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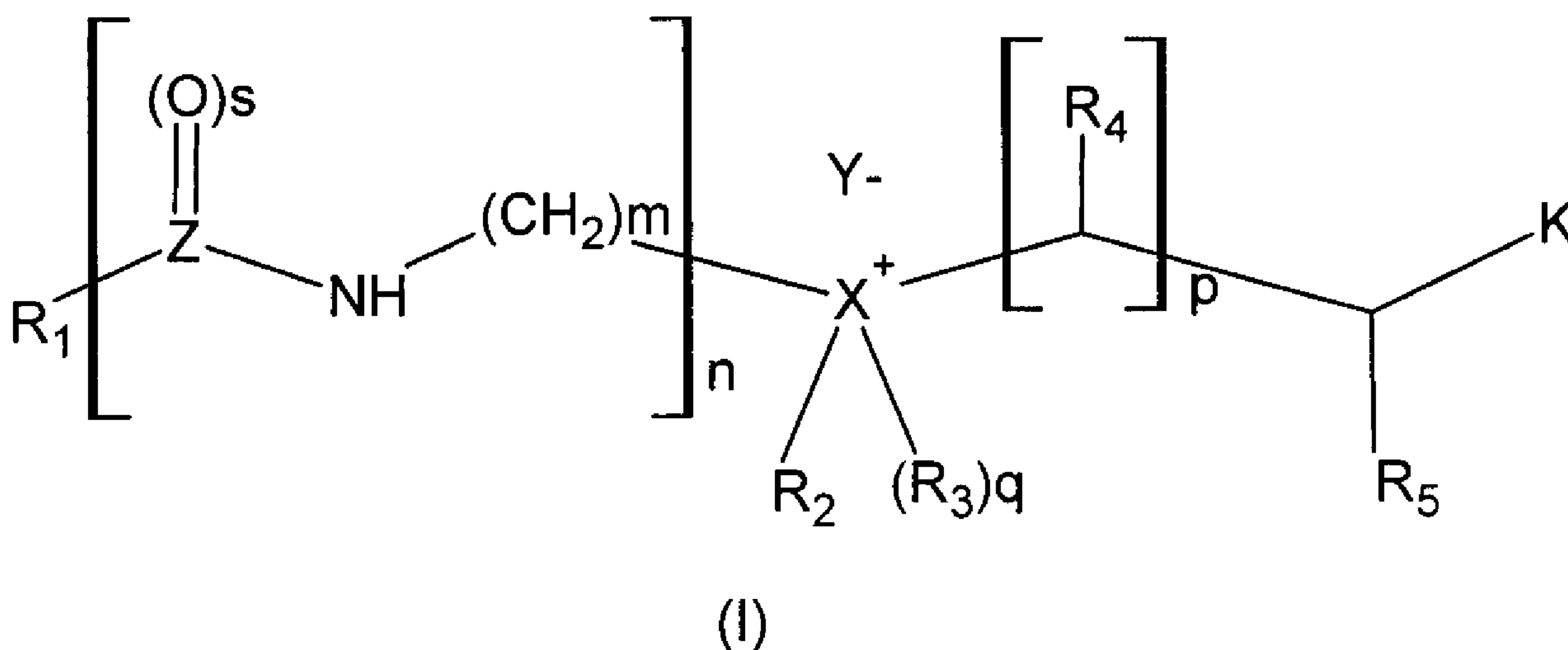
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(72) Inventeurs/Inventors:
FILIPPINI, LUCIO, IT;
GUSMEROLI, MARILENA, IT;
MORMILE, SILVIA, IT;
MIRENNA, LUIGI, IT

(73) Propriétaire/Owner:
BELCHIM CROP PROTECTION LUXEMBOURG
S.A.R.L., LU

(74) Agent: ROBIC

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(54) Title: ORGANIC DERIVATIVES, THEIR SALTS AND RELATIVE USE FOR THE CONTROL OF PHYTOPATHOGENS



(57) Abrégé/Abstract:

Organic compounds are described, which are capable of forming quaternary salts, quaternary salts thereof with a structure having general formula (I) and their use for the control of phytopathogen fungi.



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(71) Applicant (for all designated States except US): **ISAGRO S.P.A.** [IT/IT]; Via Caldera, 21, I-20153 Milan (IT).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **FILIPPINI, Lucio** [IT/IT]; Corso Vercelli, 83/S, I-28100 Novara (IT). **GUSMEROLI, Marilena** [IT/IT]; Via Don Valentini, 20, I-20052 Monza (IT). **MORMILE, Silvia** [IT/IT]; Via Vicenza, 9, I-28100 Novara (IT). **MIRENNA, Luigi** [IT/IT]; Via Gamboloita, 4, I-20139 Milan (IT).

(74) Agents: **DE GREGORI, Antonella** et al.; Barzano' & Zanardo Milano S.p.A., Via Borgonuovo, 10, I-20121 Milan (IT).

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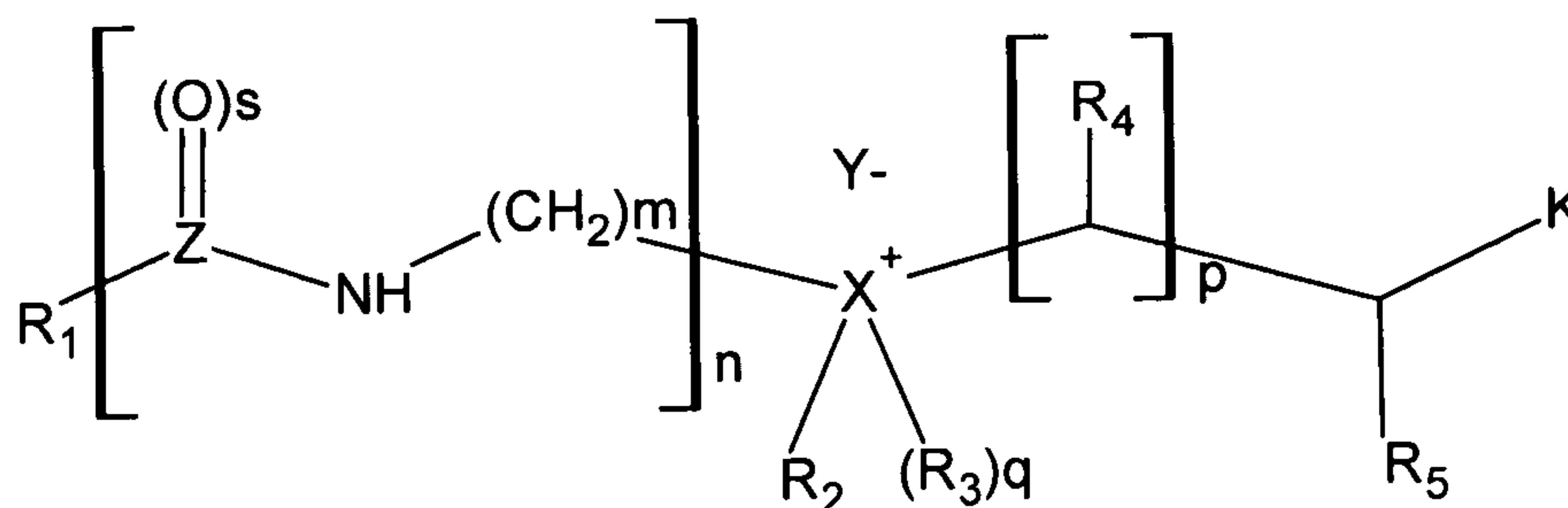
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(54) Title: ORAGANIC DERIVATIVES, THEIR SALTS AND USE FOR THE CONTROL OF PHYTO PATHOGENS



(I)

(57) Abstract: Organic compounds are described, which are capable of forming quaternary salts, quaternary salts thereof with a structure having general formula (I) and their use for the control of phytopathogen fungi.

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ORGANIC DERIVATIVES, THEIR SALTS AND RELATIVE USE FOR THE
CONTROL OF PHYTOPATHOGENS

The present invention relates to organic compounds
10 capable of forming quaternary salts, the relative quaternary salts and the relative use for the control of phytopathogens.

Quaternary salts are compounds known for being soluble in both hydrophilic and lyophilic environments. An
15 example of a quaternary salt is represented by choline, an ubiquitous substance in nature, known for being a biostimulating compound for plants as described in TG. Mason, G. Blunden (1989) Bot. Mar. 32 313-316.

Choline, moreover, in the interior of plants, is easily
20 ily oxidized to glycine betaine.

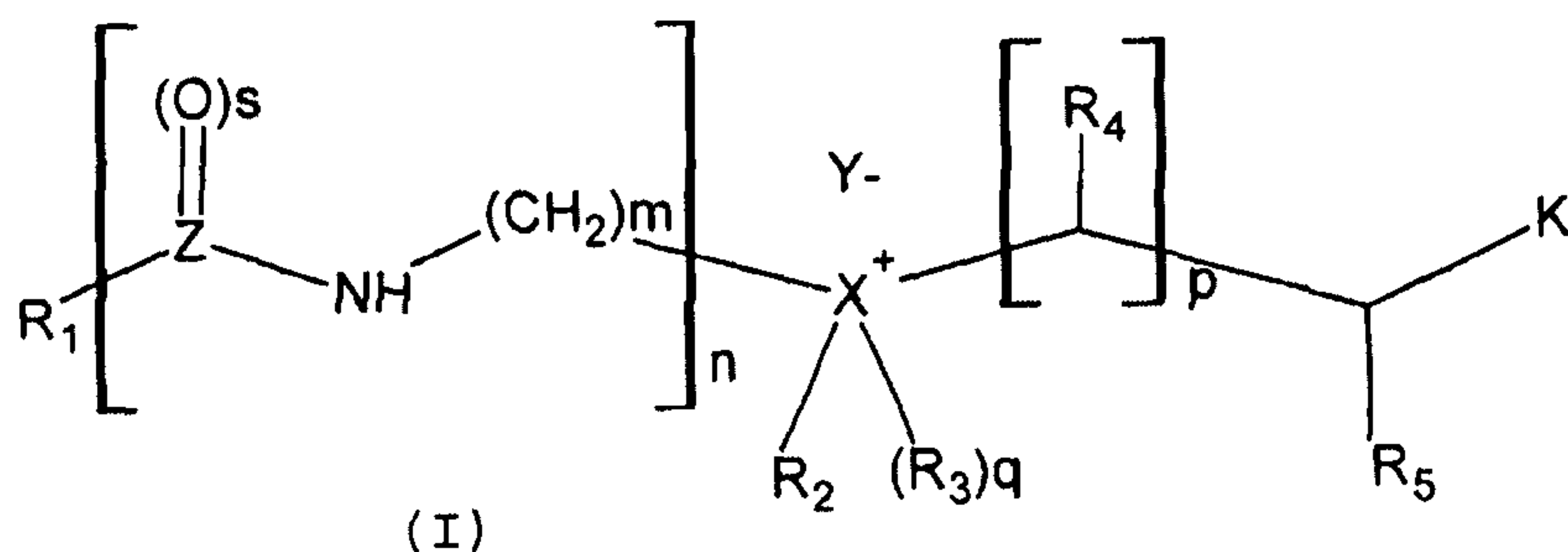
In the agronomical field, when administered, for example, to fruit plants, glycine betaine contributes to controlling abiotic and nutritional growth stress, reducing imperfections in fruit rinds and the tendency of the
25 rind to break during ripening, as described in EP-A-

0806897, by acting as an osmolyte regulator.

The Applicant has now surprisingly found that various organic compounds capable of forming quaternary salts and quaternary salts thereof have an unexpected activity as fungicidal and bactericidal products in the agronomical field, and obtain a prolonged protective action on vegetables with respect to phytopathogen fungi and bacteria.

These compounds are also capable of effectively synergizing with numerous other active principles known for having a fungicidal activity or, in turn, capable of inducing the natural defense of plants so as to be able to control both biotic and abiotic stress.

An object of the present invention as broadly disclosed therefore relates to organic compounds having general formula (I),



wherein:

- K represents a CH₂OH or COOR_a group;
- R_a represents a linear or branched C₁-C₂₆ alkyl group optionally substituted;

- R₁ represents a hydrogen or a linear or branched C₁-C₂₆ alkyl group optionally substituted; a linear or branched C₁-C₂₆ haloalkyl group optionally substituted; a linear or branched C₁-C₂₆ alkoxy group optionally substituted; a linear or branched C₁-C₂₆ alkylthio group optionally substituted; a linear or branched C₂-C₂₆ alkenyl group optionally substituted; a linear or branched C₂-C₂₆ alkynyl group optionally substituted; a C₃-C₃₀ cycloalkyl group optionally condensed or a condensed C₁₇ cycloalkyl group of the steroid type optionally substituted; a C₃-C₃₀ cycloalkoxy group optionally condensed and optionally substituted; a heterocyclic group optionally substituted; an aryl group optionally substituted; a heteroaryl group optionally substituted; a linear or cyclic C₆-C₁₂ group of the saccharide type optionally substituted; or, when n is different from 0, a C₁-C₂₆ alkylamine group or a C₂-C₂₆ dialkylamine group optionally substituted;
- R₂ and R₃, the same or different, represent a C₁-C₃ alkyl group optionally substituted;
- R₄ and R₅, the same or different, represent a hydrogen atom, or a linear or branched C₁-C₆ alkyl group optionally substituted; a linear or branched C₂-C₆ alkenyl group optionally substituted; a C₃-C₆ cyclo-

alkyl group optionally substituted; a hydroxyl group; an aryl group optionally substituted; a heteroaryl group optionally substituted; a heterocyclic group optionally substituted;

5 R_4 and R_5 can individually form a cycle together with R_2 ;

- X represents a nitrogen or sulfur atom;

- Z represents a carbon or sulfur atom;

- m represents a number ranging from 1 to 5;

10 - n and p represent a number ranging from 0 to 3;

- q has the value of 0 for X = sulfur or the value of 0 or 1 for X = nitrogen;

- Y, when q has the value of 0 for X = sulfur and when q has the value of 1 for X = nitrogen, represents a

15 halide, such as Cl^- , Br^- ; a nitrate anion (NO_3^-), a nitrite anion (NO_2^-); a phosphate anion selected from H_2PO_4^- , HPO_4^{-2} , PO_4^{-3} ; a phosphite anion selected from H_2PO_3^- , HPO_3^{-2} , PO_3^{-3} ; a carbonate anion (CO_3^{-2}); a bicarbonate anion (HCO_3^-); a sulfate anion (SO_4^{-2}); a

20 hydrogen-sulfated anion (HSO_4^-); or Y represents the salicylate anion; the acetylsalicylate anion; the saccharinate anion; the 3-aminobutanoate anion; the cyclamate anion; the taurinate anion; the ethylphosphate anion; or Y is absent when q has the value

