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(71) Applicant: SUMITOMO CHEMICAL COMPANY, LIMITED [JP/JP]; 27-1, Shinkawa 2-chome, Chuo-ku, Tokyo, 1048260 (JP).

(72) Inventors: YANAGISAWA, Kazuyuki; c/o SUMITOMO CHEMICAL COMPANY, LIMITED, 2-1, Takatsukasa 4-chome, Takarazuka-shi, Hyogo, 6658555 (JP). MIZUTANI, Motofumi; c/o SUMITOMO CHEMICAL COMPANY, LIMITED, 27-1, Shinkawa 2-chome, Chuo-ku, Tokyo, 1048260 (JP).

(74) Agents: NAKAYAMA, Tohru et al.; c/o Sumitomo Chemical Intellectual Property Service, Limited, 5-33, Ki-

tahama 4-chome, Chuo-ku, Osaka-shi, Osaka, 5418550 (JP).

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(54) Title: PESTICIDAL COMPOSITION IN THE FORM OF AQUEOUS EMULSION

(57) Abstract: This invention relates to a pesticidal composition in the form of an aqueous emulsion, comprising: (a) an agriculturally active ingredient in which a retention rate is less than 50% after irradiation with xenon light (290 nm cutoff) at an intensity of 0.68 W/m² at 340 nm for 8 hours; (b) a hydrophobic organic solvent; (c) a water-soluble polymer dispersing agent; and (d) water; wherein liquid droplets in which the agriculturally active ingredient is suspended or dissolved in the hydrophobic organic solvent are emulsified in an aqueous continuous phase, and also a weight ratio of the agriculturally active ingredient to the hydrophobic organic solvent is from 5:95 to 70:30. By the pesticidal composition of the present invention, it is possible to inhibit photodegradation of the agriculturally active ingredient.



DESCRIPTION

TITLE OF THE INVENTION

PESTICIDAL COMPOSITION IN THE FORM OF AQUEOUS EMULSION

5

Technical Field

[0001]

The present invention relates to a pesticidal composition in the form of an aqueous emulsion in which photodegradation of an agriculturally active ingredient has been inhibited.

10

BACKGROUND ART

[0002]

15

Various studies have hitherto been made for the purpose of inhibiting photodegradation of agriculturally active ingredients which are unstable against light. There have been known technologies in which stabilizers such as an ultraviolet absorber are added (see, for example, Patent Document 1). There is also known a pesticidal composition in the form of an aqueous emulsion, which contains a specific water-soluble polymer as a stabilizer in an aqueous continuous phase (see, for example, Patent Document 2).

20

[Prior Art Document]

25

[Patent Document]

[0003]

[Patent Document 1]: JP 59-39807 A

[Patent Document 2]: US 2003/0,134,910 A

DISCLOSURE OF INVENTION

5 [Problems to be Solved by the Invention]

[0004]

An object of the present invention is to provide a new
pesticidal composition in the form of an aqueous emulsion in
which photodegradation of an agriculturally active ingredient
10 has been inhibited.

[Means for Solving the Problems]

[0005]

The present inventors have intensively studied so as to
find a pesticidal composition in the form of an aqueous
15 emulsion in which photodegradation of an agriculturally
active ingredient has been inhibited, and found that, in a
pesticidal composition in the form of an aqueous emulsion,
comprising:

- (a) an agriculturally active ingredient in which a retention
20 rate is less than 50% after irradiation with xenon light (290
nm cutoff) at an intensity of 0.68 W/m^2 at 340 nm for 8 hours
(hereinafter sometimes referred to as a component a);
- (b) a hydrophobic organic solvent (hereinafter sometimes
referred to as a component b);
- 25 (c) a water-soluble polymer dispersing agent (hereinafter
sometimes referred to as a component c); and

(d) water (hereinafter sometimes referred to as a component d);

wherein liquid droplets in which the component a is suspended or dissolved in the component b are emulsified in an aqueous continuous phase, when a weight ratio of the component a to the component b is within a specific range, it is possible to inhibit photodegradation of the component a.

The present invention includes the followings:

[1] A pesticidal composition in the form of an aqueous emulsion, comprising:

(a) an agriculturally active ingredient in which a retention rate is less than 50% after irradiation with xenon light (290 nm cutoff) at an intensity of 0.68 W/m² at 340 nm for 8 hours;

(b) a hydrophobic organic solvent;

(c) a water-soluble polymer dispersing agent; and

(d) water;

wherein liquid droplets in which the agriculturally active ingredient is suspended or dissolved in the hydrophobic organic solvent are emulsified in an aqueous continuous phase, and also a weight ratio of the agriculturally active ingredient to the hydrophobic organic solvent is from 5:95 to 70:30;

[2] The pesticidal composition in the form of an aqueous emulsion according to [1], wherein the liquid droplets have an average particle diameter of 5 to 60 μm ;

[3] The pesticidal composition in the form of an aqueous emulsion according to [1] or [2], wherein the agriculturally active ingredient is an agriculturally active ingredient which is solid at normal temperature;

5 [4] The pesticidal composition in the form of an aqueous emulsion according to any one of [1] to [3], wherein the hydrophobic organic solvent(s) is/are one or more hydrophobic organic solvent(s) selected from the group consisting of esters, ketones, aromatic hydrocarbons, and paraffins; and

10 [5] The pesticidal composition in the form of an aqueous emulsion according to any one of [1] to [4], wherein the water-soluble polymer dispersing agent(s) is/are one or more water-soluble polymer dispersing agent(s) selected from the group consisting of a vinyl-based polymer, polysaccharides,
15 and derivatives thereof.

[Effects of the Invention]

[0006]

According to the present invention, it is possible to inhibit photodegradation of an agriculturally active
20 ingredient.

MODE FOR CARRYING OUT THE INVENTION

[0007]

The pesticidal composition in the form of an aqueous
25 emulsion of the present invention (hereinafter sometimes referred to as a composition of the present invention) is a

pesticidal composition in the form of an aqueous emulsion (emulsion pesticidal composition) containing an agriculturally active ingredient which is significantly unstable against light, such as the component a, and is a pesticidal composition which can effectively inhibit photodegradation of the agriculturally active ingredient.

[0008]

As used herein, xenon light means a light source in which a gas filled in a tube is xenon. Usually, xenon light means uniquely the same spectrum. Xenon light (290 nm cutoff) means xenon light having a wavelength within a range of 290 nm or more, and xenon light passed through a borosilicate filter corresponds to this.

