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- (54) **COMPOUNDS HAVING REVERSIBLE INHIBITING ACTIVITY OF CARNITINE PALMITOYL-TRANSFERASE**
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- (58) **Field of Classification Search** 514/476,
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 548/252, 254
 See application file for complete search history.

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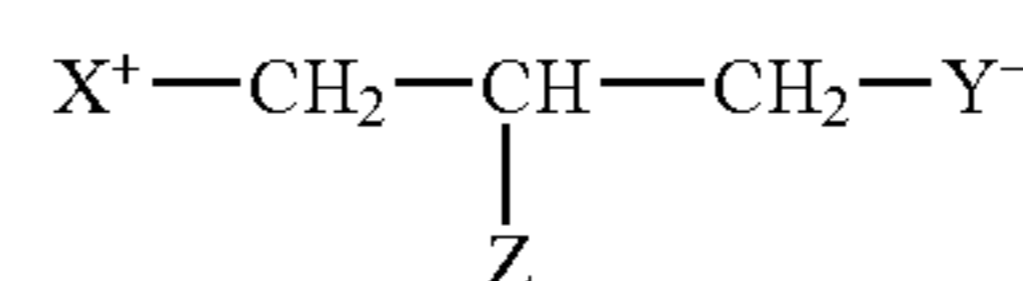
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Primary Examiner—Charanjit S Aulakh(74) *Attorney, Agent, or Firm*—Lucas & Mercanti, LLP**(57) ABSTRACT**

Compounds of formula (I)



wherein the groups are as defined in the description are
 disclosed.

The compounds of formula (I) are endowed with reversible
 inhibiting activity of carnitine palmitoyl-transferase and are
 useful in the preparation of medicaments useful in the
 pathologies related to a hyperactivity of carnitine
 polymitoyl-transferase, such as hyperglycemia, diabetes and
 pathologies related thereto, heart failure, ischemia.

13 Claims, 5 Drawing Sheets

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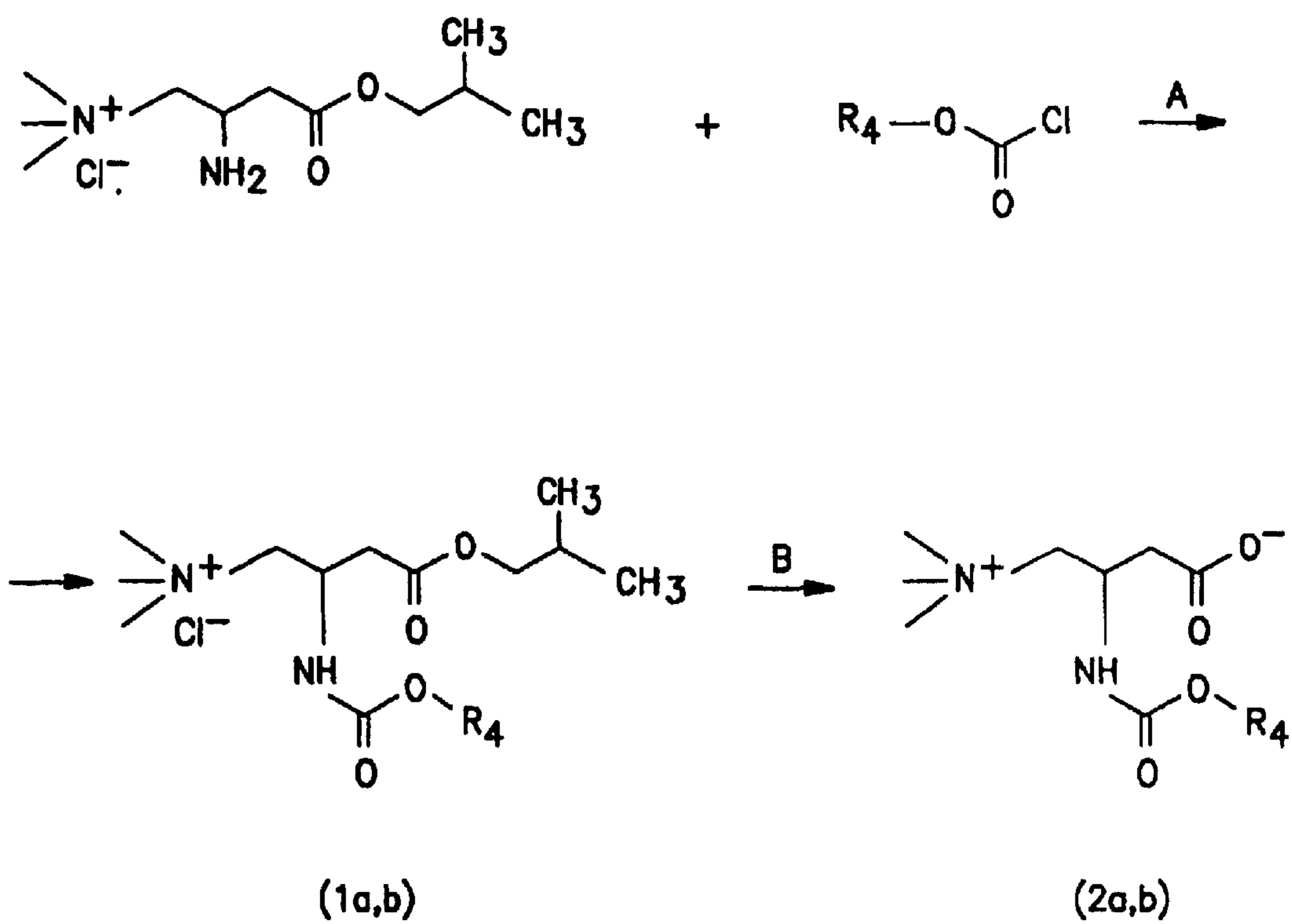
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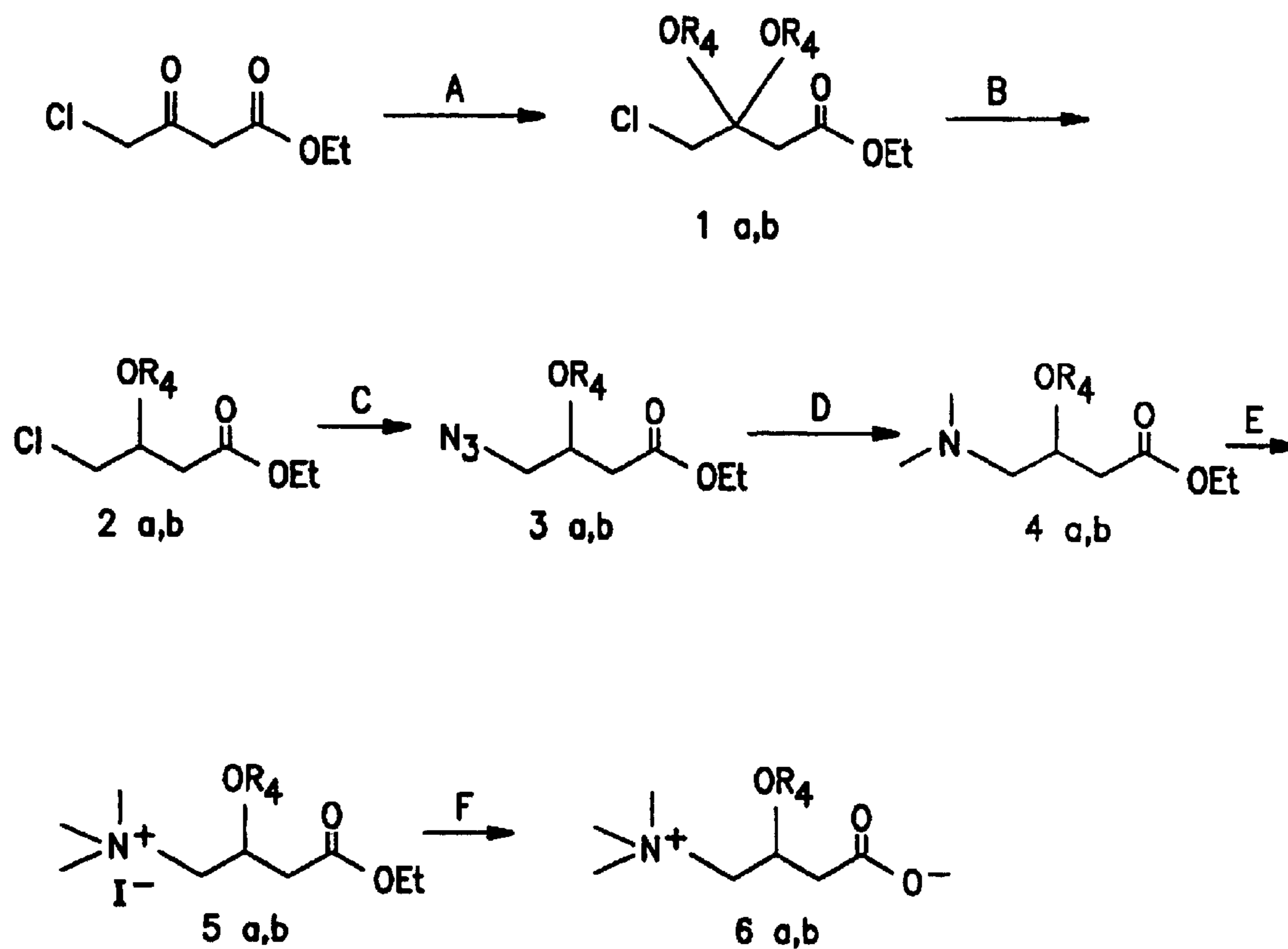
Fig. 1