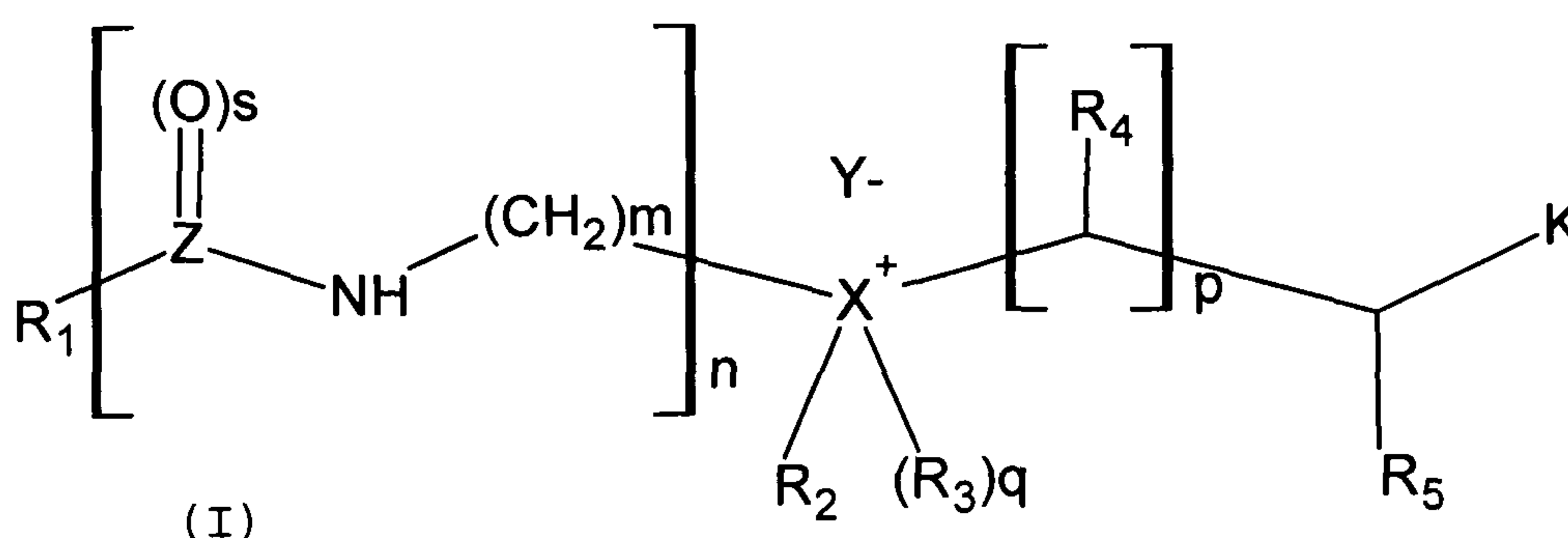
25 of 0 for X = nitrogen;

- s has the value of 1 for Z = carbon or the value of 2 for Z = sulfur.

The Applicant has also found that the compounds having general formula (I), not only have a direct fungicidal and bactericidal action, but are also capable of stimulating the natural defense systems of plants and inducing resistance in the plants themselves; this method for controlling diseases is becoming of growing interest as it is based on the amplification of a natural process already present in plants by the application of these compounds.

The Applicant has also surprisingly found that these compounds having general formula (I) are also an optimum means of controlling phytopathogens in genetically modified vegetable species to amplify the original natural defenses.

A further object of the present invention is therefore the use of compounds having general formula (I):



wherein:

- K represents a CH₂OH or COOR_a group;
- R_a represents a linear or branched C₁-C₂₆ alkyl group optionally substituted;
- 5 - R₁ represents a hydrogen or a linear or branched C₁-C₂₆ alkyl group optionally substituted; a linear or branched C₁-C₂₆ haloalkyl group optionally substituted; a linear or branched C₁-C₂₆ alkoxy group optionally substituted; a linear or branched C₁-C₂₆ alkylthio group optionally substituted; a linear or
10 cyclic C₂-C₂₆ alkenyl group optionally substituted; a linear or branched C₂-C₂₆ alkynyl group optionally substituted; a C₃-C₃₀ cycloalkyl group optionally condensed or a condensed C₁₇ cycloalkyl group of the
15 steroid type optionally substituted; a C₃-C₃₀ cycloalkoxy group optionally condensed and optionally substituted; a heterocyclic group optionally substituted; an aryl group optionally substituted; a heteroaryl group optionally substituted; a linear or
20 cyclic C₆-C₁₂ group of the saccharide type optionally substituted; a C₁-C₂₆ alkylamine group or a C₂-C₂₆ dialkylamine group optionally substituted for n different from 0;
- R₂ and R₃, the same or different, represent a C₁-C₃
25 alkyl group optionally substituted;

- R_4 and R_5 , the same or different, represent a hydrogen atom, or a linear or branched C_1 - C_6 alkyl group optionally substituted; a linear or branched C_2 - C_6 alkenyl group optionally substituted; a C_3 - C_6 cycloalkyl group optionally substituted; a hydroxyl group; an aryl group optionally substituted; a heteroaryl group optionally substituted; a heterocyclic group optionally substituted;
 R_4 and R_5 can individually form a cycle together with R_2 ;
- X represents a nitrogen or sulfur atom;
- Z represents a carbon or sulfur atom;
- m represents a number ranging from 1 to 5;
- n and p represent a number ranging from 0 to 3;
- q has the value of 0 for X = sulfur or the value of 0 or 1 for X = nitrogen;
- Y, when q has the value of 0 for X = sulfur and when q has the value of 1 for X = nitrogen, represents a halide, such as Cl^- , Br^- ; a nitrate anion (NO_3^-), a nitrite anion (NO_2^-); a phosphate anion selected from $H_2PO_4^-$, HPO_4^{-2} , PO_4^{-3} ; a phosphite anion selected from $H_2PO_3^-$, HPO_3^{-2} , PO_3^{-3} ; a carbonate anion (CO_3^{-2}); a bicarbonate anion (HCO_3^-); a sulfate anion (SO_4^{-2}); a hydrogen-sulfate anion (HSO_4^-); or Y represents the salicylate anion; the acetylsalicylate

anion; the saccharinate anion; the 3-aminobutanoate anion; the cyclamate anion; the taurinate anion; the ethylphosphonate anion; or Y is absent when q has the value of 0 for X = nitrogen;

- s has the value of 1 for Z = carbon or the value of 2 for Z = sulfur,

for the control of phytopathogen fungi and bacteria.

In particular an object of the present invention relates to the use of compounds with a structure having general formula (I) for the control of phytopathogen fungi and bacteria by stimulating the natural defense systems of plants and the induction of resistance in the plants themselves.

In particular, the use of the compounds having general formula (I) for the control of phytopathogen fungi is curative and/or preventive.

It is worth mentioning however that, in the invention as claimed hereinafter, the compounds of the general formula (I) are restricted to those where:

Y- is the phosphine anion H_2PO_3^- ;

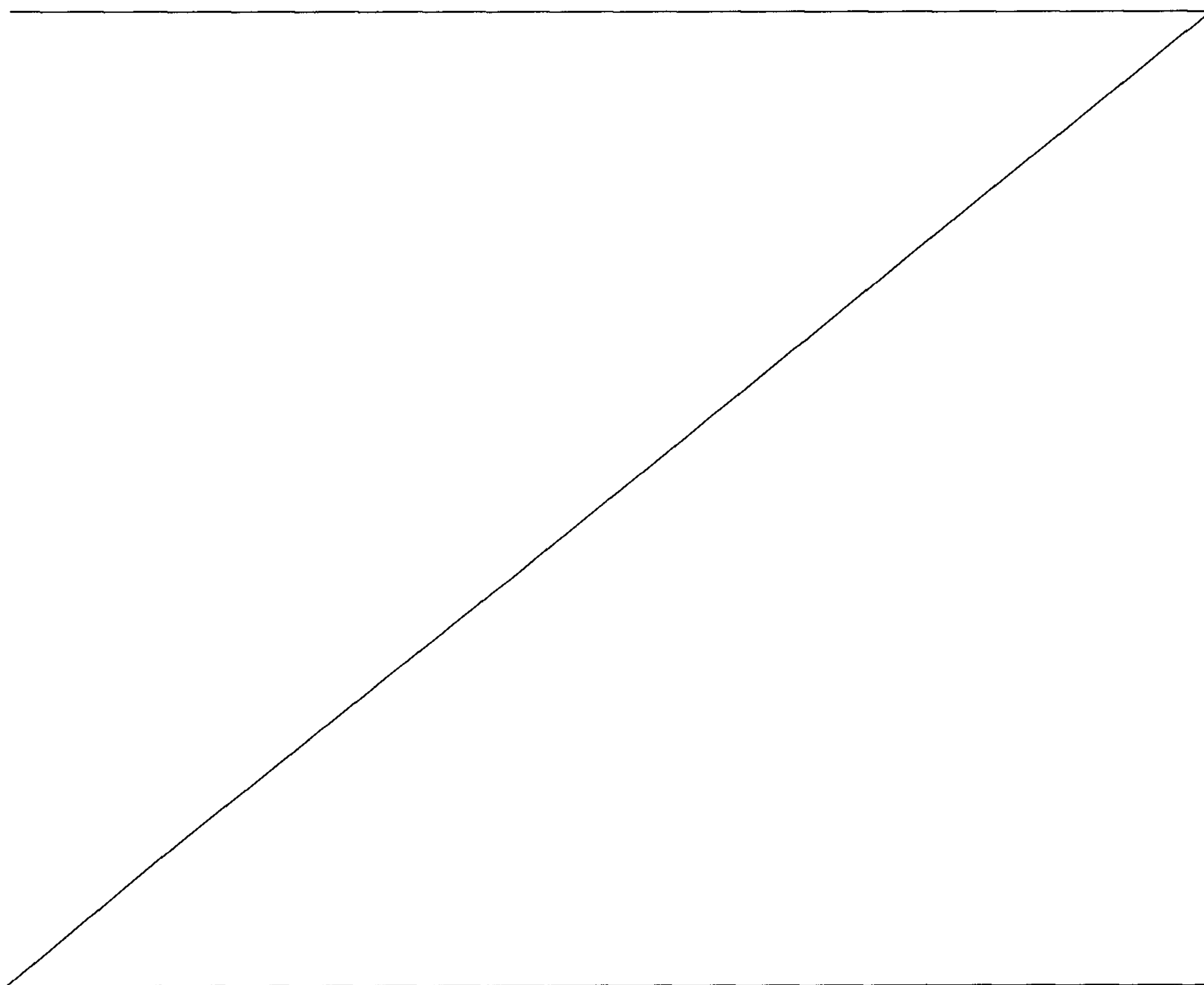
X is nitrogen; and

q is 1.

In the above definitions, a C_1 - C_{26} alkyl group refers to a linear or branched C_1 - C_{26} alkyl group, optionally substituted by one or more substituents, the same or different.

Examples of this group are: methyl, ethyl, propyl, isopropyl, butyl, isobutyl, tert-butyl, capryl, lauryl, stearyl, eicosyl, hexacosyl.

A C_1 - C_{26} haloalkyl group refers to a linear or branched alkyl group, substituted by one or more halogen atoms, the same or different.



Examples of this group are: fluoromethyl, difluoromethyl, trifluoromethyl, trichloromethyl, 2,2,2-trifluoroethyl, 2,2,2-trichloroethyl, 2,2,3,3-tetrafluoropropyl, 2,2,3,3,3-pentafluoropropyl, perfluoro-octanyl, perfluorododecyl.

A C₁-C₂₆ alkoxyl group refers to a C₁-C₂₆ alkoxyl group, wherein the aliphatic portion is a C₁-C₂₆ alkyl group, as previously defined.

Examples of this group are: methoxyl, ethoxyl, isopropoxyl, cyclopropylmethoxyl, lauryloxyl.

A C₁-C₂₆ thioalkyl group refers to a C₁-C₂₆ thioalkyl group, wherein the aliphatic portion is a C₁-C₂₆ alkyl group, as previously defined.

Examples of this group are: thiomethyl, thioethyl, thiolauryl, thiocapryl.

A C₂-C₂₆ alkenyl group refers to a linear or branched C₂-C₂₆ alkenyl group, optionally substituted by one or more substituents, the same or different.

Examples of this group are: ethenyl, propenyl, butenyl, 1-decenyl, 8-heptadecenyl, 8,11,14-heptadecatrienyl, 8,11-heptadecadienyl.

A C₂-C₂₆ alkynyl group refers to a linear or branched C₂-C₂₆ alkynyl group, optionally substituted by one or more substituents, the same or different.

Examples of this group are: ethynyl, propargyl, 1-

dodecynyl, 1-octadecynyl.

A C₃-C₃₀ cycloalkyl group optionally condensed refers to a cycloalkyl group whose ring consists of 3-30 carbon atoms, optionally substituted by one or more substituents, the same or different.

Examples of this group are: cyclopropyl, 2,2-dichlorocyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, decaline, abietyl.

A condensed C₁₇ cycloalkyl group of the steroid type refers to a steroid group consisting of 17 carbon atoms, optionally substituted by one or more substituents, the same or different.

Examples of this group are: cholanyl, or chenodeoxycholanyl, or ursodeoxycholanyl, or deoxycholanyl, or iodeoxycholanyl, or lithocholanyl.

A C₃-C₃₀ cycloalkoxyl group refers to a C₃-C₃₀ cycloalkoxyl group wherein the aliphatic portion is a C₃-C₃₀ cycloalkyl group as previously defined.

Examples of this group are: cyclopentoxy, cyclohexyloxy, cholesteryl.

A C₁-C₂₆ alkylamine or a C₂-C₂₆ dialkylamine group refers to an alkylamine or dialkylamine group wherein the aliphatic portion is respectively a C₁-C₂₆ or two C₁-C₁₃ alkyl groups as previously defined.

Examples of this group are: methylamine, dimethyla-

mine, ethylamine, isopropylamine, dibutylamine, dioctylamine, hexadecylamine, didecylamine.

An aryl group refers to an carbocyclic aromatic group optionally substituted by one or more groups, the same or different.

Examples of this group are: phenyl, naphthyl, phenanthryl.

A heteroaryl group refers to a penta- or hexa-atomic heterocyclic aromatic group also benzocondensed or heterobicyclic, containing from 1 to 4 heteroatoms selected from nitrogen, oxygen, sulfur, optionally substituted by one or more groups, the same or different.

Examples of hetero-aryl groups are: pyridine, pyrimidine, pyridazine, pyrazine, triazine, tetrazine, quinoline, quinoxaline, quinazoline, furan, thiophene, pyrol, oxazole, thiazole, isoxazole, isothiazole, oxadiazole, thiadiazole, pyrazole, imidazole, triazole, tetrazole, indole, benzofuran, benzothiophene, benzoxazole, benzothiazole, benzoxadiazole, benzothiadiazole, benzopyrazole, benzimidazole, benzotriazole, triazolepyridine, triazolepyrimidine, thiazoletriazole, cumarin.

A heterocyclic group refers to a saturated or unsaturated ring with three to twelve terms, containing at least a heteroatom selected from nitrogen, oxygen, sulfur, optionally condensed with another aromatic or non-

aromatic ring.

Examples of heterocyclic rings are: pyrrolidine, piperidine, dihydropyridine, piperazine, 2,6-diketopiperazine, 2-ketoazetidine, morpholine, thiazine,
5 indoline.