A method for the calculation of a retention rate of the agriculturally active ingredient defined above is more specifically as follows. First, 250 mg of the agriculturally active ingredient is dissolved in 50 ml of a solvent such as acetone, which can sufficiently dissolve the agriculturally active ingredient and also has high volatility, and 1 mL of the solution is added to a glass petri dish having a diameter of 6 cm using a one-mark pipette, spread entirely over the petri dish, and then air-dried at room temperature. The petri dish is covered with a lid made of quartz glass and then set in a weathering tester (manufactured by Q-Lab Corporation under the trade name of Q-SUN Xenon Accelerated Weathering Tester, Model Xe-3) equipped with a borosilicate

filter (manufactured by Q-Lab Corporation under the trade name of Daylight-BB Optical Filter) attached thereto. After irradiation with xenon light under the conditions of an intensity at 340 nm of 0.68 W/m^2 and a temperature of 35°C (35°C as measured by an insulated black panel thermometer) for 8 hours, the amount of the agriculturally active ingredient remaining on the petri dish is determined by a known determination method such as high-performance liquid chromatography. It is possible to determine a retention rate by calculating as the weight percentage relative to 5 mg of the agriculturally active ingredient used above.

[0009]

The above agriculturally active ingredient used in the composition of the present invention may be any of an insecticidally active component, a fungicidally active component, and a herbicidally active component, and is not particularly limited. It is preferred to use an agriculturally active ingredient which is solid at normal temperature (25°C). The agriculturally active ingredient includes, for example, fenpyrazamine in which a retention rate after irradiation with xenon light (290 nm cutoff) at an intensity of 0.68 W/m^2 at 340 nm for 8 hours according to the method for calculation above-mentioned is 28.1%.

[0010]

The composition of the present invention usually contains a component a in the total amount of 0.5 to 25% by

weight, and preferably 2.5 to 15% by weight.

[0011]

In the composition of the present invention, a component b means an organic solvent which is liquid at normal temperature (25°C) and is immiscible in water. Specifically, an organic solvent having water solubility of 20% by weight or less at 25°C is used. Examples of the hydrophobic organic solvent include hydrophobic organic solvents such as vegetable oils, esters, ketones, aromatic hydrocarbons, and paraffins, for example, the following hydrophobic organic solvents:

[0012]

Vegetable oils: rapeseed oil, soybean oil, linseed oil, corn oil, and olive oil;

15 [0013]

Esters: diisobutyl adipate, diisodecyl adipate, dialkyl phthalate (didecyl phthalate, etc.), octyl oleate, lauryl oleate, octyldodecyl oleate, and isopropyl myristate;

[0014]

20 Ketones: methyl isobutyl ketone, heptanone, octanone, nonanone, cyclohexanone, and acetophenone; and

[0015]

Aromatic hydrocarbons: toluene, xylene, phenylxyllethane, 1-phenyl-1-ethylphenylethane, methylnaphthalene, dimethylnaphthalene, triisopropylbiphenyl, and dimethylisopropyl naphthalene.

In the present invention, commercially available aromatic hydrocarbon solvents can be used as aromatic hydrocarbons. Examples of the commercially available aromatic hydrocarbon solvents include Hisol SAS-296 (mixture of 1-phenyl-1-xylylethane and 1-phenyl-1-ethylphenylethane, trade name of JX Nippon Oil & Energy Corporation), Hisol SAS-LH (trade name of JX Nippon Oil & Energy Corporation), CACTUS SOLVENT HP-MN (methylnaphthalene 80%, trade name of JX Nippon Oil & Energy Corporation), CACTUS SOLVENT HP-DMN (dimethylnaphthalene of 80%, trade name of JX Nippon Oil & Energy Corporation), CACTUS SOLVENT P-180 (mixture of methylnaphthalene and dimethylnaphthalene, trade name of JX Nippon Oil & Energy Corporation), CACTUS SOLVENT P-200 (mixture of methylnaphthalene and dimethylnaphthalene, trade name of JX Nippon Oil & Energy Corporation), CACTUS SOLVENT P-220 (mixture of methylnaphthalene and dimethylnaphthalene, trade name of JX Nippon Oil & Energy Corporation), CACTUS SOLVENT PAD-1 (dimethylmonoisopropylnaphthalene, trade name of JX Nippon Oil & Energy Corporation), Solvesso 100 (aromatic hydrocarbon, trade name of ExxonMobil Chemical Ltd.), Solvesso 150 (aromatic hydrocarbon, trade name of ExxonMobil Chemical Ltd.), Solvesso 200 (aromatic hydrocarbon, trade name of ExxonMobil Chemical Ltd.), Solvesso 150ND (aromatic hydrocarbon, trade name of ExxonMobil Chemical Ltd.), Solvesso 200ND (aromatic hydrocarbon, trade name of ExxonMobil Chemical Ltd.), Ruetasolv BP 4302 (manufactured by

RKS GmbH), NIKANOL (trade name of Fudow Company Limited.), SWASOL 100 (toluene, trade name of Maruzen Petrochemical CO, LTD.), and SWASOL 200 (xylene, trade name of Maruzen Petrochemical CO, LTD.).

5 [0016]

Paraffins: normal paraffin, isoparaffin, cycloparaffin, and liquid paraffin

In the present invention, commercially available paraffin solvents can be used as paraffins. Examples of the commercially available paraffin solvents include NORPAR 13 (normal paraffin, producttrade name of ExxonMobil Chemical Ltd.), NORPAR 15 (normal paraffin, producttrade name of ExxonMobil Chemical Ltd.), ISOPAR E (isoparaffin, trade name of ExxonMobil Chemical Ltd.), ISOPAR G (isoparaffin, trade name of ExxonMobil Chemical Ltd.), ISOPAR L (isoparaffin, trade name of ExxonMobil Chemical Ltd.), ISOPAR H (isoparaffin, trade name of ExxonMobil Chemical Ltd.), ISOPAR M (isoparaffin, trade name of ExxonMobil Chemical Ltd.), MORESCO-WHITE P-40 (liquid paraffin, trade name of MORESCO Corporation), MORESCO-WHITE P-70 (liquid paraffin, trade name of MORESCO Corporation), MORESCO-WHITE P-200 (liquid paraffin, trade name of MORESCO Corporation), EXXSOL D110 (mixed solvent of paraffin and cycloparaffin, trade name of ExxonMobil Chemical Ltd.), EXXSOL D130 (mixed solvent of paraffin and cycloparaffin, trade name of ExxonMobil Chemical Ltd.), and EXXSOL D160 (mixed solvent of paraffin and

cycloparaffin, trade name of ExxonMobil Chemical Ltd.).