$R_4 =$ a) $-(CH_2)_7CH_3$
 b) $-(CH_2)_8CH_3$

A) base
 B) IRA 402/ OH^- form

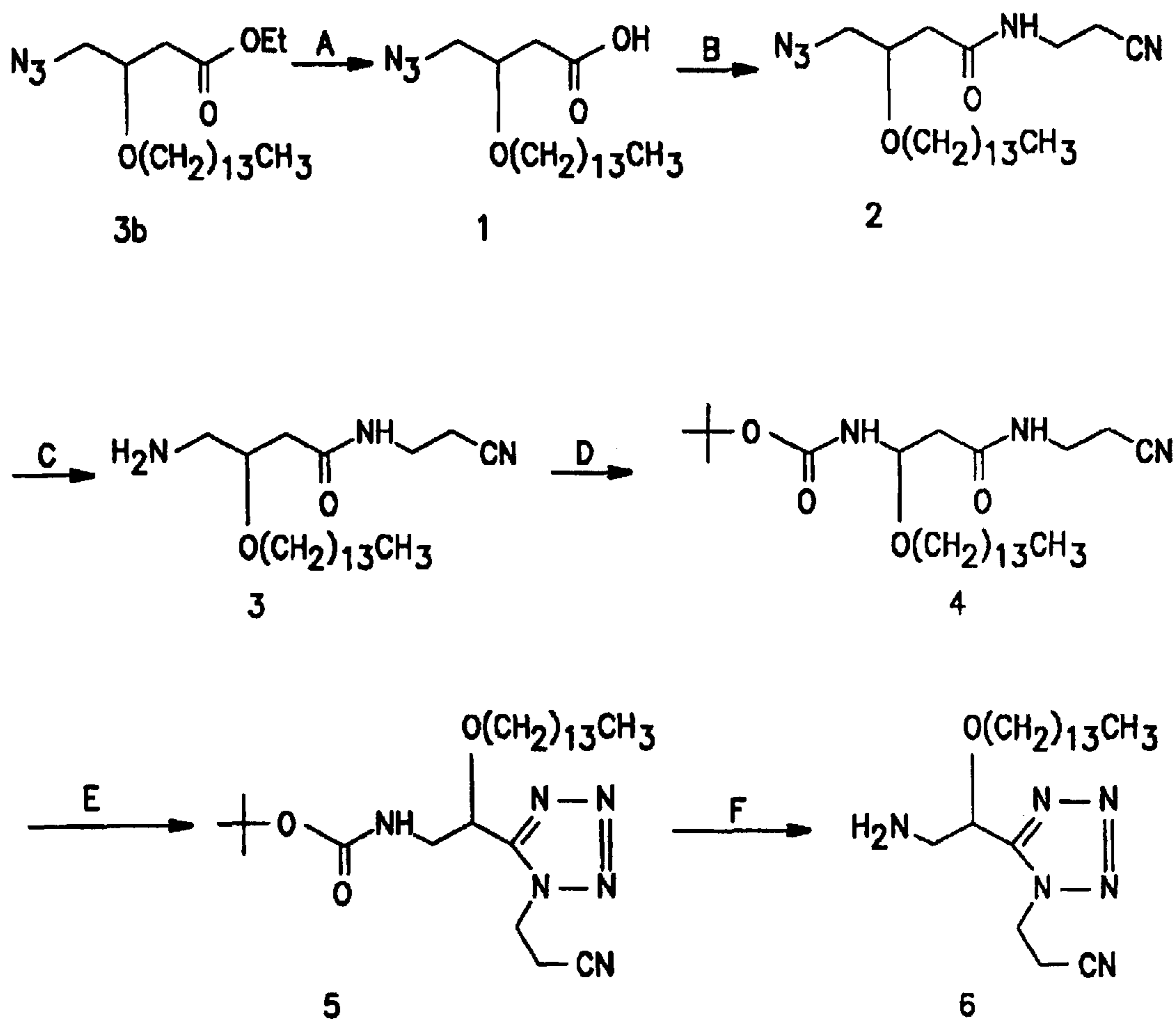
Fig. 2



$R_4 =$ a) $-(CH_2)_7CH_3$
 b) $-(CH_2)_{13}CH_3$

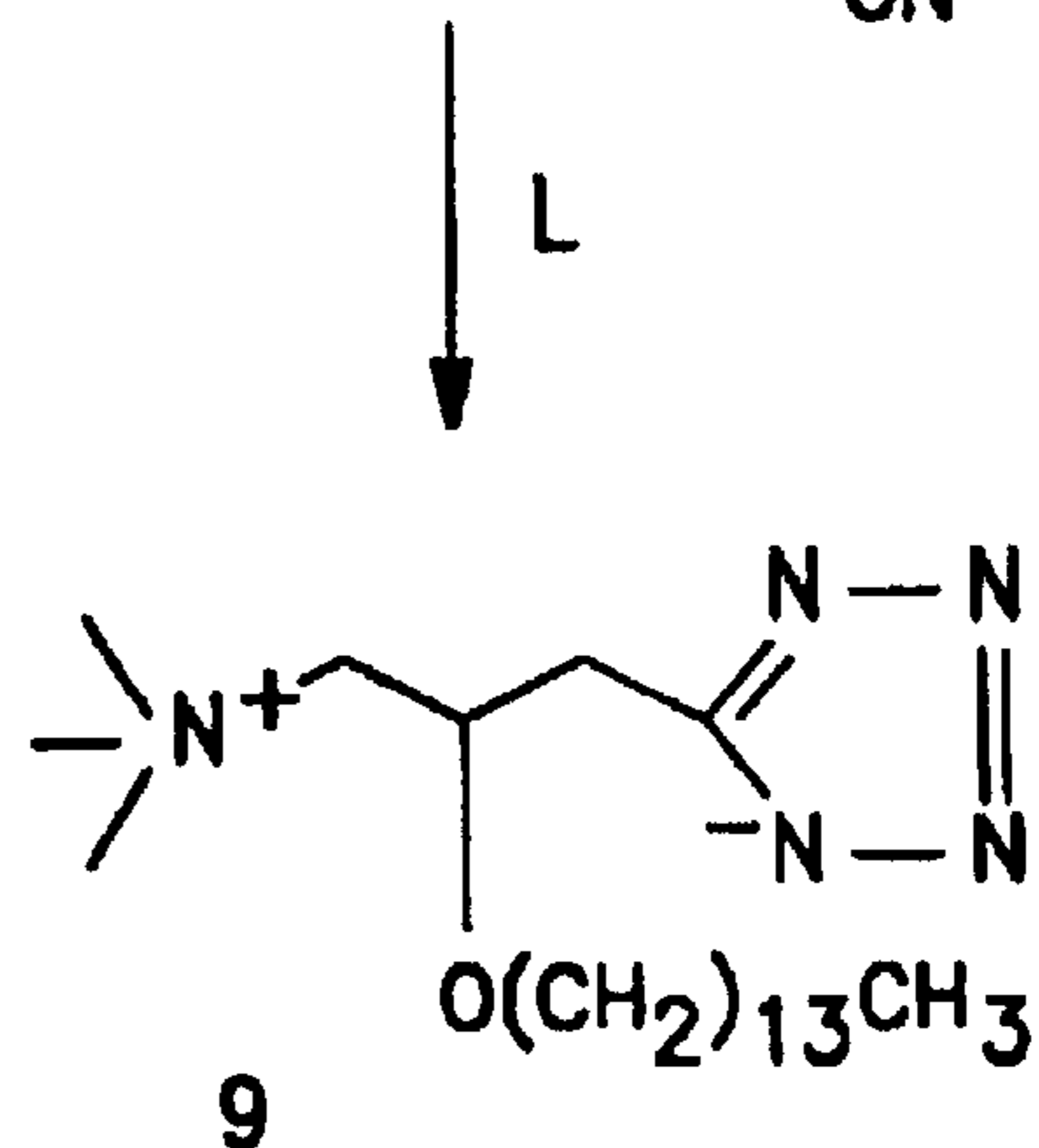