A linear or cyclic C₆-C₁₂ group of the saccharide type refers to a carbohydrate group in open or closed form.

Examples of this group are: gluconyl, glucopyranosyl,
10 syl, β -D-fructofuranosyl- α -D-glucopyranosyl, 4-O- β -D-galactopyranosyl-D-glucosyl.

Optionally substituted means, in all parts of the patent application, one or more substituents, the same or different, selected from the following groups: halogen
15 atoms; C₁-C₆ alkyls, C₁-C₆ alkoxy and C₁-C₆ alkylthio, in turn optionally substituted by halogen atoms; C₁-C₆ alkylcarbonyls and C₁-C₆ alkoxy carbonyls, optionally halogenated; aminocarbonyls, C₁-C₆ alkylaminocarbonyls, C₂-C₁₂ dialkylaminocarbonyls, optionally halogenated; carboxyl;
20 C₁-C₆ alkylcarbonyloxy optionally halogenated; cyano; nitro; formyl; hydroxyl; amino; aryl and heteroaryl optionally substituted.

Examples of compounds having general formula (I), when q has the value of 0 for X = nitrogen, which are interesting for their activity are:
25

- N,N-dimethylethanolamine;
- 3-dimethylamine-1-propanol;
- N-ethyl,N-methylethanolamine;
- 2-dimethylaminopropanol;
- 5 • N-lauryl,N-methylethanolamine;
- methyl ester of N,N-dimethyl- β -alanine;
- methyl ester of N,N-dimethylglycine.

Examples of compounds having general formula (I) when
q has the value of 0 for X = sulfur and the value of 1
10 for X = nitrogen, which are interesting for their activ-
ity are:

- Acid choline phosphite;
- Neutral choline phosphite;
- Choline ethylphosphonate;
- 15 • Acid laurylcholine phosphite;
- Neutral laurylcholine phosphite;
- Acid cocamidopropylcholine phosphite;
- Neutral cocamidopropylcholine phosphite;
- Acid stearylcholine phosphite;
- 20 • Neutral stearylcholine phosphite;
- Acid cholesterylcarbonylamidopropylcholine phos-
phite;
- Neutral cholesterylcarbonylamidopropylcholine
phosphite;
- 25 • Acid cholanylamidopropylcholine phosphite;

- Neutral cholanylamidopropylcholine phosphite;
- Acid chenodeoxycholanylamidopropylcholine phosphite;
- Neutral chenodeoxycholanylamidopropylcholine phosphite;
- Acid N,N-dimethyl,N-laurylamidopropyl [L] valinol phosphite;
- Neutral N,N-dimethyl,N-laurylamidopropyl [L] valinol phosphite;
- Acid N,N-dimethyl,N-lauryl [L] valinol phosphite;
- Neutral N,N-dimethyl,N-lauryl [L] valinol phosphite;
- Acid N-lauryl,N-methyl [L] 2-pyrrolidinemethanol phosphite;
- Neutral N-lauryl,N-methyl [L] 2-pyrrolidinemethanol phosphite;
- Choline salicylate;
- Choline acetylsalicylate;
- Choline saccharinate;
- Choline cyclamate;
- Choline taurinate;
- Laurylcholine salicylate;
- Laurylcholine acetylsalicylate;
- Laurylcholine saccharinate;
- Laurylcholine cyclamate;

- Laurylcholine taurinate;
- Laurylcholine ethylphosphonate;
- Cocamidopropylcholine salicylate;
- Cocamidopropylcholine acetylsalicylate;
- 5 • Cocamidopropylcholine saccharinate;
- Cocamidopropylcholine cyclamate;
- Cocamidopropylcholine taurinate;
- Cocamidopropylcholine ethylphosphonate;
- Stearylcholine salicylate;
- 10 • Stearylcholine acetylsalicylate;
- Stearylcholine saccharinate;
- Stearylcholine cyclamate;
- Stearylcholine taurinate;
- Cholanylamidopropylcholine salicylate;
- 15 • Cholanylamidopropylcholine acetylsalicylate;
- Cholanylamidopropylcholine saccharinate;
- Cholanylamidopropylcholine cyclamate;
- Cholanylamidopropylcholine taurinate;
- Choline carbonate;
- 20 • Choline bicarbonate;
- Laurylcholine carbonate;
- Laurylcholine bicarbonate;
- Cocamidopropylcholine carbonate;
- Cocamidopropylcholine bicarbonate;
- 25 • Stearylcholine carbonate;

- Stearylcholine bicarbonate;
- Choline sulfate;
- Choline chloride;
- Laurylcholine sulfate;
- 5 • Laurylcholine hydrogen-sulfate;
- Cocamidopropylcholine sulfate;
- Cocamidopropylcholine bromide;
- Stearylcholine sulfate;
- Stearylcholine chloride;
- 10 • Acid choline phosphate;
- Neutral choline phosphate;
- Acid laurylcholine phosphate;
- Neutral laurylcholine phosphate;
- Acid cocamidopropylcholine phosphate;
- 15 • Neutral cocamidopropylcholine phosphate;
- Acid Stearylcholine phosphate;
- Neutral Stearylcholine phosphate;
- Acid phosphite of the methyl ester of cocamido-
propylbetaine;
- 20 • Neutral phosphite of the methyl ester of cocami-
dopropylbetaine;
- Acid phosphite of betaine cetyl ester;
- Neutral phosphite of betaine cetyl ester;
- Salicylate of the methyl ester of cocamidopropyl-
25 betaine;

- Acetylsalicylate of the methyl ester of cocamidopropylbetaine;
- Salicylate of betaine cetyl ester;
- Acetylsalicylate of betaine cetyl ester;
- 5 • Cyclamate of the methyl ester of cocamidopropylbetaine;
- Saccharinate of the methyl ester of cocamidopropylbetaine;
- Cyclamate of betaine cetyl ester;
- 10 • Saccharinate of betaine cetyl ester;
- Bicarbonate of the methyl ester of cocamidopropylbetaine;
- Chloride of the methyl ester of cocamidopropylbetaine;
- 15 • Acid phosphite of the methyl ester of cholanilamidopropylbetaine;
- Neutral phosphite of the methyl ester of cholanilamidopropylbetaine;
- Acid phosphite of the methyl ester of carnitine;
- 20 • Neutral phosphite of the methyl ester of carnitine;
- Salicylate of the methyl ester of carnitine;
- Acetylsalicylate of the methyl ester of carnitine;
- 25 • Saccharinate of the methyl ester of carnitine;

- Carnitine chloride.

Particularly preferred are quaternary salts having general formula (I) when Y represents the salicylate anion; the acetylsalicylate anion; the saccharinate anion; 5 the 3-aminobutanoate anion; the cyclamate anion; the taurinate anion; the ethylphosphonate anion; or when Y represents a phosphite anion selected from H_2PO_3^- , HPO_3^{-2} , PO_3^{-3} , a halide anion or a bicarbonate anion (HCO_3^-).

The Applicant has now surprisingly found that quaternary salts having formula (I), when Y represents a phosphite anion selected from H_2PO_3^- , HPO_3^{-2} , PO_3^{-3} , or the salicylate anion; the acetylsalicylate anion; the saccharinate anion; the 3-aminobutanoate anion; the cyclamate anion; the taurinate anion; the ethylphosphonate anion, have a surprisingly higher activity than that expected deriving from the anion, which is known in literature for having its own biological efficacy, and the relative quaternary cation, demonstrating the synergism between the ionic couple of compounds having general formula (I) when q has the value of 0 for X = sulfur and the value of 1 for X = nitrogen.

An unexpected synergic effect has therefore been found between the cationic component and anionic component of quaternary salts having formula (I) when Y represents a phosphite anion selected from H_2PO_3^- , HPO_3^{-2} , PO_3^{-3} , 25

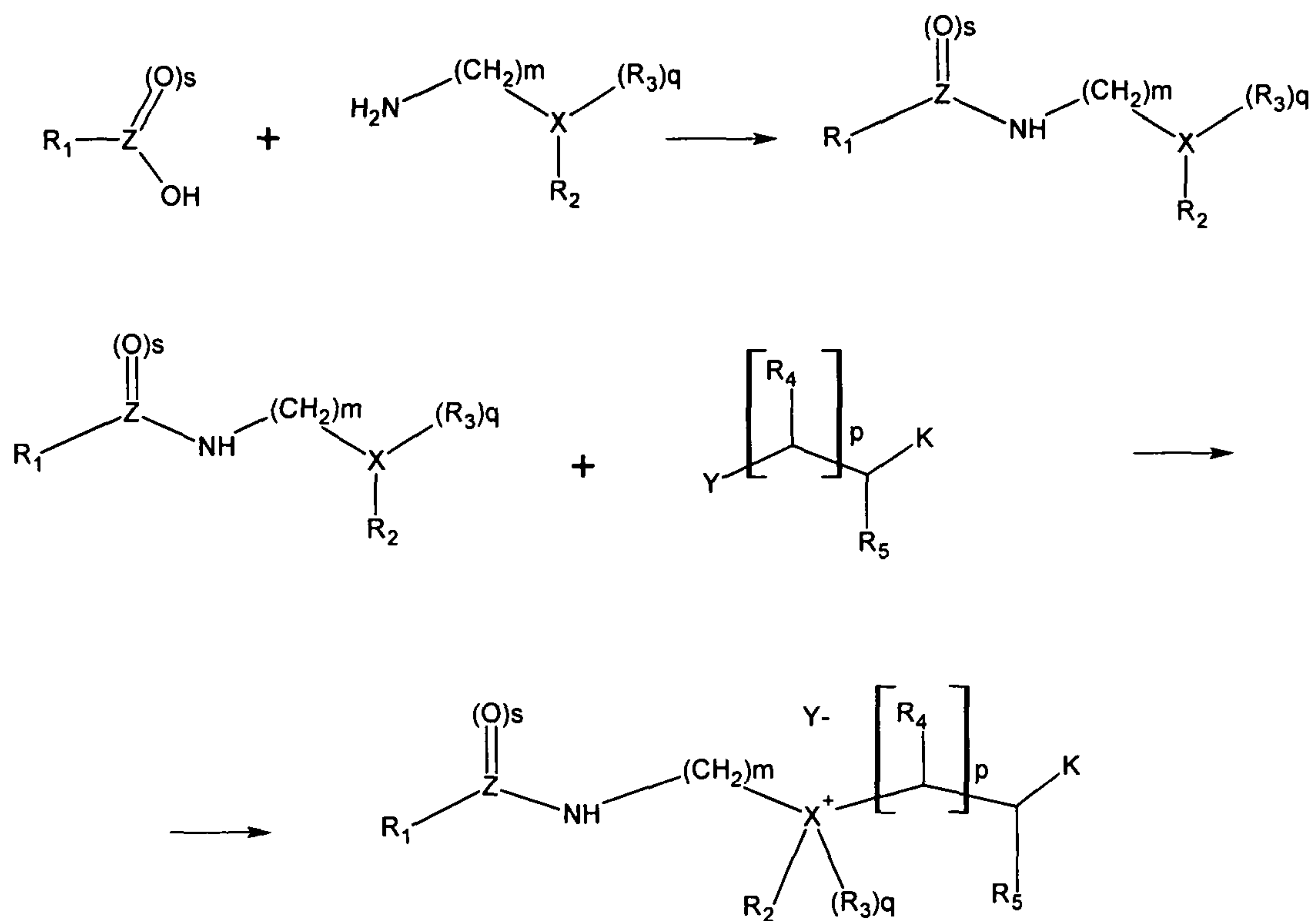
or the salicylate anion; the acetylsalicylate anion; the saccharinate anion; the 3-aminobutanoate anion; the cyclamate anion; the taurinate anion and q has the value of 0 for X = sulfur and the value of 1 for X = nitrogen.

5 When they are not commercial products, the compounds having general formula (I), when q has the value of 0 for X = nitrogen, can be easily obtained according to the methods described for example in Comprehensive Organic Transformations 1989, R. C. Larock, or in March's Advanced Organic Chemistry 2001 V^a edition, M. B. Smith, J. March.

10 The compounds having formula (I), when q has the value of 0 for X = sulfur and the value of 1 for X = nitrogen, can be obtained with different synthetic methods according to the meaning of the anion Y.

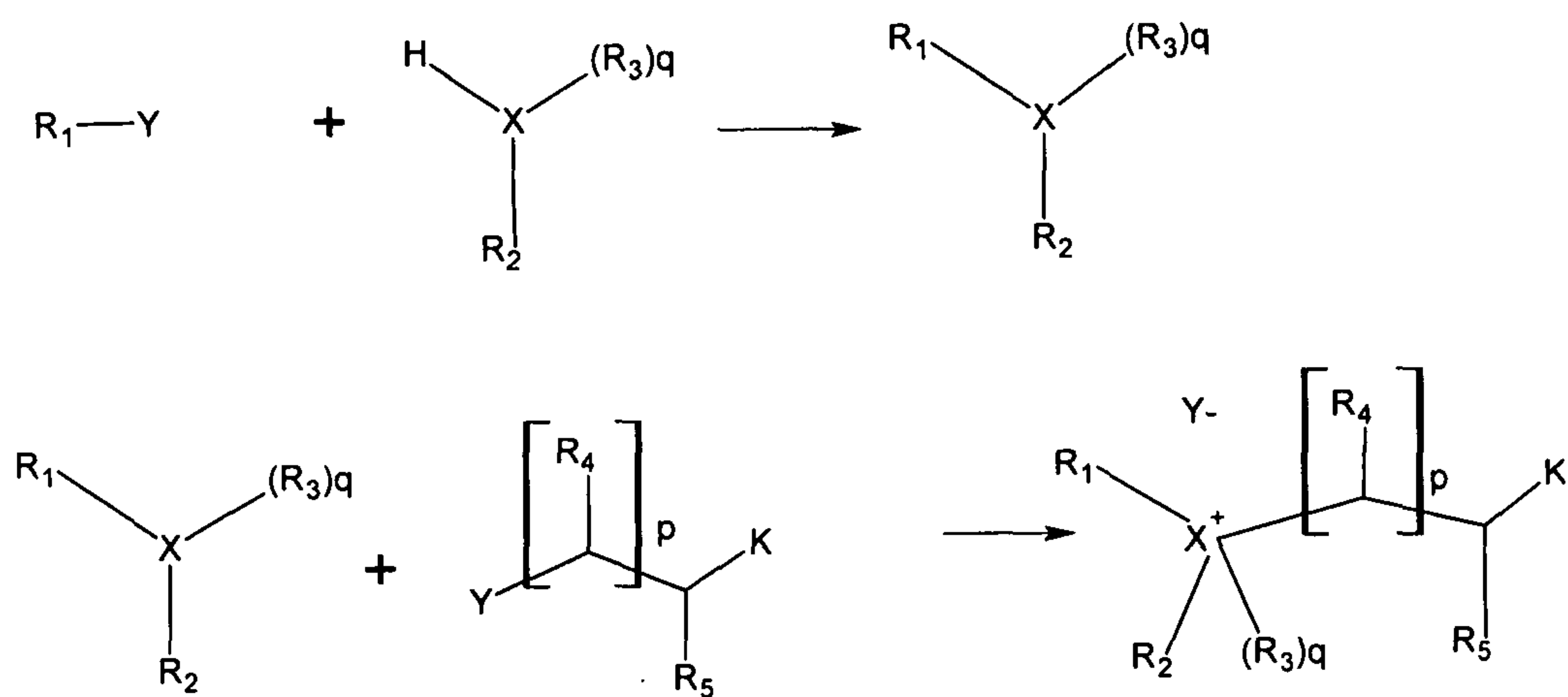
15 When Y has the meaning of a Cl⁻ e Br⁻ anion, and when R₁ has the meanings defined above with the exclusion of a C₁-C₂₆ alkoxyl group, or a C₁-C₂₆ alkylthio group, or a C₃-C₃₀ cycloalkoxyl group, or a C₁-C₂₆ alkylamine group, or a C₂-C₂₆ dialkylamine group, the quaternary salts having formula(I) can be easily obtained according to the reaction scheme A for n different from 0 and according to the reaction scheme B for n = 0:

Scheme A



Scheme B

5



wherein K, R₁, R₂, R₃, R₄, R₅, X, Z, m, p, q and s have the meanings defined above and Y represents an outgoing group such as a chlorine atom or a bromine atom which also become the counterion of the final product.