[0017]

There is no particular limitation on the component b used in the composition of the present invention, and one or
5 more hydrophobic organic solvent(s) selected from the group consisting of esters, ketones, aromatic hydrocarbons, and paraffins is/are preferably used, and aromatic ketones such as acetophenone are more preferably used.

[0018]

10 In the composition of the present invention, a weight ratio of the component a to the component b is from 5:95 to 70:30, preferably from 15:85 to 70:30, and more preferably from 30:70 to 70:30. The composition of the present invention usually contains the component b in the total
15 amount of 5 to 40% by weight, and preferably 10 to 25% by weight.

[0019]

There is no particular limitation on a component c used in the composition of the present invention, and one or more
20 water-soluble polymer dispersing agent(s) selected from the group consisting of a vinyl-based polymer, polysaccharides, and derivatives thereof is/are preferably used.

Examples of the water-soluble vinyl-based polymer include polyvinyl alcohol, polyvinylpyrrolidone, vinyl
25 acetate copolymer, and sodium polyacrylate. Examples of the water-soluble polysaccharides and derivatives thereof include

gum arabic, xanthan gum, rhamsan gum, locust bean gum, guar gum, carrageenan, welan gum, alginic acid, alginate, gum tragacanth, methyl cellulose, ethyl cellulose, sodium carboxymethyl cellulose, hydroxyethyl cellulose, and
5 hydroxypropyl cellulose. In the present invention, the vinyl-based polymer means a polymer obtained by addition polymerization of a monomer having a vinyl group ($-\text{CH}=\text{CH}_2$), and polysaccharides mean saccharides which are hydrolyzed to form two or more molecules of monosaccharides.

10 [0020]

The composition of the present invention usually contains the component c in the total amount of 1 to 10% by weight, and preferably 1 to 5% by weight.

[0021]

15 There is no particular limitation on a component d used in the composition of the present invention, and it is possible to use water which is used in a conventional pesticidal composition in the form of an aqueous emulsion (emulsion pesticidal composition), such as tap water, well
20 water, and deionized water, and deionized water is preferably used.

[0022]

The composition of the present invention usually contains the component d in the amount of 40 to 93% by weight,
25 and preferably 45 to 90% by weight. Substantially, the component c is dissolved in the component d.

[0023]

It is preferred that the composition of the present invention further contains one or more UV protectants. Examples of the UV protectants include benzophenone-based ultraviolet absorbers such as 2,4-dihydroxybenzophenone, 2-hydroxy-4-methoxybenzophenone, and 2-hydroxy-4-n-octylbenzophenone; benzotriazole-based ultraviolet absorbers such as 2-(2-hydroxy-5-methylphenyl)benzotriazole, 2-[2-hydroxy-3-(3,4,5,6-tetrahydrophthalimide-methyl)-5-methylphenyl]benzotriazole, 2-(3-tert-butyl-2-hydroxy-5-methylphenyl)-5-chlorobenzotriazole, 2-(2-hydroxy-5-tert-octylphenyl)benzotriazole, and 2-(2-hydroxy-3,5-di-tert-pentylphenyl)benzotriazole; benzoate-based ultraviolet absorbers such as 2,4-di-tert-butylphenyl 3,5-di-tert-butyl-4-hydroxybenzoate; hydroxyphenyltriazine-based ultraviolet absorber such as 2-(2-hydroxy-4-[1-octyloxycarbonylethoxy]phenyl)-4,6-bis(4-phenylphenyl)-1,3,5-triazine; photostabilizers such as non-sulfonated lignin (for example, manufactured by MeadWestvaco Corporation under the trade name of INDULIN AT) and a hindered amine-based photostabilizer (HALS); and inorganic UV shielding agents such as titanium oxide.

Substantially, the UV protectant (hereinafter sometimes referred to as a component e) is suspended or dissolved in the component b. There is no particular limitation on the component e used in the composition of the present invention,

and it is preferred to use a component e, which is miscible in the component b, from the viewpoint of emulsion stability of liquid droplets as the component b which contains the component e in the composition in the form of an aqueous
5 emulsion.

[0024]

When the composition of the present invention contains the component e, the total content is usually from 0.1 to 15% by weight, and preferably from 0.5 to 10% by weight.

10 [0025]

The composition of the present invention may further contain pesticide auxiliary agents used in a conventional pesticidal composition in the form of an aqueous emulsion. Examples of the pesticide auxiliary agents include
15 surfactants, thickeners, defoamers, preservatives, antifreezing agents, pH adjustors and the like.

[0026]

Examples of the surfactants include nonionic surfactants, cationic surfactants, anionic surfactants, and
20 amphoteric surfactants. Examples of nonionic surfactants include polyoxyethylene alkyl ether, polyoxyethylene alkyl aryl ether, polyoxyethylene lanolin alcohol, polyoxyethylene alkyl phenol formalin condensate, polyoxyethylene sorbitan fatty acid ester, polyoxyethylene glyceryl monofatty acid
25 ester, polyoxypropylene glycol monofatty acid ester, polyoxyethylene sorbitol fatty acid ester, polyoxyethylene

castor oil derivative, polyoxyethylene fatty acid ester, higher fatty acid glycerol ester, sorbitan fatty acid ester, sucrose fatty acid ester, polyoxyethylene polyoxypropylene block polymer, polyoxyethylene fatty acid amide, alkylol amide, polyoxyethylenealkylamine, and polyoxyethylene alkanediol. Examples of cationic surfactants include alkylamine hydrochlorides such as dodecylamine hydrochloride; alkyl quaternary ammonium salts such as dodecyltrimethyl ammonium salt, alkyldimethylbenzyl ammonium salt, alkylpyridinium salt, alkylisoquinolinium salt, and dialkylmorpholinium salt; benzethonium chloride, and polyalkyl vinyl pyridinium salt. Examples of anionic surfactants include fatty acid sodium such as sodium palmitate; sodium ether carboxylate such as sodium polyoxyethylene lauryl ether carboxylate; amino acid condensates of higher fatty acid, such as sodium lauroyl sarcosine and sodium N-lauroyl glutamate; higher fatty acid ester sulfonates such as higher alkyl sulfonate and lauric acid ester sulfonic acid salt; dialkyl sulfosuccinates such as dioctyl sulfosuccinate; higher fatty acid amide sulfonates such as oleic acid amide sulfonic acid; alkyl aryl sulfonates such as sodium dodecylbenzene sulfonate and diisopropyl naphthalene sulfonate; higher alcohol sulfuric acid ester salts such as formalin condensate of alkyl aryl sulfonate and pentadecane-2-sulfate; polyoxyethylene alkyl phosphates such as dipolyoxyethylene dodecyl ether phosphate; styrene-maleic

acid copolymer; and lignin sulfonate.