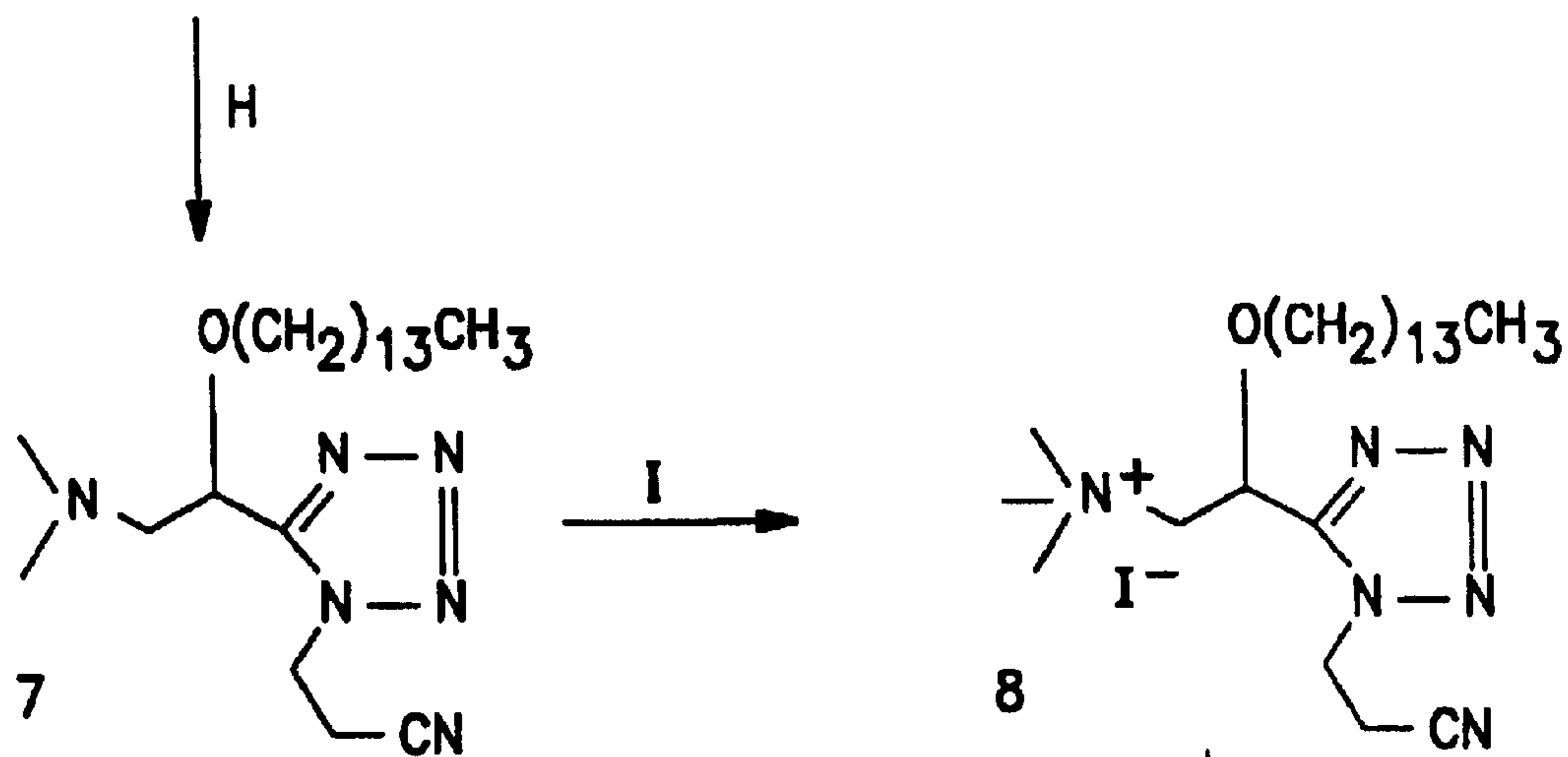
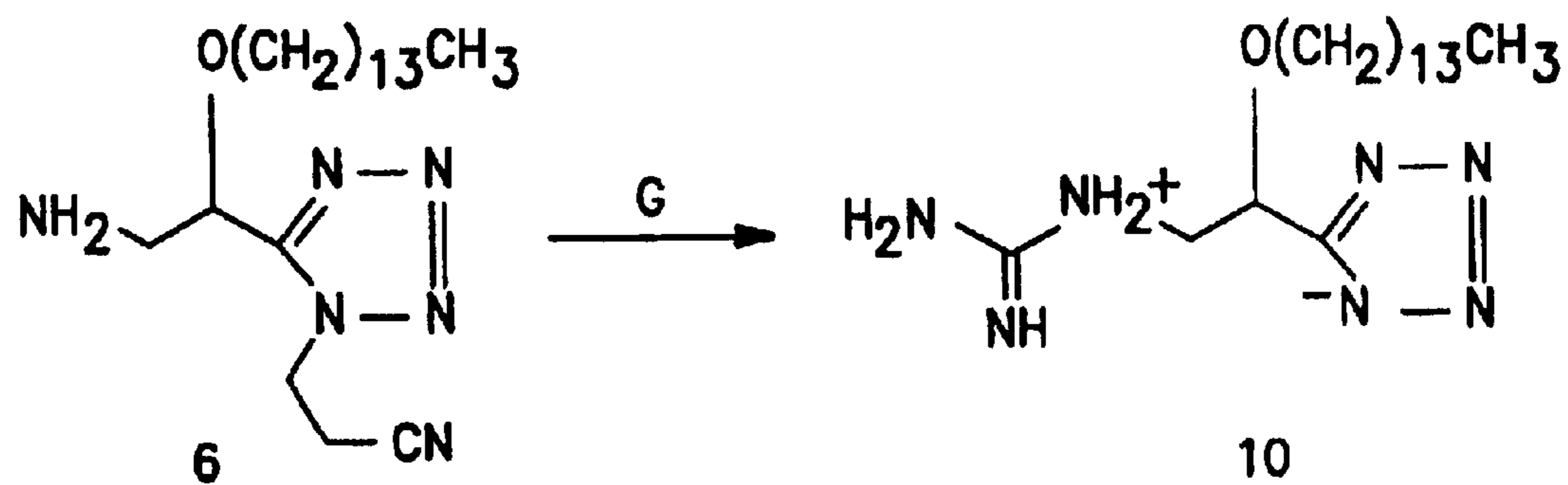
A) R_4OH , $SOCl_2$
 B) Et_3SiH , $BF_3 \cdot Et_2O$
 C) NaN_3
 D) Pd/C , H_2 , $HCHO$
 E) CH_3I
 F) IRA 402 OH^-