5 The quaternary salts having general formula (I), according to reaction scheme A, for X = nitrogen, can be obtained by condensation of the suitable N',N'-dialkylamine-N-alkylamine or, for X = sulfur, by condensation of the suitable ω-alkylthioalkylamine with car-
10 boxylic acid of a suitable R₁ residue, and a condensing agent, optionally in the presence of a base in an organic or aqueous solvent, according to methods well known in the art, for example in Comprehensive Organic Transformations 1989, R.C. Larock, so as to form the corresponding
15 amide.

The intermediate thus obtained is subsequently subjected to alkylation by reaction with the suitable halide, in water or in an organic solvent, at temperatures ranging from room temperature to 100°C, maintaining the
20 pH at values of around 7.5, by the controlled addition of a solution of a strong base.

The quaternary salts having general formula (I), according to reaction scheme B, for X = nitrogen, can be obtained by alkylation of the suitable N',N'-
25 dialkylamine-N-alkylamine or, for X = sulfur, by alkyla-

tion of the suitable ω -alkylthioalkylamine with the desired R_1 residue having the outgoing group Y, in the presence of a base in an organic or aqueous solvent, according to methods well known in the art, for example in
5 Comprehensive Organic Transformations 1989, R.C. Larock, so as to form the corresponding tertiary amine.

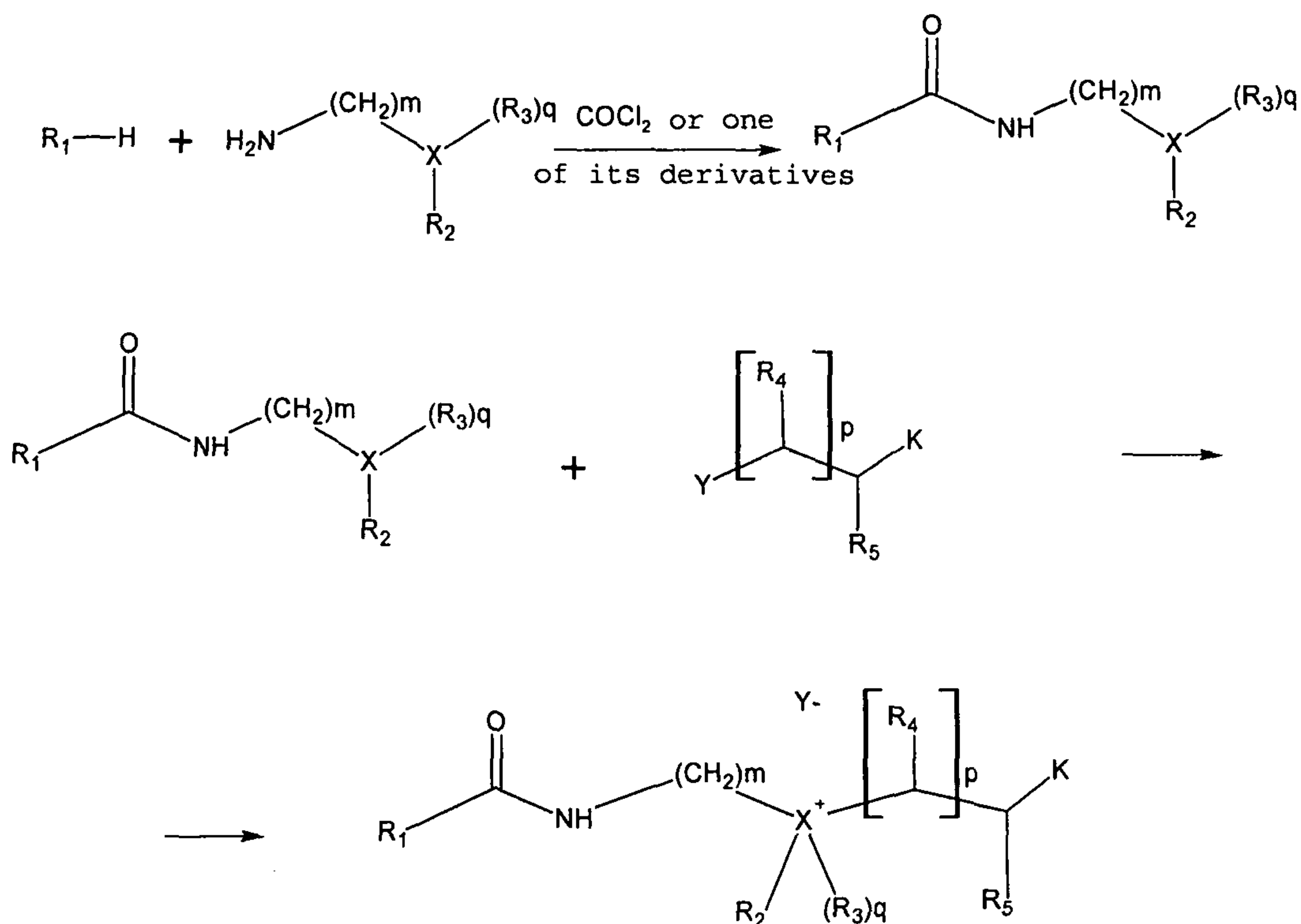
The intermediate thus obtained is subsequently subjected again to alkylation by reaction with the suitable halide, in water or in an organic solvent, at temperatures ranging from room temperature to 100°C, maintaining
10 the pH at values of around 7.5, by the controlled addition of a solution of a strong base.

The quaternary salts having formula (I), when Y has the meaning of a Cl^- and Br^- anion and when R_1 has the
15 meanings of a C_1 - C_{26} alkoxyl group, or a C_1 - C_{26} alkylthio group, or a C_3 - C_{30} cycloalkoxyl group, or a C_1 - C_{26} alkylamine group, or a C_2 - C_{26} dialkylamine group, can be easily obtained according to the reaction scheme C for n different from 0:

20

Scheme C

25



wherein K, R₁, R₂, R₃, R₄, R₅, X, Z, m, p, q and s have the meanings defined above and Y represents an outgoing group such as a chlorine atom or a bromine atom, which
 5 becomes the counterion of the final product.

The quaternary salts having general formula (I), according to reaction scheme C, for X = nitrogen, can be obtained by reaction of the suitable N',N'-dialkylamine-N-alkylamine or, for X = sulfur, by reaction of the suitable
 10 ω-alkylthioalkylamine with the desired R₁ residue having an alcoholic, or thioalcoholic, or aminic function when R₁ has the meanings of a C₁-C₂₆ alkoxyl group, or a C₃-C₃₀ cycloalkoxyl group, or a C₁-C₂₆ alkylthio group, or a C₁-C₂₆ alkylamine group, or a C₂-C₂₆ dialkylamine group
 15 respectively, in the presence of phosgene or one of its

functional substitutes, such as, for example, diphosgene, triphosgene, 1,1'-carbonyldiimidazole, in an organic or aqueous solvent, according to methods well known in the art, for example in Comprehensive Organic Transformations
5 1989, R.C. Larock, so as to form the corresponding carbamate, thiocarbamate or urea.

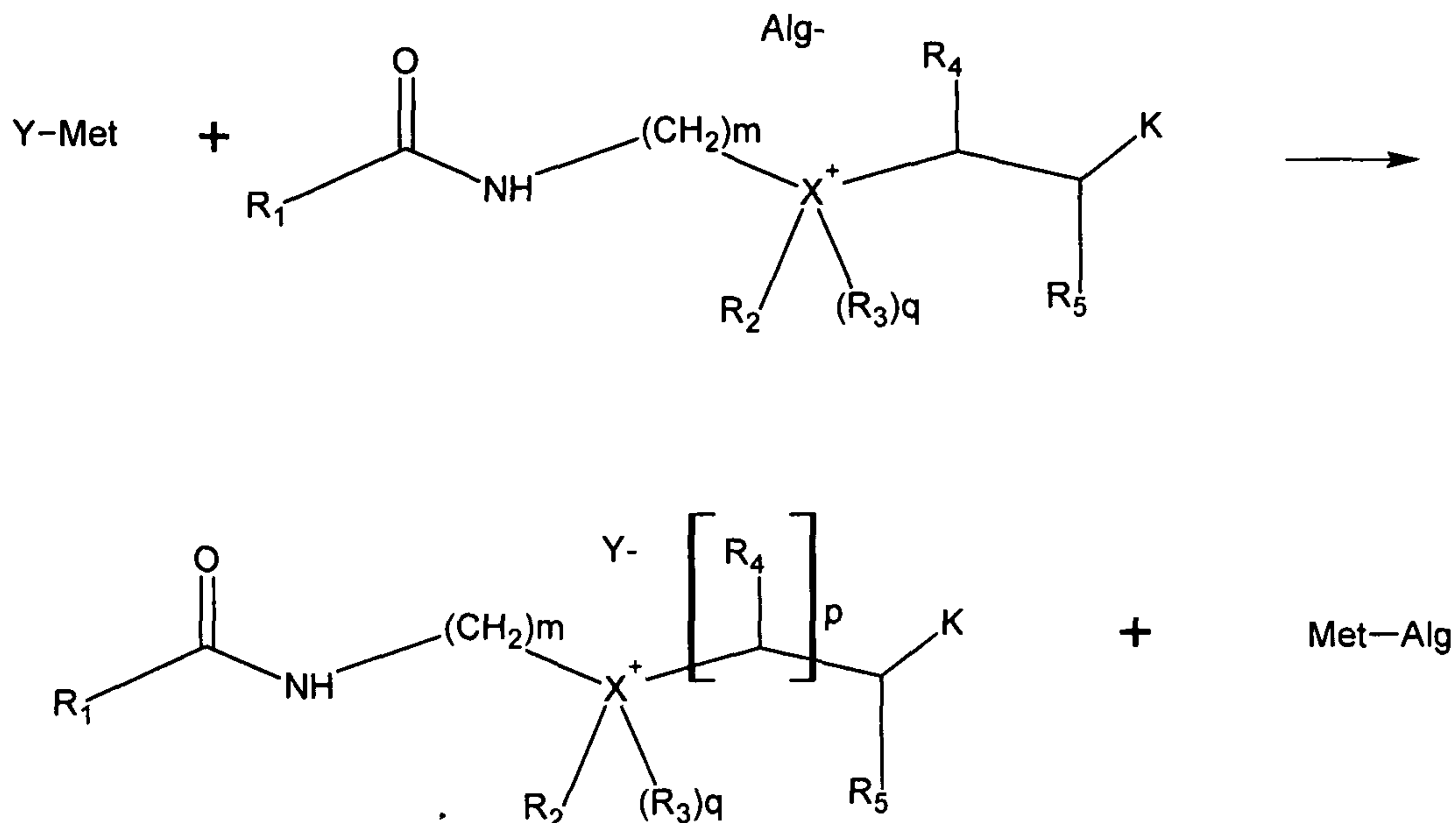
The intermediate thus obtained is subsequently subjected again to alkylation by reaction with the suitable halide, in water or in an organic solvent, at temperatures ranging from room temperature to 100°C, maintaining
10 the pH at values of around 7.5, by the controlled addition of a solution of a strong base.

When Y has a meaning different from a Cl⁻ and Br⁻ anion, the quaternary salts having formula (I) can be easily obtained by the exchange of the alkaline salts, such
15 as for example sodium and potassium, of the suitable acids YH, with the halides of the quaternary cations synthesized as described above, according to the reaction scheme D:

20

25

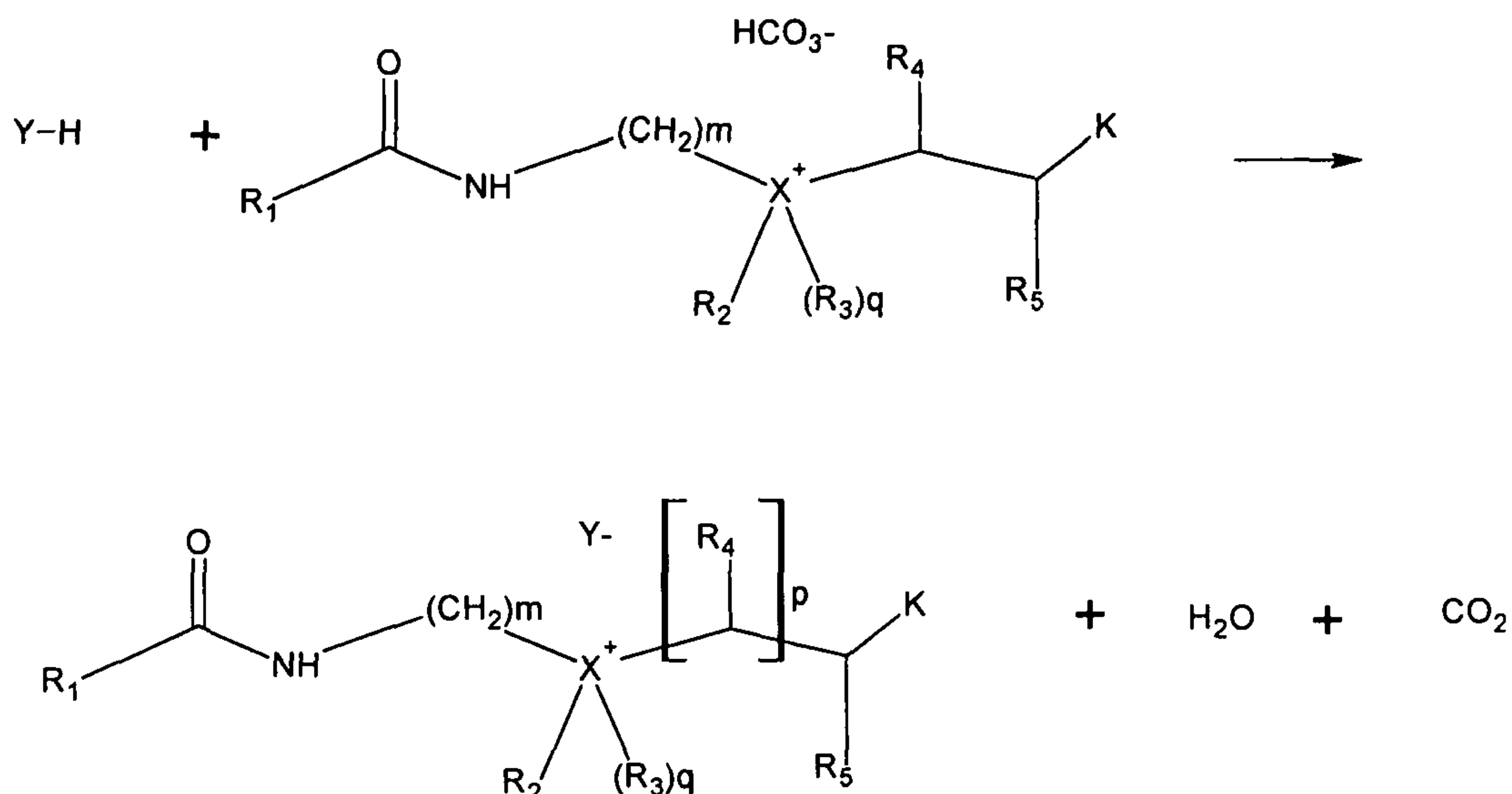
Scheme D



wherein K, R₁, R₂, R₃, R₄, R₅, X, Z, m, p, q and s have
 5 the meanings defined above and Y represents the acid
 residue which acts as counterion of the final product.