Examples of amphoteric surfactants include N-laurylalanine, N,N,N-trimethylaminopropionic acid, N,N,N-trihydroxyethylaminopropionic acid, N-hexyl-N,N-dimethylaminoacetic acid, 1-(2-carboxyethyl)pyridinium betaine, and lecithin.

When the composition of the present invention contains the surfactant, the total content is usually from 0.1 to 20% by weight, and preferably from 0.5 to 10% by weight.

10 [0027]

Examples of the thickener include natural polysaccharides such as xanthan gum, rhamsan gum, locust bean gum, guar gum, carrageenan, welan gum, alginic acid, alginate, and gum tragacanth; mineral matter powders such as aluminum silicate, magnesium aluminum silicate, smectite, bentonite, hectorite, synthetic hydrated silicic acid, and dry silica; and alumina sol. It is possible to use, as these thickeners, commercially available products as they are. Examples of commercially available products include KELZAN S (trade name of CP Kelco, Inc.) as xanthan gum, VEEGUM Granules (trade name of Vanderbilt Company, Inc.) as aluminum silicate, an Aerosil 200 (trade name of Evonik Degussa Corporation) as dry silica.

When the composition of the present invention contains the thickener, the total content is usually from 0.01 to 10% by weight, and preferably from 0.1 to 5% by weight.

[0028]

Examples of the defoamer include silicone-based defoamers such as ANTIFOAM C EMULSION (trade name of Dow Corning Toray Co., Ltd.), ANTIFOAM CE (trade name of Dow Corning Toray Co., Ltd.), TSA730 (trade name of MOMENTIVE PERFORMANCE MATERIALS JAPAN LLC), TSA731 (trade name of MOMENTIVE PERFORMANCE MATERIALS JAPAN LLC), TSA732 (trade name of MOMENTIVE PERFORMANCE MATERIALS JAPAN LLC), and YMA6509 (trade name of MOMENTIVE PERFORMANCE MATERIALS JAPAN LLC); and fluorine-based defoamers such as Fluowet PL80 (trade name of Clariant GmbH).

When the composition of the present invention contains the defoamer, the total content is usually from 0.01 to 3% by weight, and preferably from 0.05 to 1% by weight.

15 [0029]

Examples of the preservative include p-hydroxybenzoic acid ester, salicylic acid derivative, proxel (1,2-benzisothiazolin-3-one), and isothiazolin-3-one derivative (for example, BIOHOPE L (trade name of KI Chemical Industry Co., Ltd.)).

When the composition of the present invention contains the preservative, the total content is usually from 0.01 to 5% by weight, and preferably from 0.05 to 3% by weight.

[0030]

25 Examples of the antifreezing agent include water-soluble glycols such as ethylene glycol and propylene glycol.

When the composition of the present invention contains the antifreezing agent, the total content is usually from 0.5 to 30% by weight, and preferably from 1 to 20% by weight.

[0031]

5 Examples of the pH adjustor include citric acid monohydrate, sorbic acid, potassium sorbate, disodium hydrogen phosphate, dipotassium hydrogen phosphate, and sodium hydroxide.

When the composition of the present invention contains
10 the pH adjustor, the total content is usually from 0.01 to 5% by weight, and preferably from 0.5 to 3% by weight.

[0032]

A method for producing a composition of the present invention (hereinafter sometimes referred to as a present
15 production method) will be described, but the present production method is not limited only to the following methods.

[0033]

The present production method includes the following
20 first and second steps.

[0034]

The first step of the present production method is the step of suspending or dissolving a component a in a component b to prepare a suspension or a solution of the component a.

25 The component a is suspended or dissolved in the component b.

When the component a is a component a which is solid at normal temperature (25°C) (hereinafter sometimes referred to as a solid component a), it is possible to take the form of a suspension (hereinafter sometimes referred to as a present suspension a) depending on solubility of the solid component a in the component b and a weight ratio of the solid component a to the component b. The present suspension a can be prepared by finely grinding the solid component a in the component b using a wet grinding mill such as a beads mill. It is also possible to prepare by finely grinding the solid component a with or without adding other components using a dry grinding mill such as a jet mill, and adding the thus obtained finely ground powder to the component b.

The solution in which a component a is dissolved in a component b can be prepared by dissolving the component a in the component b.

[0035]

When the solid component a is dispersed (that is, suspended) in the component b in the form of fine particles, an average particle diameter of the fine particles is usually 15 μm or less, preferably 1 to 10 μm , and more preferably 1 to 5 μm .

[0036]

When the composition of the present invention contains the component e, the first step includes the step of respectively suspending or dissolving the component a and the

component e in the component b to prepare a suspension or a solution of the component a and the component e.

In the suspension or solution of the component a and the component e, the component a and the component e take any
5 one of the following forms (1) to (4).

(1) Both the component a and the component e are suspended in the component b.

(2) The component e is dissolved in the component b, and the component a is suspended in the component b in which the
10 component e has been dissolved.

(3) The component a is dissolved in the component b, and the component e is suspended in the component b in which the component a has been dissolved.

(4) Both the component a and the component e are dissolved in
15 the component b.

The solid component a can take the form (1) or (2) depending on solubility of the solid component a in the component b and a weight ratio of the solid component a to the component b. When the component e is a component e which
20 is solid at normal temperature (25°C) (hereinafter sometimes referred to as a solid component e), the component e can take the form (1) or (3) depending on solubility of the solid component e to the component b and a weight ratio of the solid component e to the component b.

25 [0037]

Suspensions of (1) to (3) can be prepared by the same

operation as in a method for preparing the present suspension a. In the case of preparing the suspension (1), the solid component a and the solid component e may be finely milled at the same time by a wet grinding mill or a dry grinding mill, 5 or the present suspension a and a suspension in which the solid component e has been suspended in the component b may be separately prepared, followed by mixing, in the method for preparing the present suspension a. In the case of preparing the suspension (2), usually, the component e is dissolved in 10 the component b and then the solid component a is added to the solution. In the case of preparing the suspension (3), usually, the component a is dissolved in the component b and then the solid component e is added to the solution.

The solution (4) can be prepared by dissolving the 15 component a and the component e in the component b.