Fig. 3A



- A) NaOH 4N, MeOH, 16h, t.a
 B) H₂NCH₂H₂CN, DMF, TEA, DEPC
 C) THF, Ph₃P
 D) (BOC)₂O, NaOH IN
 E) THF, PH₃P, DEAD, Et₃SiN₃
 F) HCl 3N, NaOH IN

Fig. 3B

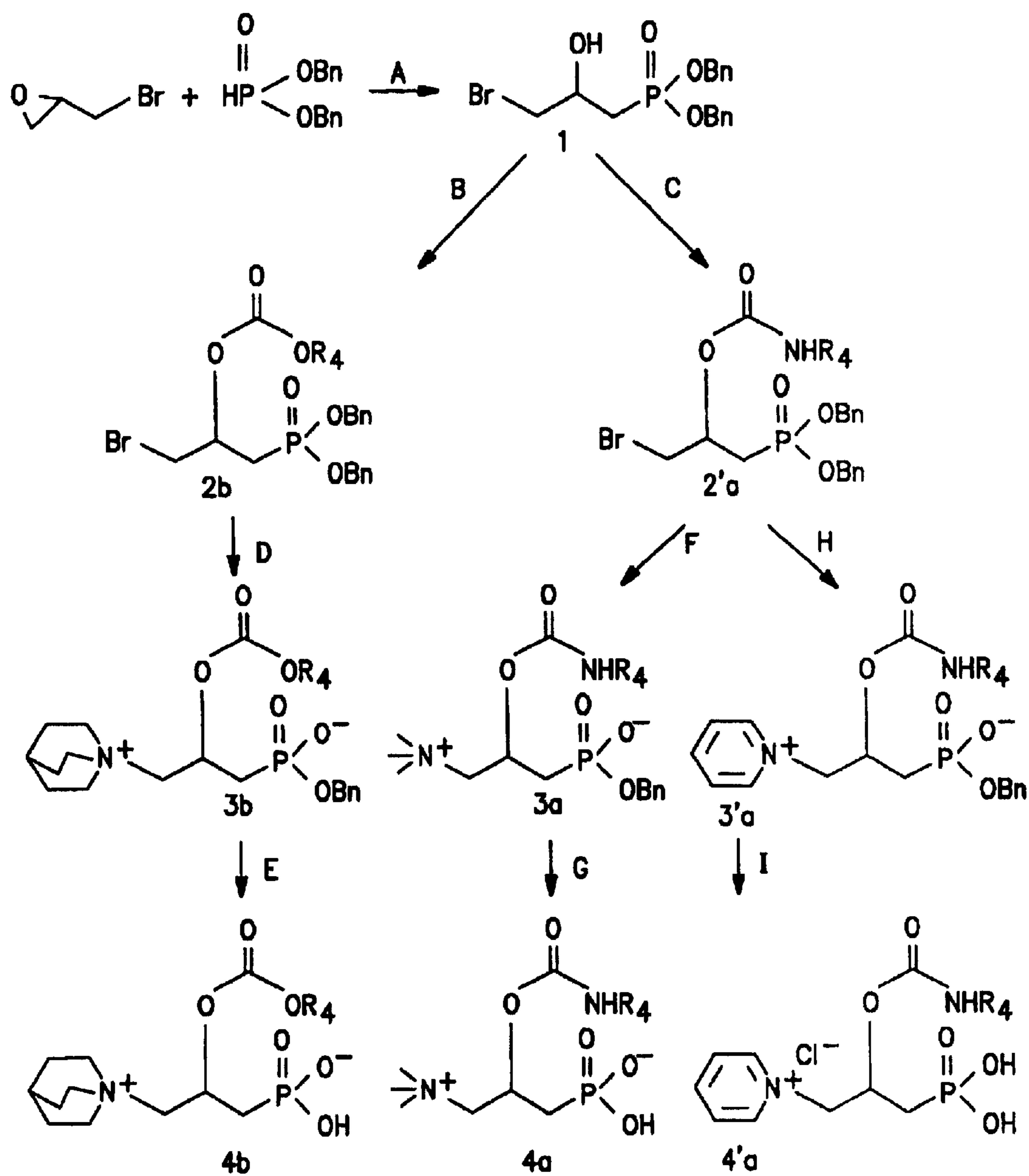


G) $\text{H}_2\text{N}-\text{C}(\text{NH})=\text{SO}_3\text{H}$, NaOH IN

H) HCOOH , H_2CO

I) CH_3I

L) IRA 402 attiv. OH^-



$R_4 =$ a) $-(\text{CH}_2)_7\text{CH}_3$
 b) $-(\text{CH}_2)_{13}\text{CH}_3$

A) 1) BuLi 2) $\text{BF}_3 \cdot \text{Et}_2\text{O}$
 B) $R_4\text{OCOC}$ l, Base
 C) $R_4\text{N}=\text{C}=\text{O}$, $\text{BF}_3 \cdot \text{Et}_2\text{O}$
 D) Quinuclidine F) Trimethylamine
 H) Pyridine
 E=G) $\text{H}_2, \text{Pd/C}$

Fig. 4

**COMPOUNDS HAVING REVERSIBLE
INHIBITING ACTIVITY OF CARNITINE
PALMITOYL-TRANSFERASE**

Matter enclosed in heavy brackets [] appears in the original patent but forms no part of this reissue specification; matter printed in italics indicates the additions made by reissue.

This application is a division of application Ser. No. 09/677,328 filed Oct. 2, 2000, now U.S. Pat. No. 6,369,073, the entire content of which is hereby incorporated by reference in this application.

The present invention relates to compounds having inhibiting activity against carnitine palmitoyl transferase. The present invention relates also to pharmaceutical compositions containing at least one of these compounds active ingredients and to the use of said compounds in the preparation of medicaments useful in the treatment of pathologies related to a hyperactivity of carnitine palmitoyl-transferase, in particular hyperglycaemic states, such as diabetes and related pathologies and of congestive heart failure.