Alternatively, the quaternary salts having formula
 (I), for Y different from HCO₃⁻, can be easily obtained
 by the appropriate molar salification of the acid YH with
 10 the bicarbonates of the relative quaternary cations, ac-
 cording to the reaction scheme E:

Scheme E



wherein K, R₁, R₂, R₃, R₄, R₅, X, Z, m, p, q and s have the meanings defined above and Y represents the acid residue which becomes the counterion of the final product.

The reactions can be conveniently carried out in an aqueous or inert organic solvent, at a temperature ranging from room temperature to the boiling point of the reaction mixture, optionally in the presence of an inorganic or organic base.

Examples of preferred solvents for effecting the reaction are ethers (ethyl ether, isopropyl ether, tetrahydrofuran, dioxane, dimethoxyethane, etc.); esters (ethyl acetate, etc.); chlorinated hydrocarbons (methylene chloride, dichloroethane, chloroform, carbon tetrachloride, etc.); aromatic hydrocarbons (benzene, toluene, xylene,

etc.); aliphatic hydrocarbons (hexane, heptane, cyclohexane, etc.); aprotic dipolar solvents (N,N-dimethylformamide, dimethylsulfoxide, sulfolane, etc.).

Examples of preferred inorganic bases are: hydroxides, carbonates of alkaline or alkaline earth metals (sodium, potassium, calcium, etc.).

Examples of preferred organic bases are: pyridine, dimethylaminopyridine, aliphatic amines (triethylamine, etc.), cyclic amines (morpholine, piperidine, etc.).

If the substituents R_1 , R_2 , R_3 , R_4 , R_5 contain optic or geometric isomerism centres, the compounds having general formula (I) can be present in all possible configurational isomeric forms.

The scope of the present invention therefore also comprises the use of the compounds having general formula (I) as isomeric mixtures in any proportion, and also the formation and use of the single isomers for the control of phytopathogen fungi in the agronomical field.

When deriving from natural extracts, the compounds having general formula (I) can also be present in mixtures of their homologous products and the scope of the present invention consequently also includes the use of the compounds having general formula (I) as mixtures of their homologous products in any proportion, for the control of phytopathogen fungi and bacteria in the agronomi-

cal field.

The compounds having general formula (I) can also be present in a hydrated form by the coordination of any number of water molecules, or obtained in aqueous solution and used directly for agronomical purposes.

The quaternary salts having general formula (I) can also contain and possibly coordinate within their structure other metallic cations, such as for example sodium, potassium, whose number can vary in relation to the preparation method used for the synthesis of the compound having general formula (I).

The scope of the present invention therefore also comprises the use of said solutions of quaternary salts having formula (I), containing said salts for the control of phytopathogen fungi and bacteria in the agronomical field.

The scope of the present invention also comprises the use of mixtures of compounds having general formula (I) in any proportion.

The Applicant has also found, in agronomical practice, that the fungicidal action of the compounds having general formula (I) is particularly effective when combined with that of numerous other fungicidal active principles, thus forming an excellent instrument for antire-sistance strategies, allowing the application dosages to

be further lowered and stimulating the natural defenses of plants.

The Applicant has again surprisingly found that the compounds having general formula (I), when q has the value of 0 for X = nitrogen, are of particular interest as they are capable of exerting a considerable synergic effect if used in a mixture with Fosetyl-Aluminium (fungicidal active principle corresponding to aluminium triethylphosphonate, described in "The Pesticide Manual", 1994, X^a edition, British Crop Protection Council Ed., page. 530 and more commonly known with the trade-name of Aliette) thus allowing, when applied together, a higher fungicidal activity to be obtained with respect to that envisaged on the basis of the activity of the single components, and also providing excellent antiresistance properties.

An object of the present invention therefore also relates to fungicidal compositions comprising one or more compounds having general formula (I), when q has the value of 0 for X = nitrogen, mixed with Fosetyl-Aluminium (Fosetyl-Al).

Preferred fungicidal compositions are:

- a). N,N-dimethylethanolamine and Fosetyl-Al;
- b). 3-dimethylamine-1-propanol and Fosetyl-Al;
- 25 c). N-ethyl,N-methylethanolamine and Fosetyl-Al;

- d) . 2-dimethylaminopropanol and Fosetyl-Al;
e) . N-lauryl,N-methylethanolamine and Fosetyl-Al;
f) . methyl ester of N,N-dimethyl- β -alanine and Fosetyl-Al;
5 g) . methyl ester of N,N-dimethylglycine and Fosetyl-Al.
Composition a) is particularly preferred for its fungicidal activity.

The additional compounds having a fungicidal activity which can be used together with the compounds having
10 general formula (I), when q has the value of 0 for X = sulfur and the value of 1 for X = nitrogen, according to the present invention, are preferably selected from:

- (1) IR5885, a dipeptide compound corresponding to diastereoisomeric mixtures of methyl [S-(R,S)]-[3-(N-isopropoxycarbonylvalinyl)-amino]-3-(4-chloro-
15 phenyl)propanoate in any proportion, or to one of the two diastereoisomeric forms S-R or S-S taken individually;
(2) IR6141, corresponding to methyl N-(phenylacetyl)-N-
20 2,6-xylyl-R- alaninate;
(3) Tetraconazole (in its racemic form or as an optically active R isomer)
(4) Salicylic acid (SA) or its derivatives such as acetylsalicylic acid (ASA), copper salts of salicylic
25 acid (SA₂Cu) or (SACu) or acetylsalicylic acid (ASA₂Cu);

- (5) A copper (I) or copper (II) salt, such as copper oxychloride, copper hydroxide, Bordeaux mixture, copper sulfate, or a mixture of copper hydroxide and oxychloride (Airone);
- (6) Benalaxyl corresponding to methyl N-(phenylacetyl)-N-2,6-xylyl-RS-alaninate;
- (7) Metalaxyl corresponding to methyl N-(2-methoxyacetyl)-N-2,6-xylyl-RS-alaninate;
- 10 (8) Metalaxyl-M corresponding to methyl N-(2-methoxyacetyl)-N-2,6-xylyl-R-alaninate;
- (9) Oxadixyl corresponding to 2-methoxy-N-(2-oxo-1,3-oxazolidin-3-yl)aceto-2',6'-xylidide;
- (10) Mandipropamid corresponding to 2-(4-chlorophenyl)-N-[2-(3-methoxy-4-prop-2-inyloxy-phenyl)ethyl]-2-prop-2-inyloxy-acetamide;
- 15 (11) Iprovalicarb corresponding to O-(1-methyl-ethyl)-N-[2-methyl-1-[[[1-(4-methyl-phenyl)-ethyl]amino]carbonyl]propyl]carbamate;
- 20 (12) Benthiavalicarb-isopropyl corresponding to O-isopropyl [(S)-1-{[(1R)-1-(6-fluoro-1,3-benzothiazol-2-yl)ethyl]-carbamoyl-2-methylpropyl}-carbamate;
- (13) Cymoxanil corresponding to 1-(2-cyano-2-methoxyimino-acetyl)-3-ethylurea;
- 25

- (14) Azoxystrobin corresponding to methyl (E)-2-[2-[6-(2-cyanophenoxy)-pyrimidin-4-yloxy]phenyl-3-methoxyacrylate;
- (15) Metominofen corresponding to N-methyl-(E)-methoxyimino-(2-phenoxyphenyl)acetamide;
- (16) Pyraclostrobin corresponding to methyl N-(2-[1-(4-chlorophenyl)pyrazol-3-yloxymethyl]-phenyl)-N-methoxycarbamate;
- (17) Acibenzolar-S-methyl corresponding to methyl benzo(1,2,3)thiadiazole-7-thiocarboxylate;
- (18) Famoxadone corresponding to 5-methyl-5-(4-phenoxyphenyl)-3-(phenylamino)oxazolidin-2,4-dione;
- (19) Fenamidone corresponding to 4-methyl-4-phenyl-1-(phenylamino)-2-methylthioimidazolidin-5-one;
- (20) Cyazofamide, corresponding to 2-cyano-4-chloro-5-(4-methylphenyl)-1-(N,N-dimethylaminosulfamoyl)-imidazole;
- (21) Fluazinam corresponding to 3-chloro-N-(3-chloro-5-trifluoromethyl-2-pyridyl)- α - α - α -trifluoro-2,6-dinitro-p-toluidine;
- (22) Dimethomorph corresponding to (E,Z)-4-[3-(4-chlorophenyl)-3-(3,4-dimethoxyphenyl)-acryloyl]morpholine; or Flumorph (SYP-L190) corresponding to (E,Z)-4-[3-(4-fluorophenyl)-3-(3,4-dimethoxyphenyl)-acryloyl]morpholine;

- (23) Flumetover corresponding to N,N-diethylamide of 4-trifluoromethyl-6-3,4-dimethoxyphenyl)-benzoic acid;
- (24) Chlorothalonil corresponding to 1,3-dicyano-2,4,5-tetrachlorobenzene;
- 5 (25) Mancozeb corresponding to the manganese and zinc salt of ethylenebis(dithiocarbamate) (polymer);
- (26) Tolyfluanide corresponding to N-dichloro-fluoromethylthio-N',N'-dimethyl-N-p-tolylsulfamide;
- (27) Folpet corresponding to N-(trichloromethyl-10 thio)phthalimide;
- (28) Etridiazole corresponding to ethyl-3-trichloromethyl-1,2,4-thiadiazolyl ether;
- (29) Hymexanol corresponding to 5-methyliso-oxazol-3-ol;
- (30) Propamocarb corresponding to propyl-(3-15 dimethylaminopropyl)carbamate;
- (31) R-3-aminobutanoic acid or RS-3-aminobutanoic acid;
- (32) Zoxamide, corresponding to 3,5-dichloro-N-(3-chloro-1-ethyl-1-methyl-2-oxopropyl)-p-toluamide;
- (33) Ethaboxam, corresponding to (RS) - (α -cyano-2-20 thienyl)-4-ethyl-2(ethylamino)-5-thiazolecarboxyamide;
- (34) Fluopicolide, corresponding to 2,6-dichloro-N-[3-chloro-5-(trifluoromethyl)-2-pyridylmethyl]benzamide;
- 25 (35) Fosetyl, corresponding to ethyl hydrogenphosphonate;

(36) Fosetyl-Al, corresponding to aluminium triethylphosphonate, more commonly known under the trade-name of Aliette.

The compounds (1) are described in Italian patent application Nr. MI98A002583.

Compound (2) is described in patent application WO 98/26654 A2.

Compound (3) is described in "The Pesticide Manual", 1997, XI^a edition, British Crop Protection Council Ed., page 1174.

The compounds (4) are commercial products and their copper salts are described in Italian patent application Nr. MI 2001A002430.

The compounds (5) are easily available on the market.

Compound (6) is described in "The Pesticide Manual", 1983, VIIth edition, British Crop Protection Council Ed., page 32.

Compound (7) is described in English patent GB 1,500,581.

Compound (8) is described in patent application WO 96/01559 A1.

Compound (9) is described in English patent GB 2,058,059.

Compound (10) is described in patent application WO 01/87822.

Compound (11) is described in patent application EP 550,788 and EP 775,696.

5 Compound (12) is described in patent application EP 775,696.

Compound (13) is described in "The Pesticide Manual", 1983, VIIth edition, British Crop Protection Council Ed., page 148.

10 Compound (14) is described in European patent application EP 382,375.

Compound (15), corresponding to the experimental abbreviation SSF-126, is described in American patent application US 5,185,242.

15 Compound (16) is described in patent application WO 96/01258.

Compound (17) is described in American patent application US 4,931,581.

20 Compound (18) is described in "Brighton Crop Protection Conference - Pests and Diseases" 1996, Congress Records.

Compound (19) is described in European patent application EP 629,616.

25 Compound (20), also called IKF916, is described in European patent application EP 705,823.

Compound (21) is described in European patent application EP 31,257.

The compounds (22) are described respectively in European patent application EP 219,756 and in "Brighton
5 Crop Protection Conference - Pests and Diseases" 2000, Congress Records.

Compound (23) is described in European patent applications EP 360,701 and EP 611,232.

Compound (24) is described in "The Pesticide Manual", 1983, VIIth edition, British Crop Protection Council Ed., page 120.
10

Compound (25) is described in "The Pesticide Manual", 1983, VIIth edition, British Crop Protection Council Ed., page 339.
15

Compound (26) is described in "The Pesticide Manual", 1983, VIIth edition, British Crop Protection Council Ed., page 537.

Compound (27) is described in "The Pesticide Manual", 1983, VIIth edition, British Crop Protection Council Ed., page 599.
20

Compound (28) is described in "The Pesticide Manual", 1983, VIIth edition, British Crop Protection Council Ed., page 252.

Compound (29) is described in "The Pesticide Manual", 1983, VIIth edition, British Crop Protection Council Ed., page 314.
25

Compound (30) is described in "The Pesticide Manual", 1983, VIIth edition, British Crop Protection Council Ed., page 471.

Compound (31) is described in European patent application EP 753,258.

Compound (32) is described in "Brighton Crop Protection Conference - Pests and Diseases" 1998, Congress Records.

Compound (33) is described in "The Pesticide Manual", 2003, XIII^a edition, British Crop Protection Council Ed.

Compound (34) is described in patent application WO 200111966.

The compounds (35) and (36) are described in "The Pesticide Manual", 1994, X^a edition, British Crop Protection Council Ed., page 530.