[0038]

The second step of the present production method is the step of mixing the suspension or solution obtained in the first step with the aqueous solution of the component c to 20 prepare liquid droplets.

[0039]

In the aqueous solution of the component c used in the second step, the amount of the component c is usually from 0.5 to 30% by weight, and preferably from 1 to 20% by weight, 25 based on the entire amount of the aqueous solution of the component c. To the aqueous solution of the component c,

defoamers may be optionally added. The weight of the aqueous solution of the component c used in the second step is usually within a range of 0.8 to 2 times more than that of the suspension or solution obtained in the first step.

5 [0040]

In the second step, the method for preparing liquid droplets in water includes, for example, a method in which the suspension or solution obtained in the first step is added to the aqueous solution of the component c, followed by
10 stirring using a stirrer. The stirrer used in this case includes, for example, a propeller stirrer, a turbine stirrer, and a high-speed shear stirrer. Specific examples of the stirrer include T.K. Homo Mixer, T.K. Homomic Line Flow, T.K. Pipeline Homo Mixer, and T.K. Filmix manufactured by PRIMIX
15 Corporation; CLEARMIX manufactured by M Technique Co., Ltd.; POLYTRON homogenizer and MEGATRON homogenizer manufactured by KINEMATICA AG; and SUPRATON manufactured by Tsukishima Kikai Co., Ltd.

[0041]

20 An average particle diameter of liquid droplets existing in water obtained in the second step, that is, liquid droplets in the composition of the present invention, in which the component a is suspended or dissolved in the component b is usually from 5 to 60 μm , preferably
25 from 10 to 60 μm , and more preferably from 10 to 40 μm .

In the present invention, the average particle diameter

means a volume median diameter. The volume median diameter refers to a particle diameter at which a cumulative frequency in a volume equivalent frequency distribution is to be 50%, and the volume median diameter can be determined, for example, 5 by wet measurement using a laser diffraction particle size distribution measuring apparatus. More specifically, liquid droplets are dispersed in water and then the volume median diameter is measured using the apparatus. The laser diffraction particle size distribution measuring apparatus 10 includes, for example, Mastersizer 2000 (manufactured by Malvern Instruments Ltd.).

[0042]

It is possible to obtain a pesticidal composition in the form of an aqueous emulsion in which liquid droplets in 15 which the component a is suspended or dissolved in the component b are emulsified in an aqueous continuous phase, by the above production method. To the pesticidal composition in the form of an aqueous emulsion, the pesticide auxiliary agents and water may be further added.

20 [0043]

The composition of the present invention can be applied to places such as paddy fields, cultivated lands, orchards, grass plot, and non-agricultural lands in the same manner as in the case of a conventional pesticidal composition in the 25 form of an aqueous emulsion. The composition of the present invention is optionally diluted with water and then the

obtained water dilution can be applied by a method in which the composition is applied to plants growing in the above places or the soil in the above places. Examples of the method for applying the water dilution include a soil surface application or foliage application method of the water dilution using a known sprinkler.

It is also possible to use the water dilution in a seed treatment, a seedling raising box treatment and the like.

[0044]

The composition of the present invention can be applied as it is without being diluted with water and, for example, the composition of the present invention is applied along from levee to levee of paddy fields under flooding. Before application, the composition of the present invention is usually mixed by slightly shaking a vessel containing the composition of the present invention.

[Example]

[0045]

The present invention will be described in further detail below by way of Examples, but the present invention is not limited only to these Examples.

[0046]

First, Preparation Examples and Comparative Preparation Examples are shown.

[0047]

Preparation Example 1

Fenpyrazamine was dry-milled by a vertical type jet mill (JOM-0101-type jet mill, manufactured by Seishin Enterprise Co., Ltd.) to prepare fenpyrazamine fine powders having an average particle diameter 5 μm or less. Twenty
5 (20) parts by weight of the obtained fenpyrazamine fine powders were mixed with 10 parts by weight of acetophenone (manufactured by Wako Pure Chemical Industries, Ltd.) to obtain a fenpyrazamine suspension. Then, 30 parts by weight of the suspension was added to 33 parts by weight of an
10 aqueous 5% by weight polyvinyl alcohol (manufactured by The Nippon Synthetic Chemical Industry Co., Ltd. under the trade name of GOHSENOL GH-17) solution. This mixture was stirred by a rotor-stator homogenizer (manufactured by KINEMATICA AG under the trade name of POLYTRON homogenizer) thereby
15 emulsifying the fenpyrazamine suspension in the aqueous polyvinyl alcohol solution to obtain a fenpyrazamine emulsion.

To 14.2 parts by weight of deionized water, 0.4 part by weight of magnesium aluminum silicate (manufactured by Vanderbilt Company, Inc. under the trade name of VEEGUM
20 Granules) was added, followed by stirring at room temperature for 15 minutes. To this solution, 0.2 part by weight of xanthan gum (manufactured by CP Kelco, Inc. under the trade name of KELZAN S) and 5 parts by weight of propylene glycol were added. This mixture was stirred at 60°C for 60 minutes.
25 The obtained dispersion was cooled to room temperature, and then 0.2 part by weight of an preservative (manufactured by

Arch Chemicals, Inc. under the trade name of proxel GXL) was added to the dispersion to give a viscosity modifying liquid. Twenty (20) parts by weight of the viscosity modifying liquid, 17 parts by weight of water, and 63 parts by weight of the above fenpyrazamine emulsion were mixed together to obtain the composition (1) of the present invention, containing 20% by weight of fenpyrazamine. Liquid droplets of the fenpyrazamine suspension had an average particle diameter of 28.5 μm .

10 [0048]

Preparation Example 2

By performing the same operation as in Preparation Example 1, except that the amount of fenpyrazamine fine powders prepared in Preparation Example 1 was changed from 20 parts to 15 parts by weight, and the amount of acetophenone (the same as mentioned above) was changed from 10 parts to 15 parts by weight, the composition (2) of the present invention, containing 15% by weight of fenpyrazamine was obtained.

15 Liquid droplets of the fenpyrazamine suspension had an average particle diameter of 31.8 μm .

[0049]

Preparation Example 3

By performing the same operation as in Preparation Example 1, except that 30 parts by weight of a fenpyrazamine solution obtained by uniformly mixing 10 parts by weight of fenpyrazamine with 20 parts by weight of acetophenone (the

25

same as mentioned above) was used in place of 30 parts by weight of the fenpyrazamine suspension, the composition (3) of the present invention, containing 10% by weight of fenpyrazamine was obtained. Liquid droplets of the fenpyrazamine solution had an average particle diameter of 25.3 μm .

