl group, 3,4-difluorophenyl group, 2,4-difluorophenyl group, 2,3,4-trifluorophenyl group, 3,4,5-trifluorophenyl group, 2,3,4,5-tetrafluorophenyl group, 2,3,4,5,6-pentafluorophenyl group, 2,3-dichlorophenyl group, 3,4-dichlorophenyl group, 2,4-dichlorophenyl group, 2,3,4-trichlorophenyl group, 3,4,5-trichlorophenyl group, 2,3,4,5-tetrachlorophenyl group, 2,3,4,5,6-pentachlorophenyl group, 2,3-dibromophenyl group, 3,4-dibromophenyl group, 2,4-dibromophenyl group, 2,3,4-tribromophenyl group, 3,4,5-tribromophenyl group, 2,3,4,5-tetrabromophenyl group, 2,3,4,5,6-penta bromophenyl group, o-methylphenyl group, m-methylphenyl group, p-methylphenyl group and the like. Suitable values for an optionally substituted 7-10C aralkyl group include a benzyl

group, o-methoxybenzyl group, m-methoxybenzyl group, p-methoxybenzyl group, o-nitrobenzyl group, m-nitrobenzyl group, p-nitrobenzyl group, o-chlorobenzyl group, m-chlorobenzyl group, p-chlorobenzyl group, o-methylbenzyl group, m-methylbenzyl group, p-methylbenzyl group and the like. Wherein preferably any of  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  is a 3,4-difluorophenyl group, and more preferably, the substituent other than 3,4-difluorophenyl group is hydrogen atom.

Moreover, \* denotes an asymmetric carbon centre. In other words, the compound represented by the formula (2) has asymmetric carbon centre. The invention includes any optically active substance or racemic mixture of the compound of formula (2). Preferably it is optically active substance, and most preferably it is a compound whose absolute configuration of asymmetric carbon centre is (1R, 2S).

When oxidant is caused to act, there proceeds a so-called Hofmann rearrangement, and the compound of formula (1) is converted to the compound of formula (2) while maintaining the stereochemistry of the asymmetric carbon centre represented by \*. For example, suitable oxidants include a high valency iodine reagent exemplified by bis(trifluoroacetoxy) phenyl iodide, halide agent such as chlorine, bromine, iodine, N-chlorosuccinimide, N-bromosuccinimide, N-iodosuccinimide, sulphuryl chloride, sulphuryl bromide or the like, hypochlorite species such as lithium hypochlorite, sodium hypochlorite, potassium hypochlorite, magnesium hypochlorite, calcium hypochlorite or the like may be proposed, and chlorine, N-chloro succinimide, hypochlorite species or the like. In general sodium hypochlorite is preferred.

The quantity of oxidant used differs depending on species of oxidant used, species of reaction solvent and reaction conditions. In particular a 1-5 fold molar ratio may be used and preferably a 2-4 fold molar ratio with respect to the compound of formula (1). Moreover, as regards the quantity used of the aforesaid oxidant, when a hypochlorite species is used as the oxidant, the quantity used is determined by effective chlorine conversion.

In the reaction of compound of formula (1) and oxidant, a base may be co-present in accordance with requirements. Base may be added after mixing the compound of formulae (1) and oxidant. Suitable bases include, for example, an organolithium compound such as methyllithium, n-butyllithium, t-butyllithium, phenyl lithium or the like, Grignard reagent  
5 such as n-butyilmagnesium chloride, methyl magnesium bromide or the like, alkaline earth metal amide or alkali metal amide such as lithium amide, sodium amide, lithium diisopropyl amide, magnesium diisopropyl amide, lithium hexamethyl disilazide, sodium hexamethyl disilazide, potassium hexamethyl disilazide or the like, alkali metal alkoxide such as sodium methoxide, sodium ethoxide, sodium-t-butoxide, lithium methoxide,  
10 lithium ethoxide, lithium-t-butoxide, potassium-t-butoxide or the like, alkaline earth metal hydride or alkali metal hydride such as lithium hydride, sodium hydride, potassium hydride, calcium hydride or the like, alkaline earth metal hydroxide or alkali metal hydroxide such as lithium hydroxide, sodium hydroxide, potassium hydroxide, cesium hydroxide, magnesium hydroxide, calcium hydroxide or the like, alkali metal carbonate  
15 such as lithium carbonate, sodium carbonate, potassium carbonate or the like, alkali metal bicarbonate such as lithium bicarbonate, sodium bicarbonate, potassium bicarbonate or the like, organic tertiary amine such as triethylamine, diisopropyl ethylamine, DBU (1,8-diazabicyclo[5,4,0]undecene) or the like.