A further object of the present invention therefore relates to fungicidal compositions comprising one or more quaternary salts having general formula (I), for q which has the value of 0 for X = sulfur and the value of 1 for X = nitrogen, and fungicidal compositions also containing other active principles.

Preferred fungicidal compositions according to the present invention can be selected from:

1. Acid choline phosphite and IR5885

2. Neutral choline phosphite and IR5885
3. Choline ethylphosphonate and IR5885
4. Choline ethylphosphonate and IR6141
5. Acid choline phosphite and IR6141
- 5 6. Neutral choline phosphite and IR6141
7. Acid choline phosphite and Benthiavalicarb-isopropyl;
8. Neutral choline phosphite and Benthiavalicarb-isopropyl;
- 10 9. Acid choline phosphite and ASA_2Cu ;
10. Acid choline phosphite and SA_2Cu ;
11. Acid choline phosphite and SACu ;
12. Neutral choline phosphite and ASA_2Cu ;
13. Neutral choline phosphite and SA_2Cu ;
- 15 14. Neutral choline phosphite and SACu ;
15. Acid laurylcholine phosphite and IR5885;
16. Neutral laurylcholine phosphite and IR5885;
17. Acid laurylcholine phosphite and IR6141;
18. Neutral laurylcholine phosphite and IR6141;
- 20 19. Acid choline phosphite and iprovalicarb;
20. Neutral choline phosphite and iprovalicarb;
21. Acid cocamidopropylcholine phosphite and IR5885;
22. Neutral cocamidopropylcholine phosphite and IR5885;
- 25 23. Acid cocamidopropylcholine phosphite and IR6141;

24. Neutral cocamidopropylcholine phosphite and IR6141;
25. Acid cocamidopropylcholine phosphite and copper oxychloride;
- 5 26. Neutral cocamidopropylcholine phosphite and copper oxychloride;
27. Acid phosphite of the methyl ester of cocamidopropylbetaine and IR5885;
28. Neutral phosphite of the methyl ester of cocamidopropylbetaine and IR5885;
- 10 29. Acid phosphite of the methyl ester of cocamidopropylbetaine and IR6141;
30. Neutral phosphite of the methyl ester of cocamidopropylbetaine and IR6141;
- 15 31. Acid phosphite of the methyl ester of cocamidopropylbetaine and Airone;
32. Neutral phosphite of the methyl ester of cocamidopropylbetaine and Airone;
33. Choline salicylate and IR5885;
- 20 34. Choline acetylsalicylate and IR5885;
35. Choline saccharinate and IR5885;
36. Choline bicarbonate and IR5885;
37. Acid choline phosphite and Mandipropamid;
38. Neutral choline phosphite and Mandipropamid;
- 25 39. Acid choline phosphite and copper oxychloride;

40. Neutral choline phosphite and copper oxychloride;
41. Acid choline phosphite and Airone;
42. Neutral choline phosphite and Airone;
43. Acid choline phosphite and Ethaboxam;
5 44. Neutral choline phosphite and Ethaboxam;
45. Choline bicarbonate and Fosetyl-Al;
46. Choline chloride and Fosetyl-Al;
47. Choline salicylate and Fosetyl-Al;
48. Choline acetylsalicylate and Fosetyl-Al;
10 49. Cocamidopropylcholine chloride and Fosetyl-Al;
50. Laurylcholine chloride and Fosetyl-Al;
51. Choline bicarbonate and Fosetyl;
52. Choline chloride and Fosetyl.

Particularly preferred are the compositions defined
15 with the following numbers: 1., 2., 3., 4., 5., 6., 7.,
8., 9., 10., 11., 12., 13., 19., 20., 33., 34., 35., 36.,
45., 46., 52.

A further object of the present invention therefore
relates to the use of fungicidal compositions comprising
20 a compound having general formula (I), when q has the
value of 0 for X = nitrogen, and the fungicidal compound
(36).

Another object of the present invention relates to
the use of fungicidal compositions comprising one or more
25 compounds having general formula (I), when q has the

value of 0 for X = sulfur and the value of 1 for X = nitrogen, and one or more fungicidal compounds, in particular one or more fungicidal compounds (1)-(36), even more in particular compounds (1), (2), (4), (5), (35) and
5 (36), for the control of phytopathogen fungi.

An object of the present invention also relates to a method for the control of phytopathogen fungi in agricultural crops by the application of the compounds having general formula (I) or fungicidal compositions comprising
10 one or more compounds having general formula (I) and one or more fungicidal compounds, in particular one or more fungicidal compounds (1)-(36), even more in particular compounds (1), (2), (4), (5), (35) e (36), for the control of phytopathogen fungi.

15 Examples of phytopathogen fungi and bacteria controlled by the above compounds having general formula (I) and by the above compositions, together with examples of application crops, are indicated below for purely illustrative and non-limiting purposes:

- 20 - *Helminthosporium spp* on cereals;
- *Erysiphe spp* on cereals;
- *Puccinia spp.* on cereals;
- *Plasmopara viticola* on vines;
- *Pythium spp* on vegetables;
25 - *Phytophthora spp.* on vegetables;

- *Rhynchosporium* on cereals;
- *Septoria* spp. on cereals;
- *Sphaerotheca fuliginea* on cucurbits (for example cucumbers);
- 5 - *Podosphaera leucotricha* on apple trees;
- *Pyricularia oryzae* on rice;
- *Uncinula necator* on vines;
- *Venturia* spp. on fruit trees;
- *Botrytis cinerea* on vines and vegetables;
- 10 - *Fusarium* spp. on cereals;
- *Alternaria* spp. on fruit trees and vegetables;
- *Cercospora* spp. on sugar beet;
- *Xantomonas*;
- *Bacillus* spp.

15 The compounds having general formula (I) and mixtures thereof with one or more fungicidal compounds are capable of exerting a bactericidal/fungicidal action of both a curative and preventive nature and have a low or zero phytotoxicity.

20 A further object of the present invention therefore relates to a method for the control of phytopathogen fungi and bacteria in agricultural crops by the application of the compounds having general formula (I) having a direct fungicidal and bactericidal activity and a method
25 for the stimulation of the natural defense systems of

plants from abiotic stress (temperature, salinity, drought, etc.) and biotic stress and the induction of resistance in the plants themselves by the application of the compounds having general formula (I).

5 The quantity of compound to be applied for obtaining the desired effect can vary in relation to various factors such as, for example, the compound used, the crop to be preserved, the type of pathogen, the degree of infection, the climatic conditions, the application method and
10 the formulation adopted.

Doses of compound ranging from 10 g to 5 kg per hectare generally provide a sufficient control.

For practical uses in agriculture it is often convenient to apply fungicidal compositions containing one
15 or more compounds having general formula (I).

The application of these compositions can be effected on all parts of the plant, for example on the leaves, stems, branches and roots, or on the seeds themselves before sowing, or on the ground in which the plant
20 grows.

Compositions can be used in the form of dry powders, wettable powders, emulsifying concentrates, micro-emulsions, pastes, granulates, solutions, suspensions, etc.: the choice of the type of formulation will depend
25 on the specific use.

The compositions are prepared in the known way, for example by diluting or dissolving the active substance with a solvent medium and/or a solid diluent, possibly in the presence of surface-active agents.

5 Solid diluents or supports which can be used are, for example: silica, kaolin, bentonite, talc, infusorial earth, dolomite, calcium carbonate, magnesia, gypsum, clays, synthetic silicates, attapulgite, sepiolite.

Liquid diluents which can be used, are for example,
10 in addition to water, aromatic organic solvents (xylols or alkyl benzol mixtures, chlorobenzene, etc.), paraffins (oil fractions), alcohols (methanol, propanol, butanol, octanol, glycerin, etc.), esters (ethyl acetate, isobutyl acetate, etc.), ketones (cyclohexanone, acetone, aceto-
15 phenone, isophorone, ethylamylketone, etc.), amides (N,N-dimethylformamide, N-methylpyrrolidone, etc.).

Surface-active agents which can be used are salts of sodium, calcium, triethylamine or triethanolamine, alkyl-sulfonates, alkylaryl-sulfonates, polyethoxylated alkyl-
20 phenols, polyethoxylated esters of sorbitol, ligninsulfonates, etc.

The compositions can also contain special additives for particular purposes, for example adhesion agents such as gum arabic, polyvinyl alcohol , polyvinylpyrrolidone,
25 polyacrylates, etc..

In the fungicidal compositions, object of the present invention, the concentration of active substances varies from 0.1% to 98% by weight, preferably from 0.5 to 90% by weight.

5 If desired, it is possible to also add compatible active principles to the compositions, object of the present invention, such as, for example, phyto-regulators, antibiotics, herbicides, insecticides, fertilizers.

The following examples are provided for a better understanding of the invention for illustrative and non-limiting purposes of the present invention.

EXAMPLE 1

Preparation of neutral choline phosphite (Compound 1).

15 A solution of 16.10 g of potash in 20 ml of water is added dropwise under cooling to a solution of 10 g of phosphorous acid in 5 ml of water.

The mixture is kept under stirring at room temperature and 34.07 g of choline chloride are added in portions.

20 At the end of the addition, the mixture is left under stirring for 3 hours and the solution thus obtained is used as such.

EXAMPLE 2

25 Preparation of acid choline phosphite (Compound 2).

20.15 g of choline bicarbonate are added in portions, under cooling, to a solution of 10 g of phosphorous acid in 3 ml of water.

At the end of the addition, the mixture is left under stirring for 4 hours and the solution thus obtained is used as such.

EXAMPLE 3

Preparation of cholesterylcarbonylamidopropyldimethylamine.

3.41 g of 3-dimethylamino-1-propylamine are added to a solution of 15 g of cholesterylchloroformate in 70 ml of methylene chloride and 3.49 ml of triethylamine. The mixture is kept under stirring at room temperature for a night. The product obtained is extracted, washed with water, anhydrified with Na_2SO_4 obtaining, after drying, 15.8 g of the desired compound (yield: 92%).

Elemental analysis [% found (theoretical)] =

C 77.0 (76.8); H 11.9 (11.2); N 5.1 (5.4).

EXAMPLE 4

Preparation of cholesterylcarbonylamidopropylcholine chloride (Compound 3).

12 g of cholesterylcarbonylamidopropyldimethylamine in 32 ml of water are charged into a reactor and 1.9 g of 2-chloroethanol are added. The reaction mixture is slowly heated to 98°C. After about 5 hours, the starting prod-

ucts are completely used up and the solution obtained is used as such.

EXAMPLE 5

Preparation of neutral cholesterylcarbonylamidopropylcholine phosphite (Compound 4).

A solution of 3.22 g of potash in 4 ml of water is added dropwise, under cooling, to a solution of 2 g of phosphorous acid in 2 ml of water.

The mixture is maintained under stirring at room temperature and 28.9 g of cholesterylcarbonylamidopropylcholine chloride are added.

At the end of the addition the mixture is left under stirring for 3 hours, and the solution thus obtained is used as such.

15 EXAMPLE 6

Preparation of acid cholesterylcarbonylamidopropylcholine phosphite (Compound 5).

A solution of 1.61 g of potash in 2 ml of water is added dropwise, under cooling to a solution of 2 g of phosphorous acid in 2 ml of water.

The mixture is maintained under stirring at room temperature and 14.45 g of cholesterylcarbonylamidopropylcholine chloride are added.

At the end of the addition the mixture is left under stirring for 3 hours, and the solution thus obtained is

used as such.

The following compounds were prepared analogously to what is described in the examples:

Table 1

Number	Compound
6	Acid laurylcholine phosphite
7	Neutral laurylcholine phosphite
8	Acid cocamidopropylcholine phosphite
9	Neutral cocamidopropylcholine phosphite
10	Acid stearylcholine phosphite
11	Neutral stearylcholine phosphite
12	Acid N,N-dimethyl,N-laurylamidopropyl [L] valinol phosphite
13	Neutral N,N-dimethyl,N-laurylamidopropyl - [L] valinol phosphite
14	Choline salicylate
15	Choline acetylsalicylate
16	Choline saccharinate
17	Choline cyclamate
18	Choline taurinate
19	Choline sulfate
20	Acid choline phosphate
21	Neutral choline phosphate
22	Laurylcholine salicylate
23	Laurylcholine acetylsalicylate

24	Laurylcholine saccharinate
25	Laurylcholine cyclamate
26	Laurylcholine taurinate
27	Cocamidopropylcholine salicylate
28	Cocamidopropylcholine acetylsalicylate
29	Cocamidopropylcholine saccharinate
30	Cocamidopropylcholine cyclamate
31	Cocamidopropylcholine taurinate
32	Acid phosphite of the methyl ester of cocamido- propylbetaine
33	Neutral phosphite of the methyl ester of cocami- dopropylbetaine
34	Acid phosphate of the methyl ester of cocamido- propylbetaine
35	Neutral phosphate of the methyl ester of cocami- dopropylbetaine
36	Bicarbonate of the methyl ester of cocamidopro- pylbetaine
37	Salicylate of the methyl ester of cocamidopropyl- betaine
38	Saccharinate of the methyl ester of cocamidopro- pylbetaine
39	Acid phosphite of the cetyl ester of cocamidopro- pylbetaine
40	Neutral phosphite of the cetyl ester of cocamido-

	propylbetaine
41	Salicylate of the cetyl ester of cocamidopropylbetaine
42	Saccharinate of the cetyl ester of cocamidopropylbetaine
43	Acetylsalicylate of the cetyl ester of cocamidopropylbetaine
44	Cyclamate of the cetyl ester of cocamidopropylbetaine
45	Acid phosphite of the methyl ester of carnitine
46	Neutral phosphite of the methyl ester of carnitine
47	Salicylate of the methyl ester of carnitine
48	Acetylsalicylate of the methyl ester of carnitine
49	Saccharinate of the methyl ester of carnitine
50	Choline ethylphosphonate
51	Laurylcholine ethylphosphonate
52	Cocamidopropylcholine ethylphosphonate

EXAMPLE 7

Determination of the fungicidal activity against peronospora of vines (*Plasmopara viticola*). Table 2

Vine leaves (cultivar Dolcetto), grown in vases in a
5 conditioned environment ($20\pm 1^{\circ}\text{C}$, 70% relative humidity)
are treated by spraying both sides of the leaves with
compounds 1 and 2, dispersed in a hydroacetone solution

at 20% by volume in acetone.

After remaining 24 hours in a conditioned environment, the plants were sprayed on both sides of the leaves with an aqueous suspension of conidia of *Plasmopara viticola* (20,000 conidia per cm³).

The plants are kept in a humidity saturated environment at 21°C for the incubation period of the fungus.

At the end of this period (7 days), the fungicidal activity is evaluated according to an evaluation percentage scale from 0 (completely infected plant) to 100 (healthy plant).

Table 2

7-day preventive activity on *Plasmopara viticola* of the compounds having general formula (I)

Compound N°	Activity 1000 ppm*	Activity 500 ppm*
1 (neutral choline phosphite)	100	90
2 (acid choline phosphite)	95	85
K ₂ HPO ₃	80	45
Choline chloride	57	35
Choline bicarbonate	84	75

* the doses in ppm refer to the quantity of equivalent phosphorous acid.