BACKGROUND OF THE INVENTION

To date, hypoglycaemic therapy is based on the use of drugs having different mechanism of action (Arch. Intern. Med., 1977, 157, 1802–1817).

Insulin and its analogues represent the most used therapy, recurring to the direct hypoglycaemic action of this hormone.

Other compounds act indirectly by stimulating insulin release (sulphonylureas). A different target of hypoglycaemic drugs is represented by the reduction of glucose intestinal absorption through the inhibition of intestinal glucosidases, or by reducing insulin resistance.

Hyperglycaemia is also treated with gluconeogenesis inhibitors, such as biguanides.

Some works have also stressed out the relationship between gluconeogenesis and fatty acid oxidation.

The membrane bound long-chain acylcarnitine transferases, also known as carnitine palmitoyltransferase (CPT), are widely represented in organs and subcellular organelles (Bieber, L. L. 1988 Ann. Rev. Biochem. 57: 261–83). The well-established role of this category of enzymes is the transport of activated long-chain fatty acids through mitochondrial membranes. In this context, the outer mitochondrial membrane CPT I catalyzes the formation of long.-chain acylcarnitines that are transported across the mitochondrial membrane by a specific carrier, and reconverted into long-chain acyl-coenzyme A esters of CPT II, which resides in the inner mitochondrial membrane. Long-chain acyl-CoAs are then oxidised to acetyl-coenzyme A, which activates a key gluconeogenetic enzyme: pyruvate carboxylase.

Other works report that diabetic patients have high blood levels of fatty acids, whose liver oxidative fate gives rise to an increase of acetyl-coenzyme A, ATP and NADH. High availability of these compounds maximally stimulates gluconeogenesis, which is in part responsible of the elevated glucose blood levels in diabetic patients. CPT inhibition indirectly reduces the extent of liver gluconeogenesis, and hence blood glucose levels.

CPT inhibitors have been disclosed in J. Med. Chem., 1995, 38(18), 3448–50 and in the corresponding European patent application EP 0 574 355 as potential derivatives with hypoglycaemic activity.

Aminocarnitines N-acylated with —COR residue, wherein R is an aliphatic residue with 1 to 19 carbon atoms are disclosed in WO85/04396 useful for investigating the role of transferases in the body, in particular the specificity of carnitine acyltransferase.

Emeriamine and its analogues are disclosed in EP 0 127 098 and J. Med. Chem. 1987, 30, 1485–1463.

Notwithstanding the mechanism of activity above outlined, to date, drugs inhibiting CPT capable to effectively counteracts hyperglycaemia do not exist. For some products, such as tetradecyl glycidic acid, or etomoxir, myocardial hypertrophy have been evidenced as side effects (Life Sci., 1989, 44, 1897–1906).

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a flow chart illustrating examples 6 and 7;

FIG. 2 is a flow chart illustrating examples 8 and 9;

FIG. 3A is a flow chart illustrating examples 10 and 11;

FIG. 3B is a flow chart illustrating examples 10 and 11; and

FIG. 4 is a flow chart illustrating examples 12-14.

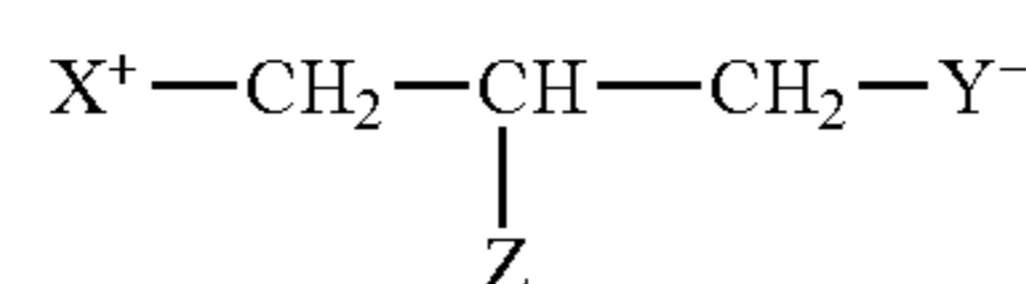
BRIEF DESCRIPTION OF THE INVENTION

None of the therapies presently used in clinic is fully satisfying, in particular due to the onset of unwanted side effects, such as severe hypoglycaemia, allergic phenomena, oedema, diarrhoea, intestinal disturbances, kidney toxicity, etc.

The necessity to obtain alternative effective therapies for hyperglycaemia still remains.

[ABSTRACT OF THE INVENTION]

It has now surprisingly been found that compounds of general formula (I):



wherein: X⁺ is selected from the group consisting of N⁺(R₁, R₂, R₃) and P⁺(R₁, R₂, R₃), wherein

(R₁, R₂, R₃), being the same or different, are selected in the group consisting of hydrogen and C₁–C₉ straight or branched alkyl groups, —CH=NH(NH₂), —NH₂, —OH; or two or more R₁, R₂ and R₃ together with the nitrogen atom, which they are linked to, form a saturated or unsaturated, monocyclic or bicyclic heterocyclic system; with the proviso that at least one of the R₁, R₂ and R₃ is different from hydrogen;

Z is selected from

—OR₄,

—OCOOR₄,

—OCONHR₄,

—OCSNHR₄,

—OCSOR₄,

—NHR₄,

—NHCOR₄,

—NHCSR₄,

—NHCOOR₄,

—NHCSOR₄,

—NHCONHR₄,

—NHCSNHR₄,

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—NHSOR₄, p1 —NHSONHR₄,
 —NHSO₂R₄,
 —NHSO₂NHR₄,
 —SR₄,

wherein —R₄ is a C₁–C₂₀ saturated or unsaturated, straight or branched alkyl group, optionally substituted with a A₁ group, wherein A₁ is selected from the group consisting of halogen atom, C₆–C₁₄ aryl, heteroaryl, aryloxy or heteroaryloxy group, said aryl, heteroaryl, aryloxy or heteroaryloxy groups being optionally substituted with one or more C₁–C₂₀ saturated or unsaturated, straight or branched alkyl or alkoxy group and/or halogen atom;

Y⁻ is selected from the group consisting of —COO⁻, PO₃H⁻, —OPO₃H⁻, tetrazolate-5-yl;

with the proviso that when Z is —NHCOR₄, X⁺ is trimethylammonium, Y is —COO⁻, then R₄ is C₂₀ alkyl;

with the proviso that when Z is —NHSO₂R₄, X⁺ is trimethylammonium and Y⁻ is —COO⁻, then R₄ is not tolyl;

with the proviso that when Z is —NHR₄, X⁺ is trimethylammonium and Y⁻ is —COO⁻, then R₄ is not C₁–C₆ alkyl.

The present invention further comprises the use of the compounds of the above-mentioned formula (I) as active ingredients for medicaments, in particular for medicaments useful for the treatment of pathologies related to a hyperactivity of carnitine palmitoyl carnitine, such as and in particular hyperglycemic states, diabetes and related pathologies, congestive heart failure and dilatative cardiopathy.

The present invention comprises pharmaceutical compositions containing compounds of formula (I) as active ingredients, in admixture with pharmaceutically acceptable vehicles and excipients.

The present invention comprises also processes for the preparation of compounds of formula (I).

[ABSTRACT OF THE INVENTION]

Within the scope of the present invention, as examples of C₁–C₂₀ linear or branched alkyl group, methyl, ethyl, propyl, butyl, pentyl, hexyl, heptyl, octyl, nonyl, decyl, undecyl, dodecyl, tridecyl, tetradecyl, pentadecyl, hexadecyl, heptadecyl, octadecyl, nonadecyl and eicosyl and their possible isomers are meant, such as for example isopropyl, isobutyl, tert-butyl.