20 In general an alkali metal hydroxide such as sodium hydroxide is the preferred.

The quantity of base used differs depending on species of base used, species of solvent and reaction conditions. In particular the reaction may be caused to proceed in high yield by using a 5-30 fold molar ratio, preferably 5-20 fold molar ratio with respect to compound  
25 represented by general formula (2).

In particular the concentration of the base in the reaction may be in the range of 5-30 wt %, more particularly in the range of 15-25 wt %.

30 In general a solvent is usually used in the reaction. Suitable solvents include, for example, water, dichloromethane, chloroform, dichloroethane, benzene, toluene, diethyl ether, methyl-t-butyl ether, tetrahydrofuran, 1,4-dioxane, N,N-dimethylformamide, N-

methylpyrrolidone, 1,3-dimethyl imidazolidinone, dimethylsulfoxide, acetone, acetonitrile, ethyl acetate, acetic acid-t-butyl, t-butanol and the like.

The solvent may be used alone or as a mixture. In the case of a mixture the proportion is  
5 not limited. In general, water is preferred.

Suitable reaction temperatures include those selected from the range of -30°C to boiling point of solvent used and preferably it is selected from the range of 20°C-60°C. The reaction time required is usually 30 mins to 24 hours.

10

On completion of the reaction the solvent may be removed by distillation. The reaction mixture may be added to water or water to it, and then the mixture is acidified by addition of acid. The Compound (2) is transferred to the aqueous layer, and after having been caused to undergo liquid separation and washing with organic solvent such as toluene,  
15 ethyl acetate, isopropyl acetate, diethyl ether, dichloromethane, chloroform or the like, the aqueous layer is made basic using a base. The Compound of formula (2) is obtained using procedures such as extraction with an organic solvent such as toluene, ethyl acetate, isopropyl acetate, diethyl ether, dichloromethane, chloroform or the like, washing with water and concentration. Usually, on completion of the reaction, solvent is removed by  
20 distillation, and the compound of formula (2) may be obtained via procedures such as extraction with organic solvent such as toluene, ethyl acetate, isopropyl acetate, diethyl ether, dichloromethane, chloroform or the like, washing with water and concentration without the step of transferring to the aqueous layer. The compound (2) may be obtained in the form of a salt of an acid. The compound may be further purified by column  
25 chromatography, distillation or crystallisation, or it may be separated and purified in the form of a salt of an acid.

Suitable acids used after completion of the reaction include, for example, an organic carboxylic acid such as formic acid, acetic acid, propionic acid, trifluoroacetic acid,  
30 chloroacetic acid, dichloroacetic acid, trichloroacetic acid, oxalic acid, benzoic acid, phthalic acid, fumaric acid, mandelic acid or the like, optically active organic carboxylic acid such as tartaric acid, lactic acid, ascorbic acid, amino acid or the like, organic sulfonic

acid such as methanesulfonic acid, trifluoromethanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, camphor sulfonic acid or the like, inorganic acid such as hydrochloric acid, sulphuric acid, nitric acid, phosphoric acid, carbonic acid. Hydrochloric acid or sulfuric acid are generally preferred.