EXAMPLE 8

Determination of the fungicidal activity against perono-

spora of potatoes (*Phytophthora infestans*). Table 3

Leaves of potato plants, Primura variety, grown in vases in a conditioned environment ($20\pm 1^{\circ}\text{C}$, 70% relative humidity) were infected with an aqueous suspension of
5 spores of *Phytophthora infestans* (100,000 spores per cm^3).

After remaining 24 hours in a conditioned environment, both sides of the leaves were sprayed with the products under examination dissolved in a hydroacetone solution at 20% of acetone (vol./vol.).

10 After drying, the plants were transferred for the incubation period of the fungus (4 days) in a conditioned environment at 70% of relative humidity and 24°C .

The intensity of the disease was finally evaluated according to an evaluation percentage scale from 100
15 (healthy plant) to 0 (completely infected plant).

Table 3

1-day curative activity on *Phytophthora infestans* of the compounds having general formula (I)

Compound	Activity
20 Choline chloride	15 (2690 ppm*)
K_2HPO_3	40 (1575 ppm**)
2(acid choline phosphite)	65 (1575 ppm**)

* the dose in ppm refers to the quantity of choline con-
25 tained in Compound N°2

** the doses in ppm refer to the quantity of equivalent phosphorous acid.

From the data indicated in Table 3, it is possible to assert the synergic effect of the ionic couple present in
5 Compound Nr. 2.

When in fact the fungicidal activity found experimentally (65) is greater than that expected (49), calculated considering the contribution of potassium phosphite and choline chloride and using the Limpel formula ("Pesticide
10 Science" (1987), vol. 19, pages 309-315), then this activity should be considered as being a synergic effect with a synergy factor equal to 1.4 (experimental activity/activity calculated according to Limpel).

EXAMPLE 9

15 Determination of the fungicidal activity of fungicidal mixtures against peronospora of potatoes (*Phytophthora infestans*). Tables 4-6

Leaves of potato plants, Primura variety, grown in vases in a conditioned environment ($20\pm 1^{\circ}\text{C}$, 70% relative
20 humidity) were infected with an aqueous suspension of spores of *Phytophthora infestans* (100,000 spores per cm^3).

After remaining 24 hours in a conditioned environment, both sides of the leaves were sprayed with the fungicidal mixtures under examination dissolved in a hydro-
25 acetone solution at 20% of acetone (vol./vol.).

After drying, the plants were transferred for the incubation period of the fungus (4 days) in a conditioned environment at 70% of relative humidity and 24°C.

The intensity of the disease was finally evaluated according to an evaluation percentage scale from 100 (healthy plant) to 0 (completely infected plant).

From the data indicated in tables 4-6, it is possible to assert the synergic effect of the mixtures, consisting of the mixtures under examination, compared with the efficacy expected using the Limpel formula ("Pesticide Science" (1987), vol. 19, pages 309-315):

$$E = x + y - (xy/100)$$

wherein:

- E is the fungicidal activity expected, without synergic effects, from a mixture obtained by mixing g.x of compound X with g.y of compound Y;

- x is the activity of compound X when used alone at a dose of g.x;

- y is the activity of compound Y when used alone at a dose of g.y.

When the fungicidal activity found experimentally is greater than the value of E, this activity should be considered a synergic effect.

Table 4

1-day curative activity on *Phytophthora infestans* of the fungicidal mixtures consisting of Fosetyl-Al which at 1600 ppm* (g.x) is 0 (x) with compounds having general formula (I) when q has the value of 0 for X = nitrogen.

Mixture	Dose ppm (g.y)	Activ- ity (y)	Activity mixture according to Limpel (E)	Experimen- tal activity of mixture	Synergy factor
10 Mixture 1. (N,N-dimethylethanolamine and Fosetyl-Al)	1736	30	30	90	3
Mixture 2. (3-dimethylamine-1-propanol and Fosetyl-Al)	1745	25	25	65	2.6

* the doses in ppm refer to the quantity of equivalent phosphorous acid.

Table 5

1-day curative activity on *Phytophthora infestans* of fungicidal mixtures consisting of IR5885 which at 150 ppm (g.x) is 15 (x) with compounds having general formula (I) when q has the value of 0 for X = sulfur and the value of 1 for X = nitrogen.

Mixture	Dose ppm* (g. <u>y</u>)	Activity (y)	Activity of mixture ac- cording to Limpel (E)	Experimen- tal activ- ity of mix- ture	Synergy factor
Mixture 3. (IR5885 and acid cho- line phos- phite, compound 1)	1575	65	70.25	98	1.4
Mixture 4. (IR5885 and neutral choline phosphite, compound 2)	1575	65	70.25	90	1.3

* the doses in ppm refer to the quantity of equivalent phosphorous acid.

Table 6

1-day curative activity on *Phytophthora infestans* of
 5 fungicidal mixtures consisting of Fosetyl-Al which at
 1600 ppm* (g.x) is 0 (x) with other compounds having gen-
 eral formula (I) when q has the meaning of 0 for X = sul-
 fur and the value of 1 for X = nitrogen.

Mixture	Dose ppm (g.y)	Activity (y)	Activity of mixture ac- cording to Limpel (E)	Experimen- tal activ- ity of mix- ture	Synergy factor
Mixture 5. (Choline chloride and Fose- tyl-Al)	2730	20	70.25	75	1.4

* the doses in ppm refer to the quantity of equivalent phosphorous acid.

EXAMPLE 10

Determination of the fungicidal activity of fungi-
5 cidal mixtures against peronospora of vines (*Plasmopara viticola*). Table 7.

Leaves of vines (cultivar Dolcetto), grown in vases
in a conditioned environment ($20 \pm 1^\circ\text{C}$), 70% relative hu-
midity) were infected with an aqueous suspension of
10 spores of *Plasmopara viticola* (200,000 conidia per cm^3).

After remaining 24 hours in a conditioned environ-
ment, both sides of the leaves were sprayed with the fun-
gicidal mixtures under examination dissolved in a hydro-
acetone solution at 20% of acetone (vol./vol.).

15 After drying, the plants were transferred for the in-
cubation period of the fungus (6 days) to a conditioned
at 70% relative humidity and 24°C .

At the end of this period (7 days), the fungicidal activity is evaluated according to an evaluation percentage scale from 0 (completely infected plant) to 100 (healthy plant).

5 From the data indicated in table 7, it is possible to assert the synergic effect of the mixtures under examination, compared with the expected efficacy using the Limpel formula already described in example 9.

Table 7

10 1-day curative activity on *Plasmopara viticola* of fungicidal mixtures consisting of Fosetyl-Al which at 1200 ppm* (g.x) is 23 (x) with other compounds having general formula (I).

Mixture	Dose ppm (g.y)	Activity (y)	Activity of mixture ac- cording to Limpel (E)	Experimen- tal activ- ity of mix- ture	Synergy factor
Mixture 1. (N,N- dimethy- lethano- lamina and Fosetyl-Al)	1302	25	42.25	100	2.4
Mixture 5. (Choline chloride and Fose- tyl-Al)	2047	20	34.4	100	2.9

15 * the doses in ppm refer to the quantity of equivalent phosphorous acid.

EXAMPLE 11

Determination of the fungicidal activity of fungicidal mixtures against peronospora of tobacco (*Plasmopara tabacina*). Table 8.

Leaves of tobacco plants (cultivar Barley), grown in
5 vases in a conditioned environment ($20\pm 1^{\circ}\text{C}$), 70% relative humidity) were infected with an aqueous solution of spores of *Plasmopara tabacina* (200,000 conidia per cm^3).

After remaining 24 hours in a conditioned environment, both sides of the leaves were sprayed with the fungicidal mixtures under examination dissolved in a hydro-
10 acetone solution at 20% of acetone (vol./vol.).

After drying, the plants were transferred for the incubation period of the fungus (6 days) to a conditioned environment at 70% relative humidity and 24°C .

15 At the end of this period (7 days), the fungicidal activity is evaluated according to an evaluation percentage scale from 0 (completely infected plant) to 100 (healthy plant).

From the data indicated in table 8, it is possible to
20 assert the synergic effect of the mixtures under examination, compared with the expected efficacy using the Limpel formula already described in example 9.

Table 8

1-day curative activity on *Plasmopara tabacina* of
25 fungicidal mixtures consisting of Fosetyl-Al which at

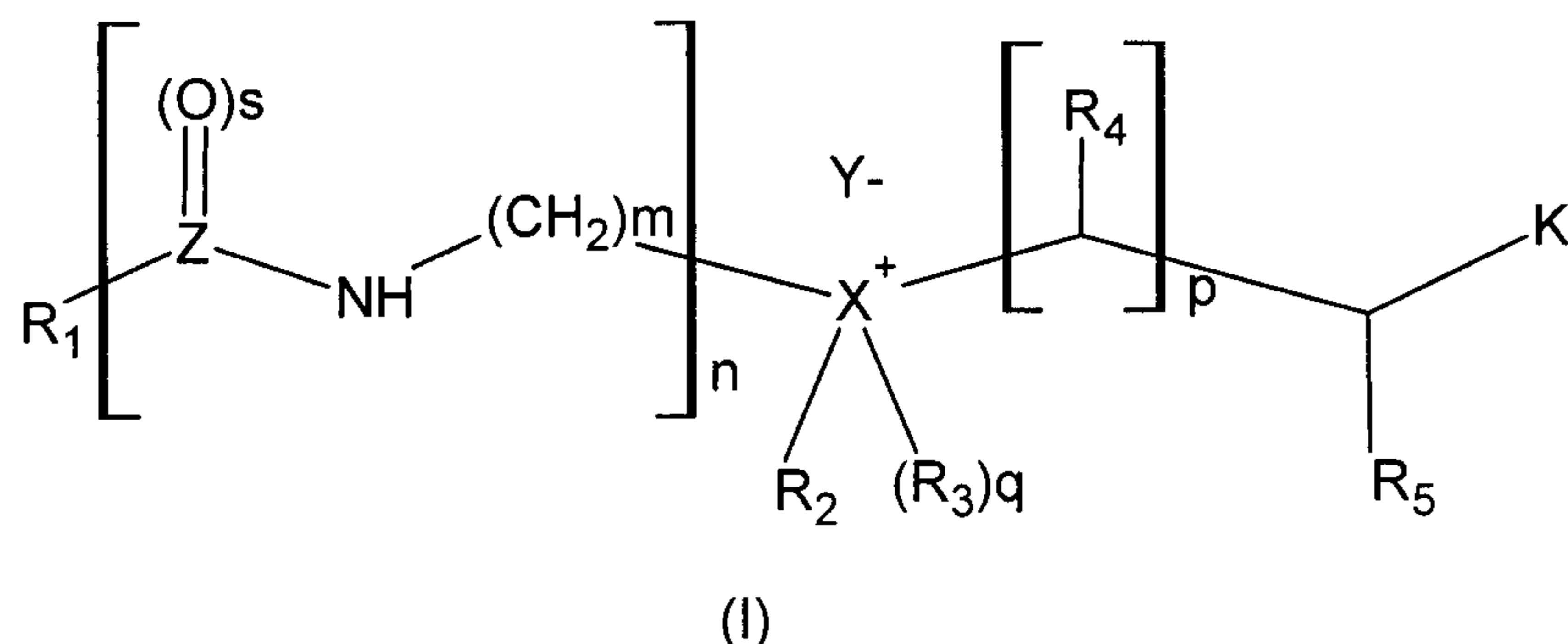
1200 ppm* (g.x) is 15 (x) with other compounds having general formula (I).

Mixture	Dose ppm (<u>g.y</u>)	Activity (y)	Activity of mixture ac- cording to Limpel (E)	Experimen- tal activ- ity of mix- ture	Synergy factor
Mixture 1. (N,N- dimethy- lethano- lamina and Fosetyl-Al)	1302	5	19	100	5.3
Mixture 5. (Choline chloride and Fose- tyl-Al)	2047	10	23.5	100	4.3

* the doses in ppm refer to the quantity of equivalent
5 phosphorous acid.

CLAIMS

1. An organic compound of general formula (I),



5 wherein:

K represents a CH₂OH or COOR_a group.

R_a represents a linear or branched C₁-C₂₆ alkyl group optionally substituted;

R₁ represents a hydrogen or a linear or branched C₁-C₂₆ alkyl group optionally substituted; a linear or branched C₁-C₂₆ haloalkyl group optionally substituted; a
 10 linear or branched C₁-C₂₆ alkoxy group optionally substituted; a linear or branched C₁-C₂₆ alkylthio group optionally substituted; a linear or branched C₂-C₂₆ alkenyl group optionally substituted; a linear or branched C₂-C₂₆ alkynyl group optionally substituted; a C₃-C₃₀ cycloalkyl group optionally condensed or a condensed C₁₇ cycloalkyl group of the steroid type optionally substituted; a
 15 C₃-C₃₀ cycloalkoxy group optionally condensed and optionally substituted; a heterocyclic group optionally substituted; an aryl group optionally substituted; a heteroaryl group optionally substituted; a linear or cyclic C₆-C₁₂ group of the saccharide type optionally substituted; or, when n is different from 0, a C₁-C₂₆ alkylamine group or a C₂-C₂₆ di-alkylamine group optionally substituted;

20 R₂ and R₃, the same or different, represent a C₁-C₃ alkyl group optionally substituted;

R₄ and R₅, the same or different, represent a hydrogen atom, or a linear or branched C₁-C₆ alkyl group optionally substituted; a linear or branched C₂-C₆ alkenyl group optionally substituted; a C₃-C₆ cycloalkyl group optionally substituted; a hydroxyl group; an aryl group optionally substituted; a heteroaryl group optionally substituted; or a heterocyclic group optionally substituted;

R₄ and R₅ can individually form a cycle together with R₂;

X represents a nitrogen atom;

Z represents a carbon or sulfur atom;

m represents a number ranging from 1 to 5;

n and p represent a number ranging from 0 to 3 ;

q has the value of 1;

Y⁻ represents a phosphite anion H₂PO₃⁻ ; and

s has the value of 1 for Z = carbon or the value of 2 for Z = sulfur.

2. The compound according to claim 1, characterized in that the C₁-C₂₆ alkyl group is methyl, ethyl, propyl, isopropyl, butyl, isobutyl, tert-butyl, capryl, lauryl, stearyl, eicosyl or hexacosyl.

3. The compound according to claim 1, characterized in that the C₁-C₂₆ haloalkyl group is fluoromethyl, difluoromethyl, trifluoromethyl, trichloromethyl, 2,2,2-trifluoroethyl, 2,2,2-trichloroethyl, 2,2,3,3-tetrafluoropropyl, 2,2,3,3,3-pentafluoro-propyl, perfluorooctanyl or perfluorododecyl .

4. The compound according to claim 1, characterized in that the C₁-C₂₆ alkoxy group is methoxy, ethoxy, isopropoxy, cyclopropylmethoxy or lauryloxy.

5. The compound according to claim 1, characterized in that the C₁-C₂₆ thioalkyl group is thiomethyl, thioethyl, thiolauryl or thiocapryl .