Examples of C₁–C₂₀ linear or branched alkenyl group are methylene, ethylidene, vinyl, allyl, propargyl, butylene, pentylene, wherein the carbon—carbon double bond, optionally in the presence of other carbon—carbon unsaturations, can be situated in the different possible positions of the alkyl chain, which can also be branched within the allowed isometry.

Examples of (C₆–C₁₄) aryl group are phenyl, 1- or 2-naphthyl, anthryl, optionally substituted as shown in the general definitions above-mentioned.

Examples of heterocyclic groups thienyl, quinolyl, pyridyl, N-methylpiperidyl, 5-tetrazolyl, optionally substituted as shown in the general definitions above-mentioned.

As halogen atom it is intended fluorine, chlorine, bromine, iodine.

The compounds of formula (I) can be also in the form of inner salts.

A first group of preferred compounds comprises the compounds of formula (I) wherein N⁺(R₁,R₂,R₃) is trimethyl ammonium.

A second group of preferred compounds comprises the compounds of formula (I) wherein two or more R₁, R₂ and R₃, together with the nitrogen atom, which they are linked

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to, form a saturated or unsaturated, monocyclic or bicyclic heterocyclic system; for example morpholinium, pyridinium, pyrrolidinium, quinolinium, quinuclidinium.

A third group of preferred compounds comprises the compounds of formula (I) wherein R₁ and R₂ are hydrogen and R₃ is selected from the group consisting of —C=NH(NH₂), —NH₂ and —OH.

Within the different embodiments of the present invention, the R₄ group is preferably a C₇–C₂₀ saturated or unsaturated, straight or branched alkyl group. In fact, it has been observed the length of the alkyl chain R₄ significantly increases the selectivity against CPT. Preferred R₄ groups are selected from the group consisting of heptyl, octyl, nonyl, decyl, undecyl, dodecyl, tridecyl, tetradecyl, pentadecyl, hexadecyl, heptadecyl, octadecyl, nonadecyl and eicosyl.

Preferred examples of Z group are ureido (—NHCONHR₄), and carbamate (—NHCOOR₄, —OCONHR₄) ones.

In particular, compounds of formula (I) wherein X⁺, R₁, R₂, R₃, have the above disclosed meanings, Z is ureido (—NHCONHR₄) or carbamate (—NHCOOR₄, —OCONHR₄), R₄ is C₇–C₂₀, preferably a C₉–C₁₈ saturated or unsaturated, straight or branched alkyl group, are preferred.

The compounds of formula (I) have an asymmetry center on carbon atom bound to a Z group. For the purposes of the present invention, each compound of formula (I) can exist both as R,S racemic mixture and as separated R/S isomeric form.

The compounds of formula (I) are quaternary ammonium or phosphonium derivatives always containing a Y⁻ anionic group. Dependently on pH, each compounds of formula (I) can exist indifferently as amphoteric (inner salt) or as a compound wherein Y⁻ is preferably in the YH form. In such a case, X⁺ is salified with a pharmacologically acceptable acid. Formula (I) covers all these different possibilities. In case of nitrogen atoms having basic character, the salts with pharmaceutically acceptable acids, both inorganic and organic, such as for example, hydrochloric acid, sulfuric acid, acetic acid, or, in the case of acid group, such as carboxyl, the salts with pharmaceutically acceptable bases, both inorganic and organic, such as for example, alkaline and alkaline-earth hydroxides, ammonium hydroxide, amine, also heterocyclic ones. Examples of pharmaceutically acceptable salts are chloride; bromide; iodide; aspartate; acid aspartate; citrate; acid citrate; tartrate; acid tartrate; phosphate, acid phosphate; fumarate; acid fumarate; glycerophosphate; glucosephosphate; lactate; maleate; acid maleate; mucate; orotate; oxalate; acid oxalate; sulfate; acid sulfate; trichloroacetate; trifluoroacetate; methanesulfonate; pamoate and acid pamoate.

A first group of particularly preferred compounds comprises:

R,S-4-trimethylammonium-3-(nonylcarbamoyl)-aminobutyrate;

R,S-4-quinuclidinium-3-(tetradecyloxycarbonyl)-oxybutyrate;

R,S-4-trimethylammonium-3-(nonylcarbamoyl)-oxybutyrate;

R,S-4-trimethylammonium-3-(nonyloxycarbonyl)-oxybutyric acid chloride;

R,S-4-trimethylphosphonium-3-(nonylcarbamoyl)-oxybutyrate;

R,S-4-trimethylammonium-3-(octyloxycarbonyl)-aminobutyrate;

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R,S-4-trimethylammonium-3-(nonyloxycarbonyl)-aminobutyrate;
 R,S-4-trimethylammonium-3-octyloxybutyrate;
 R,S-4-trimethylammonium-3-tetradecyloxybutyrate;
 R,S-1-guanidinium-2-tetradecyloxy-3-(tetrazolate-5-yl) propane;
 R,S-1-trimethylammonium-2-tetradecyloxy-3-(tetrazolate-5-yl)-propane;
 R,S-3-quinuclidinium-2-(tetradecyloxycarbonyl)-oxy-1-propanephosphonate monobasic;
 R,S-3-trimethylammonium-2-(nonylaminocarbonyl)-oxy-1-propanephosphonate monobasic;
 R,S-3-pyridinium-2-(nonylaminocarbonyl)-oxy-1-propanephosphonic acid chloride;
 R-4-trimethylammonium-3-(tetradecylcarbamoyle)-aminobutyrate;
 R-4-trimethylammonium-3-(undecylcarbamoyle)-aminobutyrate;
 R-4-trimethylammonium-3-(heptylcarbamoyle)-aminobutyrate;
 R,S-4-trimethylammonium-3-(nonylthiocarbamoyle)-aminobutyrate;
 R-4-trimethylammonium-3-(nonylcarbamoyle)-aminobutyrate;
 S-4-trimethylammonium-3-(nonylcarbamoyle)-aminobutyrate;
 S-4-trimethylammonium-3-(tetradecylcarbamoyle)-aminobutyrate;
 R,S-4-trimethylammonium-3-tetradecylaminobutyrate;
 R,S-4-trimethylammonium-3-octylaminobutyrate;
 R,S-4-trimethylammonium-3-(decansulfonyl)-aminobutyrate;
 R,S-4-trimethylammonium-3-(nonylsulfamoyle)-aminobutyrate;
 S-4-trimethylammonium-3-(dodecanesulfonyl)-aminobutyrate;
 R-4-trimethylammonium-3-(dodecansulfonyl)-aminobutyrate;
 S-4-trimethylammonium-3-(undecylsulfamoyle)-aminobutyrate;
 R-4-trimethylammonium-3-(undecylsulfamoyle)-aminobutyrate;
 R-4-trimethylammonium-3-(dodecylcarbamoyle)-aminobutyrate;
 R-4-trimethylammonium-3-(10-phenoxydecylcarbamoyle)-aminobutyrate; R-4-trimethylammonium-3-(trans- β -styrenesulfonyl) aminobutyrate.

The compounds of formula (I) can be prepared with reactions that are well known in the state of the art.

A process for the preparation of the compounds of claim 1, wherein Z is $-\text{NHR}_4$ comprising the reaction of $\text{X}^+-\text{CH}_2-\text{CH}(\text{NH}_2)-\text{CH}_2-\text{Y}^-$, wherein X^+ and Y^- have the same meanings as in claim 1, of the desired structure, optionally protected on the acid Y^- group, with alkane carbaldehydes, wherein the alkyl moiety is a one-term lower homologue of the desired R_4 and subsequent reduction.