5

Suitable bases used after completion of the reaction include, for example, an organolithium compound such as methyllithium, n-butyllithium, t-butyllithium, phenyl lithium or the like, Grignard reagent such as n-butyilmagnesium chloride, methyl magnesium bromide or the like, alkaline earth metal amide or alkali metal amide such as lithium amide, sodium  
10 amide, lithium diisopropyl amide, magnesium diisopropyl amide, lithium hexamethyl disilazide, sodium hexamethyl disilazide, potassium hexamethyl disilazide or the like, alkali metal alkoxide such as sodium methoxide, sodium ethoxide, sodium-t-butoxide, lithium methoxide, lithium ethoxide, lithium-t-butoxide, potassium-t-butoxide or the like, alkaline earth metal hydride or alkali metal hydride such as lithium hydride, sodium  
15 hydride, potassium hydride, calcium hydride or the like, alkaline earth metal hydroxide or alkali metal hydroxide such as lithium hydroxide, sodium hydroxide, potassium hydroxide, cesium hydroxide, magnesium hydroxide, calcium hydroxide or the like, alkali carbonate metal salt such as lithium carbonate, sodium carbonate, potassium carbonate or the like, alkali metal bicarbonate such as lithium bicarbonate, sodium bicarbonate, potassium  
20 bicarbonate or the like, organic tertiary amine or the like such as triethylamine, diisopropyl ethylamine, DBU (1,8-diazabicyclo[5,4,0]undecene).

In general an alkali metal hydroxide, alkaline earth metal hydroxide, alkali carbonate metal salt, alkaline earth metal carbonate, alkali metal bicarbonate alkaline earth metal  
25 carbonate, organic tertiary amine are preferred.

### Examples

Below this invention will be described in greater detail by reference to Examples.

30 However, this invention is not restricted only to these Examples.

**Example 1****Preparation of (2*S*)-2-(3,4-difluorophenyl)oxirane**

A mixture of (1*S*)-2-chloro-1-(3,4-difluorophenyl)-1-ethanol (net 11.47 g, 59.5 mmol), toluene (25.23 g), sodium hydroxide (2.53 g, 1.06 molar equivalents) and water (24.25 g) was stirred and heated at 40°C for 1 hour. The organic layer was separated, washed with water, and concentrated under reduced pressure. (2*S*)-2-(3,4-difluorophenyl)oxirane was obtained as resultant concentrate (net 8.94 g, yield: 96%).

<sup>1</sup>H-NMR in (400MHz, CDCl<sub>3</sub>)

δ 2.71 - 2.73(1H, dd, J=2.44Hz, 5.37Hz), 3.13 - 3.15(1H, m), 3.82 - 3.83(1H, m), 7.01 - 7.27(4H, m).

**Example 2****Preparation of ethyl (1*R*, 2*R*)-2-(3,4-difluorophenyl)-1-cyclopropanecarboxylate**

Sodium t-butoxide (32.22 g, 1.25 molar equivalents) and toluene (243.0 g) were charged into a reaction vessel. Triethyl phosphonoacetate (78.06 g, 1.04 molar equivalents to sodium t-butoxide) was added to the mixture with stirring. A toluene solution of (2*S*)-2-(3,4-difluorophenyl) oxirane (32.8 wt % solution, net 41.83 g, 267.9 mmol) was added drop-wise to the mixture keeping the internal temperature between 60 to 80 °C. After completion of addition, stirring was continued for 11 hours at 80°C. After cooling to room temperature, the mixture was washed with water, and the organic layer was concentrated under reduced pressure. Ethyl (1*R*, 2*R*)-2-(3,4-difluorophenyl)-1-cyclopropanecarboxylate was obtained as resultant concentrate (net 49.11 g, yield: 81%).