[0050]

Preparation Example 4

By performing the same operation as in Preparation Example 3, except that the amount of fenpyrazamine was changed from 10 parts by weight to 5 parts by weight, the amount of acetophenone (the same as mentioned above) was changed from 20 parts by weight to 25 parts by weight, and the aqueous 2.5% by weight polyvinyl alcohol (the same as mentioned above) solution was used in place of the aqueous 5% by weight polyvinyl alcohol (the same as mentioned above) solution, the composition (4) of the present invention, containing 5% by weight of fenpyrazamine was obtained. Liquid droplets of the fenpyrazamine solution had an average particle diameter of 29.3 μm .

[0051]

Preparation Example 5

By performing the same operation as in Preparation Example 3 performed, except that the amount of fenpyrazamine was changed from 10 parts by weight to 2.5 parts by weight, the amount of acetophenone (the same as mentioned above) was

changed from 20 parts by weight to 27.5 parts by weight, and the aqueous 1.0% by weight polyvinyl alcohol (the same as mentioned above) solution was used in place of the aqueous 5% by weight polyvinyl alcohol (the same as mentioned above) solution, the composition (5) of the present invention, containing 2.5% by weight of fenpyrazamine was obtained. Liquid droplets of the fenpyrazamine solution had an average particle diameter of 31.9 μm .

[0052]

10 Preparation Example 6

By performing the same operation as in Preparation Example 3, except that the average particle diameter of liquid droplets of the fenpyrazamine solution was changed from 25.3 μm to 44.0 μm in the case of stirring using the rotor-stator homogenizer (the same as mentioned above), the composition (6) of the present invention, containing 10% by weight of fenpyrazamine was obtained.

[0053]

Preparation Example 7

20 By performing the same operation as in Preparation Example 3, except that the average particle diameter of liquid droplets of the fenpyrazamine solution was changed from 25.3 μm to 14.9 μm in the case of stirring using the rotor-stator homogenizer (the same as mentioned above), the composition (7) of the present invention, containing 10% by weight of fenpyrazamine was obtained.

[0054]

Preparation Example 8

By performing the same operation as in Preparation Example 3, except that the average particle diameter of liquid droplets of the fenpyrazamine solution was changed from 25.3 μm to 7.8 μm in the case of stirring using the rotor-stator homogenizer (the same as mentioned above), the composition (8) of the present invention, containing 10% by weight of fenpyrazamine was obtained.

10 [0055]

Preparation Example 9

By performing the same operation as in Preparation Example 3, except that the aqueous 2.5% by weight polyvinyl alcohol (the same as mentioned above) solution was used in place of the aqueous 5% by weight polyvinyl alcohol (the same as mentioned above) solution, and the average particle diameter of liquid droplets of the fenpyrazamine solution was changed from 25.3 μm to 50.1 μm in the case of stirring using the rotor-stator homogenizer (the same as mentioned above), the composition (9) of the present invention, containing 10% by weight of fenpyrazamine was obtained.

[0056]

Preparation Example 10

By performing the same operation as in Preparation Example 1, except that the amount of fenpyrazamine fine powders prepared in Preparation Example 1 was changed from 20

parts by weight to 10 parts by weight, and 10 parts by weight of acetophenone (the same as mentioned above) was replaced by 20 parts by weight of liquid paraffin (manufactured by MORESCO Corporation under the trade name of MORESCO-WHITE
5 P-40), the composition (10) of the present invention, containing 10% by weight of fenpyrazamine was obtained. Liquid droplets of the fenpyrazamine suspension had an average particle diameter of 34.9 μm .

[0057]

10 Preparation Example 11

By performing the same operation as in Preparation Example 10, except that the average particle diameter of liquid droplets of the fenpyrazamine suspension was changed from 34.9 μm to 53.1 μm in the case of stirring using the
15 rotor-stator homogenizer (the same as mentioned above), the composition (11) of the present invention, containing 10% by weight of fenpyrazamine was obtained.

[0058]

Preparation Example 12

20 By performing the same operation as in Preparation Example 1, except that the amount of fenpyrazamine fine powders prepared in Preparation Example 1 was changed from 20 parts by weight to 10 parts by weight, and 10 parts by weight of acetophenone (the same as mentioned above) was replaced by
25 20 parts by weight of an aromatic hydrocarbon (manufactured by ExxonMobil Chemical Ltd. under the trade name of Solvesso

150ND), the composition (12) of the present invention, containing 10% by weight of fenpyrazamine was obtained. Liquid droplets of the fenpyrazamine suspension had an average particle diameter of 27.3 μm .

5 [0059]

Preparation Example 13

By performing the same operation as in Preparation Example 12, except that the average particle diameter of liquid droplets of the fenpyrazamine suspension was changed
10 from 27.3 μm to 42.6 μm in the case of stirring using the rotor-stator homogenizer (the same as mentioned above), the composition (13) of the present invention, containing 10% by weight of fenpyrazamine was obtained.

[0060]

15 Preparation Example 14

By performing the same operation as in Preparation Example 1, except that the amount of fenpyrazamine fine powders prepared in Preparation Example 1 was changed from 20 parts by weight to 10 parts by weight, and 10 parts by weight
20 of acetophenone (the same as mentioned above) was replaced by 20 parts by weight of diisobutyl adipate (manufactured by Kao Corporation under the trade name of Vinisizer 40), the composition (14) of the present invention, containing 10% by weight of fenpyrazamine was obtained. Liquid droplets of the
25 fenpyrazamine suspension had an average particle diameter of 30.1 μm .

[0061]

Preparation Example 15

By performing the same operation as in Preparation Example 3, except that the aqueous 5% by weight polyvinyl alcohol (the same as mentioned above) solution was replaced by an aqueous 20% by weight gum arabic (under the trade name of Arabiccol SS, manufactured by SAN-EI YAKUHIH BOEKI K.K.) solution, the composition (15) of the present invention, containing 10% by weight of fenpyrazamine was obtained. Liquid droplets of the fenpyrazamine solution had an average particle diameter of 21.3 μm .

[0062]

Preparation Example 16

By performing the same operation as in Preparation Example 1, except that the amount of fenpyrazamine fine powders prepared in Preparation Example 1 was changed from 20 parts by weight to 10 parts by weight, and 10 parts by weight of acetophenone (the same as mentioned above) was replaced by 20 parts by weight of isoparaffin (manufactured by ExxonMobil Chemical Ltd. under the trade name of ISOPAR E), the composition (16) of the present invention, containing 10% by weight of fenpyrazamine was obtained. Liquid droplets of the fenpyrazamine suspension had an average particle diameter of 31.0 μm .