Generally, the compounds of formula (I), wherein Z is carbonate ($-\text{OCOOR}_4$), carbamate ($-\text{OCONHR}_4$, $-\text{NHCOOR}_4$), thiocarbamate ($-\text{OCSNHR}_4$, $-\text{NHCSOR}_4$) or thiocarbonate ($-\text{OCSOR}_4$), are obtained by reacting a compound of formula $\text{X}^+-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{Y}^-$, wherein X^+ and Y^- are as above defined, of the desired structure, optionally protected on the acid Y^- group, respectively with alkyl chloroformates, alkyl isocyanates, alkyl isothiocyanates, alkyl thiochloroformates, containing the desired R_4 alkyl group.

Compounds of formula (I), wherein Z is amide ($-\text{NHCOR}_4$), thioamide ($-\text{NHCSR}_4$), carbamate

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($-\text{NHCOOR}_4$, $-\text{OCONHR}_4$), thiocarbamate ($-\text{NHCSOR}_4$, $-\text{OCSNHR}_4$), ureido ($-\text{NHCONHR}_4$), thioureido ($-\text{NHCSNHR}_4$), sulfinamide ($-\text{NHSOR}_4$), sulfonamide ($-\text{NHSO}_2\text{R}_4$), sulfinamoylamino ($-\text{NHSONHR}_4$), and sulfamide ($-\text{NHSO}_2\text{NHR}_4$), are obtained by reacting $\text{X}^+-\text{CH}_2-\text{CH}(\text{NH}_2)-\text{CH}_2-\text{Y}^-$, wherein X^+ and Y^- are as above defined, of the desired structure, optionally protected on the acid Y^- group, respectively with acetyl chlorides, thioacyl chlorides, alkyl chloroformates, alkyl thiochloroformates, alkyl isocyanates, alkyl thioisocyanates, alkyl sulfinyl chlorides, alkyl sulfonyl chlorides, SOCl_2 and alkyl amines, alkyl sulfamoyl chlorides (or SO_2Cl_2 and alkyl amines), containing the desired R_4 alkyl group.

Compounds of formula (I), wherein Z is $-\text{OR}_4$ or $-\text{SR}_4$ are obtained by the reaction of carbonyl compounds of formula $\text{Hal}-\text{CH}_2-\text{CO}-\text{CH}_2-\text{COOR}'$, wherein Hal is a halogen atom, preferably chlorine, and R' is the residue of a suitable ester, such as for example a lower alkyl ester (an ethyl or a tert-butyl ester) with respectively alcohols and thiols R_4OH or R_4SH , wherein R_4 is as above defined, to give the respective ketal or thioketal, followed by the transformation of the respective ketal or thioketal into the respective ether to thioether, subsequent substitution of the Hal atom with a nucleophilic group, such as azido, phthalimido, nitro, amino, alkyl amino group, and transformation of the nucleophilic group into the X^+ group, wherein X^+ is $\text{N}^+(\text{R}_1, \text{R}_2, \text{R}_3)$ or, alternatively the Hal atom is substituted with a $(\text{R}_1, \text{R}_2, \text{R}_3)$ -substituted phosphine to obtain the compounds of formula (I) wherein X^+ is $\text{P}^+(\text{R}_1, \text{R}_2, \text{R}_3)$.

Compounds of formula (I), wherein Z is $-\text{NHR}_4$ are obtained by reacting $\text{X}^+-\text{CH}_2-\text{CH}(\text{NH}_2)-\text{CH}_2-\text{Y}^-$, wherein X^+ and Y^- have the same meanings as in claim 1, of the desired structure, optionally protected on the acid Y^- group, with alkane carbaldehydes, wherein the alkyl moiety is a one-term lower homologue of R_4 , and subsequent reduction.

Regarding the various meanings of R_4 , present in the different reactives, these reactives are available in the market, or can be prepared according to well-known methods in literature, which the experts in the field can resort to, completing with their own knowledge of the argument.

Pharmaceutically acceptable salts are obtained with conventional methods found in the literature, and do not necessitate of further disclosure.

The compounds disclosed in the present invention have reversible inhibiting activity of carnitine palmitoyl-transferase (CPT). This activity allows their use as active ingredients in the preparation of medicaments useful for the treatment and prevention of hyperglycaemia, diabetes and disorders related thereto, such as, for example diabetic retinopathy, diabetic neuropathy. The compounds of the present invention are also useful as active ingredient for the treatment and prevention of cardiovascular disorders, such as congestive heart failure. The compounds of formula (I) are also applicable for medicaments for the prevention and treatment of ketonic states, wherein it is intended the pathological conditions characterized by high levels of ketone bodies in blood.

Inhibiting activity mainly occurs on the isoform I of palmitoyl carnitine transferase (CPT-I).

A further object of the present invention relates to pharmaceutical compositions comprising at least a compound of formula (I), in an amount such as to produce a significant therapeutical effect. The compositions according to the present invention are conventional and are obtained with commonly used methods in the pharmaceutical industry.

According to the desired administration route, the compositions shall be in solid or liquid form, suitable to the oral, parenteral, intravenous or transdermal route. The compositions according to the present invention comprise together with the active ingredients at least a pharmaceutically acceptable vehicle or excipient. Formulation co-adjuvants, for example solubilizing, dispersing, suspending, emulsifying agents can be particularly useful. Examples of suitable oral pharmaceutical compositions are capsules, tablets, granulates, powders, syrups, elixirs. Examples of suitable parenteral pharmaceutical compositions are solutions, emulsions, suspensions. Examples of suitable transdermal pharmaceutical compositions are patches, subcutaneous implants.

The compounds of formula (I) can also be used in combination with other well-known active ingredients.

The dose of the active ingredients will vary depending on the kind of active ingredient used, the administration route, the grade of pathology to be treated and the general conditions of the subject. The dosage and posology shall be determined by the clinic expert or the physician. Generally, a therapeutic effect can be obtained at dosages comprised between 1 and 100 mg/kg body weight.

The compounds according to the present invention are useful as medicaments with hypoglycaemic activity. A further object of the present invention is the preparation of a pharmaceutical composition comprising admixing at least a compound of formula (I) with suitable pharmaceutically acceptable excipients and/or vehicles.

The following examples further illustrate the invention.

EXAMPLE 1

R,S-4-trimethylammonium-3-(nonylcarbamoyl)aminobutyrate (ST 1251)

Nonyl Isocyanate

A solution of decanoyl chloride (20 g, 104.8 mmoles) in acetone (30 ml) was dropped into a solution of sodium azide (9.53 g, 146.6 mmoles) in water (30 ml), cooled in an ice bath. The temperature of the azide solution was kept between 10 and 15° C. after one hour, the solution was transferred in a separatory funnel and the lower phase (the aqueous one) was eliminated. The higher phase was transferred into a flask containing 100 ml of toluene, previously warmed at 65° C. After 1.5 hours, the solution was evaporated to dryness, giving 13.37 g crude product, which after vacuum distillation gave 8.3 g pure product in the form of colorless liquid.

Yield 47%.

¹H-NMR (300 MHz; CDCl₃):

δ: 3.3 (t, 2H), 1.6 (m, 2H), 1.45-1.2 (m, 12H), 0.9 (brt, 3H).