<sup>1</sup>H-NMR in (400MHz, CDCl<sub>3</sub>)

δ 1.22 - 1.26(1H, m), 1.26 - 1.30(3H, t, J=7.1Hz), 1.57 - 1.62(1H, m), 1.82 - 1.87(1H, m), 2.45 - 2.50(1H, m), 4.14 - 4.20(2H, q, J=7.1Hz), 6.82 - 6.91(2H, m), 7.02 - 7.09(1H, m)

**Example 3****Preparation of (1*R*, 2*R*)-2-(3,4-difluorophenyl)-1-cyclopropanecarboxylic acid**

Methanol (322.2 g) and 30% sodium hydroxide aqueous solution (65.5 g, 1.8 molar equivalents) were added to a solution of ethyl (1*R*, 2*R*)-2-(3,4-difluorophenyl)-1-cyclopropanecarboxylate (48.2 wt% toluene solution, net 61.22 g, 270.6 mmol). The mixture was heated at 65°C with stirring for 2 hours. The resultant mixture was

concentrated under reduced pressure, then toluene and water were added to the concentrate. The mixture was acidified with 35% hydrochloric acid. The organic layer was separated and concentrated under reduced pressure. (1*R*, 2*R*)-2-(3,4-difluorophenyl)-1-cyclopropanecarboxylic acid was obtained as resultant concentrate (net 52.55 g, yield: 98%).

<sup>1</sup>H-NMR in (400MHz, CDCl<sub>3</sub>)

δ 1.33 - 1.38(1H, m), 1.64 - 1.69(1H, m), 1.83 - 1.88(1H, m), 2.54 - 2.59(1H, m), 6.83 - 6.93(2H, m), 7.04 - 7.10(1H, m).

#### Example 4

##### Preparation of (1*R*, 2*R*)-2-(3,4-difluorophenyl)-1-cyclopropanecarboxamide

Thionyl chloride (72.65g, 1.21 molar equivalents) was added to the stirred toluene solution of (1*R*, 2*R*)-2-(3,4-difluorophenyl)-1-cyclopropanecarboxylic acid (18wt%, net 100.00g, 504.62 mmol). The mixture was stirred at 35°C for 6 hours, then concentrated under reduced pressure to give a solution of (1*R*, 2*R*)-2-(3,4-difluorophenyl)-1-cyclopropanecarbonyl chloride. To a mixture of 28% ammonia aqueous solution (122.55g, 4.00 molar equivalents), water (300.4g) and ethyl acetate (700.2g), the solution of (1*R*, 2*R*)-2-(3,4-difluorophenyl)-1-cyclopropanecarbonyl chloride obtained above was gradually added with stirring below 10°C. The reaction mixture was allowed to stir below 10°C for 1 hour. The mixture was neutralized with 35% hydrochloric acid, then the organic layer was separated and washed with water. The resultant solution was concentrated azeotropically under reduced pressure to give a slurry of (1*R*, 2*R*)-2-(3,4-difluorophenyl)-1-cyclopropanecarboxamide. The resultant slurry was heated to obtain a clear solution, and cooled for crystallization. Hexane was added to the slurry, then the precipitates were collected by filtration and dried to give (1*R*, 2*R*)-2-(3,4-difluorophenyl)-1-cyclopropanecarboxamide (net 91.12 g, Yield: 92%).

<sup>1</sup>H-NMR in (400MHz, CDCl<sub>3</sub>)

δ 1.21 - 1.27(1H, m), 1.56 - 1.64(3H, m), 2.47 - 2.49(1H, m), 5.45(1H, br), 5.63(1H, br), 6.83 - 6.90(2H, m), 7.03 - 7.10(1H, m).

**Example 5****Preparation of (1*R*, 2*S*)-2-(3,4-difluorophenyl)-1- cyclopropanamine**

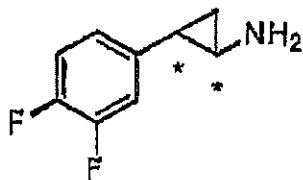
(1*R*, 2*R*)-2-(3,4-difluorophenyl)-1-cyclopropanecarboxamide (net 9.00g, 45.64 mmol) and 30% sodium hydroxide aqueous solution (54.77g, 9.00 molar equivalents) were charged  
5 into a reaction vessel and the mixture was stirred. Aqueous 12% sodium hypochlorite solution (29.53g, 2.25 mol equivalents) was added to the stirred slurry maintaining the internal temperature at 30°C. The resultant mixture was stirred at 30°C for 14 hours, then at 40 °C for 2 hours. After completion of the reaction, isopropyl acetate was poured to the resultant mixture, then the organic layer was separated, washed with water, and  
10 concentrated under reduced pressure. (1*R*, 2*S*)- 2-(3,4-difluorophenyl)-1-cyclopropanamine was obtained as resultant concentrate (net 6.89 g, yield: 89%).