[0063]

Preparation Example 17

Ten (10) parts by weight of fenpyrazamine, 19 parts by weight of acetophenone (the same as mentioned above), and 1 part by weight of a benzophenone-based ultraviolet absorber (manufactured by BASF Corporation under the trade name of Uvinul 3000) were mixed together to prepare a fenpyrazamine solution. Then, 30 parts by weight of the solution was added to 33 parts by weight of the aqueous 5% by weight polyvinyl alcohol (the same as mentioned above) solution. This mixture was stirred by the rotor-stator homogenizer (the same as mentioned above) thereby emulsifying the fenpyrazamine solution in the aqueous polyvinyl alcohol solution to obtain a fenpyrazamine emulsion.

To 14.2 parts by weight of deionized water, 0.4 part by weight of magnesium aluminum silicate (the same as mentioned above) was added, followed by stirring at room temperature for 15 minutes. To this solution, 0.2 part by weight of xanthan gum (the same as mentioned above) and 5 parts by weight of propylene glycol were added. This mixture was stirred at 60°C for 60 minutes. The obtained dispersion was cooled to room temperature, and then 0.2 part by weight of a preservative (the same as mentioned above) was added to the dispersion to give a viscosity modifying liquid. Twenty (20) parts by weight of the viscosity modifying liquid, 17 parts by weight of water, and 63 parts by weight of the above fenpyrazamine emulsion were mixed together to obtain the composition (17) of the present invention, containing 10% by

weight of fenpyrazamine. Liquid droplets of the fenpyrazamine solution had an average particle diameter of 17.8 μm .

[0064]

5 Preparation Example 18

By performing the same operation as in Preparation Example 17, except that 30 parts by weight of the fenpyrazamine suspension obtained by mixing 10 parts by weight of the fenpyrazamine fine powders prepared in

10 Preparation Example 1, 15 parts by weight of acetophenone (the same as mentioned above), and 5 parts by weight of a benzotriazole-based ultraviolet absorber (manufactured by BASF Corporation under the trade name of TINUVIN571) was used in place of 30 parts by weight of the fenpyrazamine solution,
15 the composition (18) of the present invention, containing 10% by weight of fenpyrazamine was obtained. Liquid droplets of the fenpyrazamine suspension had an average particle diameter of 28.4 μm .

[0065]

20 Comparative Preparation Example 1

By performing the same operation as in Preparation Example 3, except that the amount of fenpyrazamine was changed from 10 parts by weight to 0.8 part by weight, and the amount of acetophenone (the same as mentioned above) was
25 changed from 20 parts by weight to 29.2 parts by weight, the comparative composition (1), containing 0.8% by weight of

fenpyrazamine was obtained. Liquid droplets of the fenpyrazamine solution had an average particle diameter of 33.9 μm .

[0066]

5 Comparative Preparation Example 2

By performing the same operation as in Preparation Example 1, except that the amount of fenpyrazamine fine powders prepared in Preparation Example 1 was changed from 20 parts by weight to 25 parts by weight, and the amount of acetophenone (the same as mentioned above) was changed from 10 parts by weight to 5 parts by weight. As a result, viscosity of the fenpyrazamine suspension significantly increased, and thus a fenpyrazamine emulsion could not be produced.

15 [0067]

Next, Test Example will be shown.

[0068]

Test Example 1

The compositions (1) to (18) of the present invention, and the comparative composition (1) were respectively diluted with water so as to make each active ingredient concentration to 1,120 ppm. Each of the dilutions (100 μL) was added to a glass petri dish having a diameter of 6 cm, uniformly spread and then air-dried at room temperature. The petri dish was covered with a lid made of quartz glass and placed in a weathering tester (manufactured by Q-Lab Corporation under

the trade name of Q-SUN Xenon Accelerated Weathering Tester, Model Xe-3) equipped with a borosilicate filter (manufactured by Q-Lab Corporation under the trade name of Daylight-BB Optical Filter) attached thereto, followed by irradiation with xenon light under the conditions of an intensity at 340 nm of 0.68 W/m^2 and a temperature of 34°C (34°C as measured by an insulated black panel thermometer) for 6 hours. After irradiation, fenpyrazamine remaining on the petri dish was extracted with acetonitrile, followed by quantitative analysis through high-performance liquid chromatography. A retention rate was determined by calculating as the weight percentage relative to the amount of fenpyrazamine before irradiation.

The results are shown in Table 1.

15 [0069]

[Table 1]

Compositions used	Component b	Component e	Component a:Component b (weight ratio)	Average particle diameter (μm)	Retention rate (%)
Composition (1) of the present invention	Acetophenone	-	67:33	28.5	46.1
Composition (2) of the present invention	Acetophenone	-	50:50	31.8	49.2
Composition (3) of the present invention	Acetophenone	-	33:67	25.3	49.9
Composition (4) of the present invention	Acetophenone	-	17:83	29.3	37.2
Composition (5) of the present invention	Acetophenone	-	8:92	31.9	3.3
Composition (6) of the present invention	Acetophenone	-	33:67	44.0	46.1
Composition (7) of the present invention	Acetophenone	-	33:67	14.9	38.1
Composition (8) of the present invention	Acetophenone	-	33:67	7.8	8.6
Composition (9) of the present invention	Acetophenone	-	33:67	50.1	63.1
Composition (10) of the present invention	Liquid paraffin	-	33:67	34.9	13.1

[Table 1] (continued)

Compositions used	Component b	Component e	Component a:Component b (weight ratio)	Average particle diameter (μm)	Retention rate (%)
Composition (11) of the present invention	Liquid paraffin	-	33:67	53.1	22.8
Composition (12) of the present invention	Aromatic hydrocarbon	-	33:67	27.3	24.2
Composition (13) of the present invention	Aromatic hydrocarbon	-	33:67	42.6	58.2
Composition (14) of the present invention	Diisobutyl adipate	-	33:67	30.1	11.2
Composition (15) of the present invention	Acetophenone	-	33:67	21.3	24.7
Composition (16) of the present invention	Isoparaffin	-	33:67	31.0	24.3
Composition (17) of the present invention	Acetophenone	Benzophenone-based ultraviolet absorber	34:66	17.8	78.6
Composition (18) of the present invention	Acetophenone	Benzotriazole-based ultraviolet absorber	40:60	28.4	75.7
Comparative composition (1)	Acetophenone	-	3:97	33.9	0.6

CLAIMS

1. A pesticidal composition in the form of an aqueous emulsion, comprising:

- 5 (a) an agriculturally active ingredient in which a retention rate is less than 50% after irradiation with xenon light (290 nm cutoff) at an intensity of 0.68 W/m^2 at 340 nm for 8 hours;
- (b) a hydrophobic organic solvent;
- 10 (c) a water-soluble polymer dispersing agent; and
- (d) water;

wherein liquid droplets in which the agriculturally active ingredient is suspended or dissolved in the hydrophobic organic solvent are emulsified in an aqueous continuous phase, and also a weight ratio of the
15 agriculturally active ingredient to the hydrophobic organic solvent is from 5:95 to 70:30.