6. The compound according to claim 1, characterized in that the C₂-C₂₆ alkenyl group is ethenyl, propenyl, butenyl, 1-decenyl, 8-heptadecenyl, 8,11,14-heptadecatrienyl or 8, 11-heptadecadienyl .
7. The compound according to claim 1, characterized in that the C₂-C₂₆ alkynyl group is ethynyl, propargyl, 1-dodecynyl or 1-octadecynyl.
8. The compound according to claim 1, characterized in that the C₃-C₃₀ cycloalkyl group optionally condensed is cyclopropyl, 2, 2-dichlorocyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, decaline or abietyl .
9. The compound according to claim 1, characterized in that the condensed C₁₇ cycloalkyl group of the steroid type is cholanyl, chenodeoxycholanyl , ursodeoxycholanyl, deoxycholanyl, iodeoxycholanyl or lithocholanyl .
10. The compound according to claim 1, characterized in that the C₃-C₃₀ cycloalkoxy group is cyclopentoxy, cyclohexyloxy or cholesteryl .
11. The compound according to claim 1, characterized in that the C₁-C₂₆ alkylamine or C₂-C₂₆ dialkylamine groups are methylamine, dimethylamine, ethylamine, isopropylamine, dibutylamine, dioctylamine, hexadecylamine or didecylamine.
12. The compound according to claim 1, characterized in that the aryl group is phenyl, naphthyl or phenanthryl .
13. The compound according to claim 1, characterized in that the heteroaryl group is pyridine, pyrimidine, pyridazine, pyrazine, triazine, tetrazine, quinoline, quinoxaline, quinazoline, furan, thiophene, pyrrol, oxazole, thiazole, isoxazole, isothiazole, oxadiazole, thiadiazole, pyrazole, imidazole, triazole, tetrazole, indole, benzofuran, benzothiophene, benzoxazole, benzothiazole, benzoxadiazole, benzothiadiazole, benzopyrazole, benzimidazole, benzotriazole, triazolepyridine, triazolepyrimidine, thiazoletriazole or cumarin.

14. The compound according to claim 1, characterized in that the heterocyclic group is pyrrolidine, piperidine, dihydropyridine, piperazine, 2,6-diketopiperazine, 2-ketoazetidine, morpholine, thiazine or indoline.

15. The compound according to claim 1, characterized in that the linear or cyclic C₆-C₁₂ group of the saccharide type is gluconyl, glucopyranosyl, β-D-fructofuranosyl-α-D-glucopyranosyl or 4-O-β-D-galactopyranosyl-D-glucosyl.

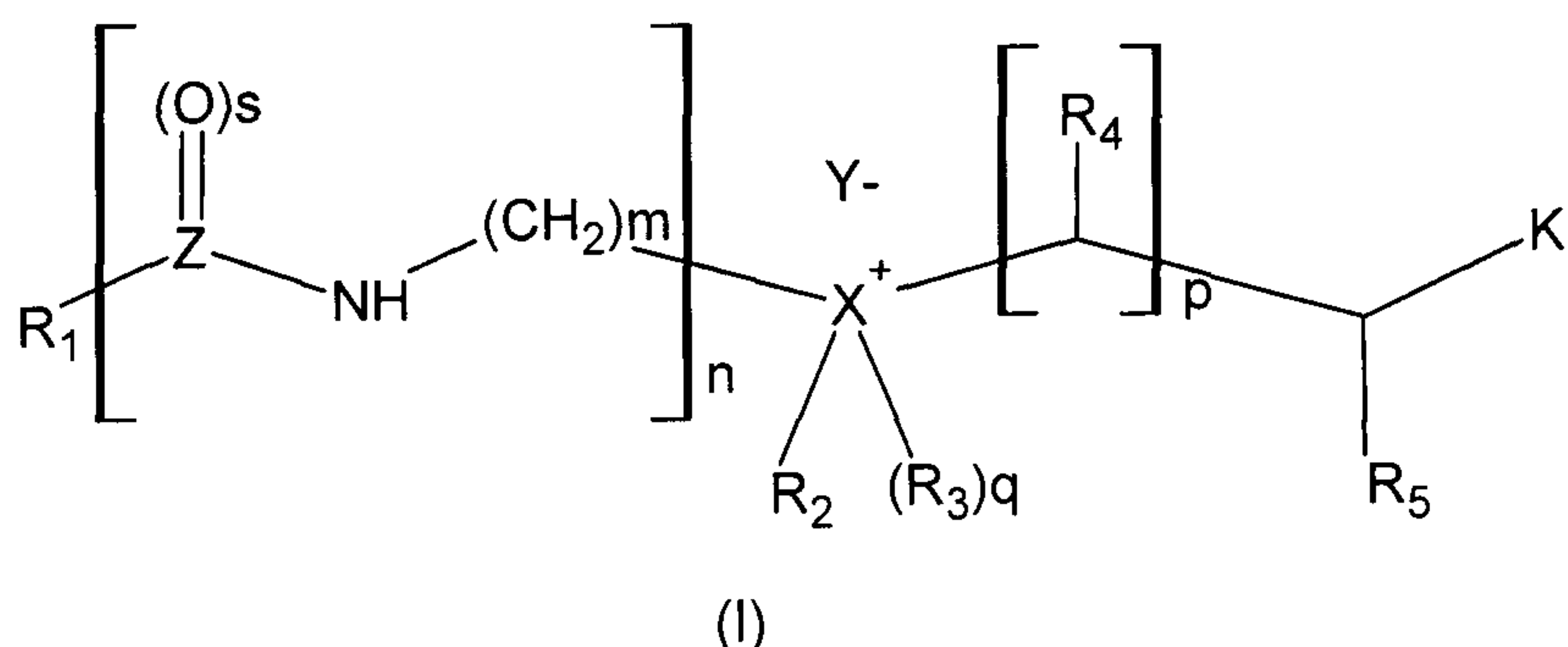
16. The compound according to claim 1, characterized in that it is:

- acid choline phosphite;
- acid laurylcholine phosphite;
- 10 • acid cocamidopropylcholine phosphite;
- acid stearylcholine phosphite;
- acid cholesterylcarbonylamidopropylcholine phosphite;
- acid cholanylamidopropylcholine phosphite;
- acid chenodeoxycholanylamidopropylcholine phosphite;
- 15 • acid N,N-dimethyl,N-laurylamidopropyl[L]valinol phosphite;
- acid N,N-dimethyl,N-lauryl[L]valinol phosphite;
- acid N-lauryl,N-methyl[L]2-pyrrolidinemethanol phosphite;
- acid phosphite of the methyl ester of cocamidopropylbetaine ;
- acid phosphite of betaine cetyl ester;
- 20 • acid phosphite of the methyl ester of cholanylamidopropylbetaine; or
- acid phosphite of the methyl ester of carnitine.

17. The compound according to claim 1, characterized in that wherein the substituents R_1 , R_2 , R_3 , R_4 , R_5 contain optic or geometric isomerism centres, they can be present in all possible configurational isomeric forms.

18. Use of a compound of general formula (I)

5



wherein

K represents a CH_2OH or COOR_a group.

R_a represents a linear or branched $\text{C}_1\text{-C}_{26}$ alkyl group optionally substituted;

- 10 R_1 represents a hydrogen or a linear or branched $\text{C}_1\text{-C}_{26}$ alkyl group optionally substituted; a linear or branched $\text{C}_x\text{-C}_{26}$ haloalkyl group optionally substituted; a linear or branched $\text{C}_1\text{-C}_{26}$ alkoxy group optionally substituted; a linear or branched $\text{C}_1\text{-C}_{26}$ alkylthio group optionally substituted; a linear or branched $\text{C}_2\text{-C}_{26}$ alkenyl group optionally substituted; a linear or branched $\text{C}_2\text{-C}_{26}$ alkynyl group optionally substituted; a $\text{C}_3\text{-C}_{30}$ cycloalkyl group optionally condensed or
- 15 a condensed C_{17} cycloalkyl group of the steroid type optionally substituted; a $\text{C}_3\text{-C}_{30}$ cycloalkoxy group optionally condensed and optionally substituted; a heterocyclic group optionally substituted; an aryl group optionally substituted; a heteroaryl group optionally substituted; a linear or cyclic $\text{C}_6\text{-C}_{12}$ group of the saccharide type optionally substituted; a $\text{C}_1\text{-C}_{26}$ alkylamine group or a $\text{C}_2\text{-C}_{26}$ di-
- 20 alkylamine group optionally substituted for n different from 0;

R_2 and R_3 , the same or different, represent a C_1 - C_3 alkyl group optionally substituted;

R_4 and R_5 , the same or different, represent a hydrogen atom, or a linear or branched C_1 - C_6 alkyl group optionally substituted; a linear or branched C_2 - C_6 alkenyl group optionally substituted; a C_3 - C_6 cycloalkyl group optionally substituted; a hydroxyl group; an aryl group optionally substituted; a heteroaryl group optionally substituted or a heterocyclic group optionally substituted;

R_4 and R_5 can individually form a cycle together with R_2 ;

X represents a nitrogen atom;

Z represents a carbon or sulfur atom;

m represents a number ranging from 1 to 5;

n and p represent a number ranging from 0 to 3 ;

q has the value of 1;

Y^- represents a phosphite anion $H_2PO_3^-$; and

s has the value of 1 for Z = carbon or the value of 2 for Z = sulfur;

for the control of phytopathogenic fungi and bacteria.

19. Use of a compound according to any one of claims 2 to 17 for the control of phytopathogenic fungi and bacteria.

20. Use of a compound according to any one of claims 1 to 17 for the control of phytopathogenic fungi and bacteria by the stimulation of the natural defense systems of plants and the induction of resistance in the plants themselves.

21. Use according to any one of claims 18 to 20, characterized in that said use is curative and/or preventive.

22. Use according to any one of the claims 18 to 21 wherein the compound of general formula (I) is used in a quantity ranging from 10 g to 5 kg per hectare.

23. Use according to any one of claims 18 to 21 wherein the compound of general formula (I) is in the form of single isomer or as isomeric mixtures in any
5 proportion.

24. A method for the control of phytopathogenic fungi and bacteria in agricultural crops, for the stimulation of the natural defense systems of plants from abiotic and biotic stresses, and for the induction of resistance in plants themselves, by the application of a compound of general formula (I) according
10 to any one of claims 1 to 17.

25. A fungicidal composition comprising one or more compounds of general formula (I) according to claim 1, and a liquid or solid diluent.

26. The composition according to claim 25, characterized in that it contains other active principles.

15 27. The composition according to claim 26 characterized in that it contains as other active principles, at least one of the following products:

IR5885, a dipeptide compound corresponding to diastereoisomeric mixtures of methyl [S- (R, S)]- [3- (N- isopropoxycarbonylvalinyl) -amino] -3- (4-
chlorophenyl)-propanoate in any proportion, or to one of the two
20 diastereoisomeric forms S-R or S-S taken individually;

IR6141, corresponding to methyl N- (phenylacetyl) -N- 2,6-xylyl-R-alaninate;

salicylic acid (SA) or one of its derivatives selected from acetylsalicylic acid (ASA) , copper salts of salicylic acid (SA₂Cu) or (SACu) or acetylsalicylic acid (ASA₂Cu);

a copper (I) or copper (II) salt, selected from copper oxychloride, copper hydroxide, Bordeaux mixture, copper sulfate, or a mixture of copper hydroxide and oxychloride;

Fosetyl, corresponding to ethyl hydrogenphosphonate; and

- 5 Fosetyl -Al, corresponding to aluminum triethylphosphonate, commonly known with the trade-name of Aliette.

28. The composition according to claim 26 characterized in that it is selected from:

acid choline phosphite and IR5885;

- 10 acid choline phosphite and IR6141;

acid choline phosphite and Benthiavalicarb-isopropyl;

acid choline phosphite and ASA_2Cu ;

acid choline phosphite and SA_2Cu ;

acid choline phosphite and $SACu$; and

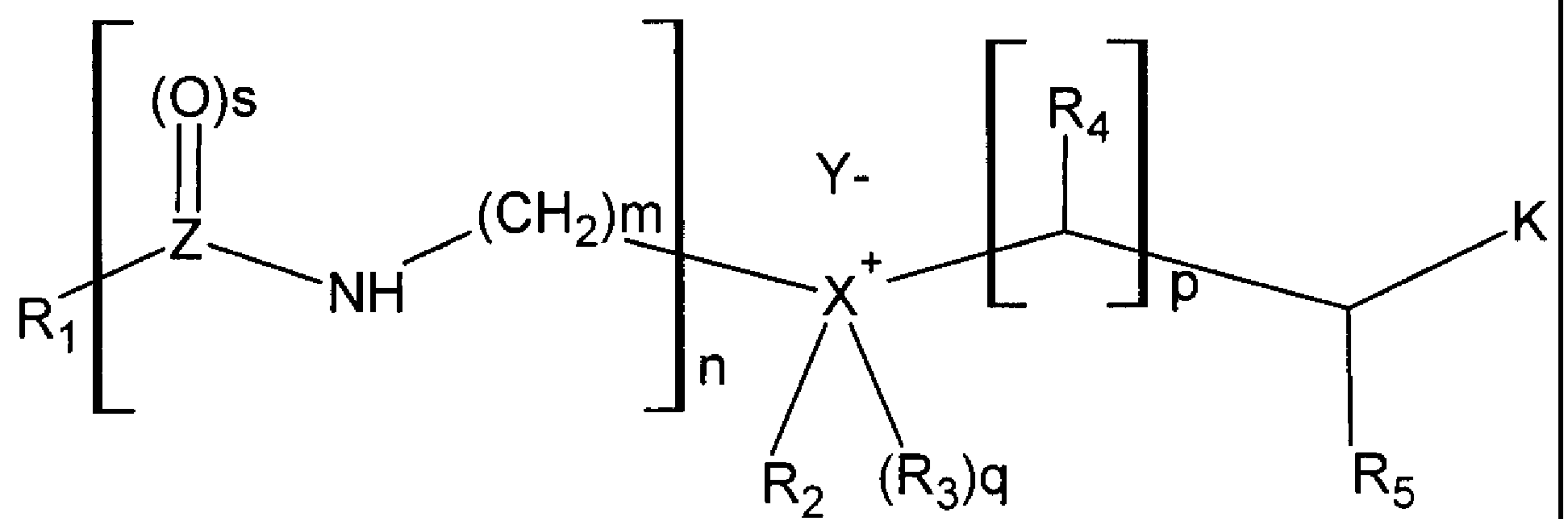
- 15 acid choline phosphite and iprovalicarb.

29. The composition according to any one of claims 25 to 28 wherein the concentration of active principle ranges from 1% to 90%.

30. Use of the composition according to any one of claims 25 to 29, for the control of phytopathogenic fungi and bacteria.

- 20 31. Use of the composition according to any one of claims 25 to 29, for the stimulation of the natural defense systems of plants from abiotic stress and biotic stress and the induction of resistance in the plants themselves.

32. Use according to claim 30 or 31, wherein the application of the composition is effected on all parts of the plant, on the leaves, stems, branches and roots, or on the seeds themselves before planting, or on the ground in which the plant grows.



(I)