<sup>1</sup>H-NMR in (400MHz, CDCl<sub>3</sub>)

δ 0.88 - 0.93(1H, m), 1.03 - 1.08(1H, m), 1.70(2H, s), 1.79 - 1.84(1H, m), 2.47 - 2.51(1H, m), 6.72 - 6.79(2H, m), 7.00 - 7.02(1H, m).

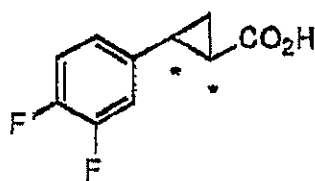
**Claims**

1. A process for the production of an optically active 2-aryl cyclopropylamine derivative represented by general formula (9) or a salt thereof



( 9 )

(wherein \* denotes an asymmetric carbon centre), wherein an optically active 2-aryl

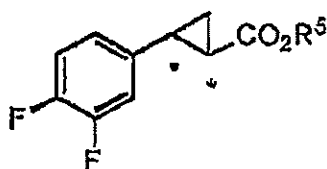


( 7 )

cyclopropanecarboxylic acid derivative represented by general formula (7)

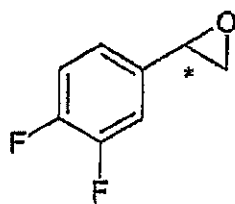
(wherein \* denotes an asymmetric carbon centre) is obtained,

by de-esterifying an optically active 2-aryl cyclopropane carboxylate ester derivative represented by general formula (6)



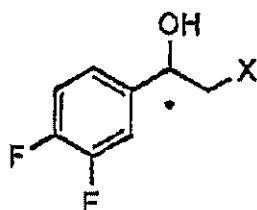
( 6 )

(wherein R<sup>5</sup> denotes optionally substituted 1-10C cyclic or acyclic alkyl group, optionally substituted 6-10C aryl group or optionally substituted 7-10C aralkyl group, and \* denotes an asymmetric carbon centre), wherein the compound of formula (6) is obtained by reacting an optically active styrene oxide derivative represented by general formula (3)



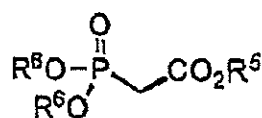
( 3 )

(wherein \* denotes an asymmetric carbon centre) or optically active halohydrin derivative represented by general formula (4)



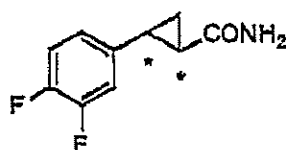
( 4 )

(wherein X denotes a halogen atom, and \* denotes an asymmetric carbon centre), with a phosphonoacetic acid ester derivative represented by general formula (5)



( 5 )

(wherein, R<sup>5</sup> or R<sup>6</sup> denote optionally substituted 1-10C cyclic or acyclic alkyl group, optionally substituted 6-10C aryl group or optionally substituted 7-10C aralkyl group) in the presence of base, and thereafter the 2-aryl cyclopropanecarboxylic acid derivative of formula (7) obtained is activated by reaction with a carboxylic acid activator and thereafter reacting with ammonia to give the optically active 2-aryl cyclopropane carboxamide derivative represented by obtained general formula (8)



( 8 )

(wherein \* denotes an asymmetric carbon centre), which is reacted with an oxidant to give the compound of formula (9).

2. A process as claimed in Claim 1, wherein the compound of formula (3) is a (S)-styrene oxide derivative, the compound of formula (4) is a (S)-halohydrin derivative and a (1R, 2S)-2-aryl cyclopropylamine derivative of formula (9) is prepared.