2. The pesticidal composition in the form of an aqueous emulsion according to claim 1, wherein the liquid droplets have an average particle diameter of 5 to 60 μm .
20

3. The pesticidal composition in the form of an aqueous emulsion according to claim 1 or 2, wherein the agriculturally active ingredient is an agriculturally active ingredient which is solid at normal temperature.
25

4. The pesticidal composition in the form of an aqueous emulsion according to any one of claims 1 to 3, wherein the hydrophobic organic solvent(s) is/are one or more hydrophobic organic solvent(s) selected from the group consisting of esters, ketones, aromatic hydrocarbons, and paraffins.

5. The pesticidal composition in the form of an aqueous emulsion according to any one of claims 1 to 4, wherein the water-soluble polymer dispersing agent(s) is/are one or more water-soluble polymer dispersing agent(s) selected from the group consisting of a vinyl-based polymer, polysaccharides, and derivatives thereof.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP2013/067585

A. CLASSIFICATION OF SUBJECT MATTER		
Int.Cl. A01N25/04 (2006.01) i, A01N25/22 (2006.01) i, A01N47/16 (2006.01) i, A01P3/00 (2006.01) i		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols)		
Int.Cl. A01N25/04, A01N25/22, A01N47/16, A01P3/00		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Published examined utility model applications of Japan 1922-1996 Published unexamined utility model applications of Japan 1971-2013 Registered utility model specifications of Japan 1996-2013 Published registered utility model applications of Japan 1994-2013		
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)		
CAPlus/REGISTRY (STN), JSTPlus/JMEDPlus/JST/580 (JDreamIII), Thomson Innovation		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X A	JP 2008-081430 A (SUMITOMO CHEMICAL COMPANY LIMITED) 2008.04.10, claim 1, 9, paragraph [0010], [0057], [0061]-[0064], examples 5-6 & CN 101977498 A & EP 2245929 A1 & KR 10-2010-0114506 A & US 2010/0331186 A1 & WO 2009/093742 A1	1-4 5
X A	JP 2008-081431 A (SUMITOMO CHEMICAL COMPANY LIMITED) 2008.04.10, claim 1, 9, paragraph [0010], [0056], [0058]-[0063], examples 3-4 & CN 101516187 A & KR 10-2009-0060307 A & US 2009/0239750 A1 & WO 2008/047570 A2	1-4 5
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.		
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family		
Date of the actual completion of the international search		Date of mailing of the international search report
26.07.2013		06.08.2013
Name and mailing address of the ISA/JP		Authorized officer
Japan Patent Office		Teruki AMANO
3-4-3, Kasumigaseki, Chiyoda-ku, Tokyo 100-8915, Japan		4H 5084
		Telephone No. +81-3-3581-1101 Ext. 3443

INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP2013/067585

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X A	JP 2009-173568 A (SUMITOMO CHEMICAL COMPANY LIMITED) 2009.08.06, claim 1, 9, paragraph [0010], [0072], [0075]-[0081], examples 3-4 & CN 101980596 A & EP 2245930 A1 & KR 10-2010-0114504 A & US 2011/0009271 A1 & WO 2009/093744 A1	1-4 5
X A	JP 2009-173578 A (SUMITOMO CHEMICAL COMPANY LIMITED) 2009.08.06, claim 1, 9, paragraph [0010], [0071], [0074]-[0079], examples 3-4 & CN 101980597 A & KR 10-2010-0114505 A & US 2011/0021359 A1 & WO 2009/093743 A1	1-4 5
X A	JP 2009-173579 A (SUMITOMO CHEMICAL COMPANY LIMITED) 2009.08.06, claim 1, 10, paragraph [0010], [0074], [0077], [0081], [0085], [0090]-[0091], examples 3-4, 17-18, 31-32 & CN 101977498 A & EP 2245929 A1 & KR 10-2010-0114506 A & US 2010/0331186 A1 & WO 2009/093742 A1	1-4 5
X A	JP 2009-173596 A (SUMITOMO CHEMICAL COMPANY LIMITED) 2009.08.06, claim 1, 9, paragraph [0010], [0117], [0120], [0124], [0129]-[0130], examples 3-4, 17-18 (no family)	1-4 5
A	JP 2011-522005 A (BASF SE) 2011.07.28, paragraph [0124], [0133]-[0134], example 1-4 & CN 102105060 A & KR 10-2011-0014712 A & US 2011/0177944 A1 & WO 2009/147205 A2	1-5
A	JP 59-039807 A (SUMITOMO CHEMICAL COMPANY LIMITED) 1984.03.05, whole document (no family)	1-5
A	US 2003/0134910 A1 (CIBA SPECIALTY CHEMICALS CORPORATION) 2003.07.17, whole document & CN 1438837 A & WO 2002/000023 A1	1-5

<Subject of search>

Claims 1-5 relate to a pesticidal composition, which comprises an agriculturally active ingredient (a) that is defined by a specific parameter "a retention rate is less than 50% after irradiation with xenon light (290 nm cutoff) at an intensity of 0.68 W/m² at 340 nm for 8 hours". The active ingredient (a) of claims 1-5 includes all of compounds having the property.

However, the compound disclosed in the meaning within PCT Article 5 is just one of the claimed active ingredients. Furthermore, it is impossible for a person skilled in the art to specify the scope of the compounds having the property, because the above mentioned parameter is neither commonly used nor accepted. Therefore, it cannot be considered that these claims are supported by the disclosure of the description in the meaning within PCT Article 6.

Such being the case, the search was made on a pesticidal composition comprising the compound that is described specifically in the description as agriculturally active ingredient (a), i.e., "fenpyrazamine" (paragraph[0009], page 6, line 13-24).