R,S-4-trimethylammonium-3-(nonylcarbamoyl)aminobutyrate

Nonyl isocyanate (15.42 g, 91.12 mmoles) was added to a solution of aminocarnitine, inner salt (7.3 g, 45.56 mmoles) in anhydrous DMSO (350 ml) and the solution was left to stand for 60 hours at 40° C. The resulting mixture was transferred in a 3 l Erlenmeyer flask, containing ethyl ether (2.5 l) and the solvent was separated by decanting the formed precipitate, which was then transferred into a flask and precipitated again with ethyl ether. The so obtained crude product was washed several times with ethyl ether and purified on a silica gel chromatographic column, using a CHCl₃:

MeOH 9:1 to CHCl₃: MeOH 3:7 gradient until elution of impurities with higher R_f, then eluting the product of interest with MeOH only. 9.7 g of pure product were obtained.

Yield 68%.

M.p.: 145–147° C.

¹H-NMR (300 MHz; D₂O):

δ: 4.4 (m, 1H), 3.45 (dd, 1H), 3.30 (d, 1H), 3.05 (s, 9H), 2.9 (t, 2H), 2.3 (d, 2H), 1.3 (m, 2H), 1.15 (brs, 12H), 0.8 (brt, 3H).

FAB Mass=330, [(M+H)⁺].

Elemental analysis: responding to the expected formula C₁₇H₃₅N₃O₃.

K.F.=2.5% water.

TLC silica gel CHCl₃:iPrOH:MeOH:H₂O:CH₃COOH 42:7:28:10.5:10.5;

R_f=0.55.

HPLC: SGE-SCX column (5 μm, 250×4 mm), T=30° C., mobile phase 0.2 M KH₂PO₄:CH₃CN 85:15, pH as such, flow 0.75 ml/min, detector: RI, UV 205 nm, RT=12.63 min.

EXAMPLE 2

R,S-4-quinuclidinium-3-(tetradecyloxycarbonyl)oxybutyrate (ST 1265)

ter-Butyl R,S-4-quinuclidinium-3-hydroxybutyrate iodide

Quinuclidine (2.40 g, 21.60 mmoles) was added to ter-Butyl R,S-4-iodo-3-hydroxybutyrate (6.18 g, 21.60 mmoles) in acetonitrile (60 ml) and the solution was warmed to 60° C. for 20 hours under stirring. After evaporation of the solvent, the residue was dissolved in acetonitrile and precipitated with ethyl ether several times to give 7.2 g of product, contaminated with about 13% by weight of quinuclidine iodide (as from NMR). After repeated crystallization from CH₃CN/Et₂O, 4.3 g of pure product were obtained.

Yield 50%.

M.p.: 124–127° C.

¹H-NMR (300 MHz; D₂O):

δ: 4.50 (m, 1H), 3.40 (m, 2H), 2.42 (m, 2H), 2.08 (m, 1H), 1.88 (m, 6H), 1.34 (m, 9H).

FAB Mass=270, [M⁺].

Elemental analysis: responding to the expected formula C₁₅H₂₈INO₃.

K.F.=0.5% water.

The preparation of ter-butyl 4-iodo-3-hydroxybutyrate was carried out as described in J. Pharm. Science 64/7, 1262–1264, 1975.

Tetradecyl Chloroformate

29 ml of a 20% toluene solution of phosgene (55.98 mmoles) was added to tetradecyl alcohol (4 g, 18.66 mmoles) and the reaction mixture was left to stand for 20 hours under stirring at room temperature. After solvent evaporation, the residue was taken up with hexane and evaporated to dryness (several times) to give 5.1 g product as colorless liquid.

Yield 98%.

¹H-NMR (300 MHz, CDCl₃):

δ: 4.30 (t, 2H), 1.72 (m, 2H), 1.30 (m, 22H), 0.85 (brt, 3H).

ter-butyl R,S-4-guinaclidinium-3-(tetradecyloxycarbonyl)-oxy butyrate chloride

Dimethylaminopyridine (922 mg, 7.55 mmoles) and tetradecyl chloroformate (2.09 g, 7.55 mmoles) were added to ter-butyl R,S-4-guinaclidinium-3-hydroxybutyrate (2 g, 5.03 mmoles) in anhydrous CH_2Cl_2 (20 ml). The solution was left to stand at room temperature for 20 hours under stirring. After this time, the solution was diluted with CHCl_3 saturated with NaCl, and dried over anhydrous sodium sulfate. The dry residue obtained after evaporation was taken up with ethyl ether and the undissolved residue was filtered off. After solvent evaporation a crude product was obtained. Flash-chromatography (CHCl_3 : MeOH 9:1) and elution with MeOH on Amberlyst A-21 resin (activated in HCl from), gave 1.6 g product as chloride.

Yield 58%.

M.p.: 59–60° C.

$^1\text{H-NMR}$ (300 MHz, CDCl_3):

δ : 5.50 (m, 1H), 4.55 (d, 2H), 3.80 (m, 7H), 2.90 (dd, 1H), 2.75 (dd, 1H), 2.22 (m, 1H), 2.05 (d, 6H), 1.65 (m, 2H), 1.41 (s, 9H), 1.25 (m, 22H), 0.85 (brt, 3H).

FAB Mass=510, $[\text{M}^+]$.

Elemental analysis: responding to the expected formula $\text{C}_{30}\text{H}_{56}\text{ClNO}_5$.

K.F.=1.5% water.

R,S-4-guinaclidinium-3-(tetradecyloxycarbonyl)oxybutyrate

Trifluoroacetic acid (6 ml) was added to ter-butyl R,S-4-guinaclidinium-3-(tetradecyloxycarbonyl)-oxybutyrate chloride (1.05 g, 1.92 mmoles) and the solution was left to stand for 1 hour at room temperature under stirring. After vacuum-evaporation of trifluoroacetic acid, the residue was taken up with cyclohexane and evaporated to dryness several times, then transferred on an Amberlyst IRA 402 resin (Cl^- form) and eluted with water. The crude product, obtained by freeze-drying was purified through silica gel flash-chromatography (CHCl_3 : MeOH 8:2) giving 480 mg product as inner salt.

Yield 55%.

M.p.: 132–134° C.

$^1\text{H-NMR}$ (300 MHz; D_2O):

δ : 5.35 (m, 1H), 4.05 (m, 2H), 3.40 (m, 8H), 2.55 (dd, 1H), 2.35 (dd, 1H), 2.08 (m, 1H), 1.90 (m, 6H), 1.55 (m, 2H), 1.20 (m, 22H), 0.75 (brt, 3H).

FAB Mass=454, $[(\text{M}+\text{H})^+]$.

Elemental analysis: responding to the expected formula $\text{C}_{26}\text{H}_{47}\text{NO}_5$

K.F.=1.5% water.

TLC silica gel CHCl_3 :MeOH 7:3.

Rf=0.34.

HPLC: SGE-SCX column (5 μm , 250 \times 4 mm), T=30° C., mobile phase 0.05 M $(\text{NH}_4)\text{H}_2\text{PO}_4$: CH_3CN 60:40, pH 4.0, flow 0.75 ml/min, detector: RI, UV 205 nm, RT=6.72 min.

EXAMPLE 3

R,S-4-trimethylammonium-3-(nonylcarbamoyl)-oxybutyrate (ST 1298)

Benzyl ester of R,S-4-trimethylammonium-3-(nonylcarbamoyl)-oxybutyric acid Perchlorate

Nonyl isocyanate (7.39 g, 43.36 mmoles) was added to a solution of R,S-carnitine perchlorate, benzyl ester (7.69 g,

21.86 mmoles) in toluene (100 ml) and the solution was refluxed for 5 days under stirring. Nonyl isocyanate (1.84 g, 10.86 mmoles) was further added and the reaction mixture was left under reflux for other 5 days. The solvent was vacuum-evaporated and the residue was washed with ethyl ether and subsequently taken up with chloroform, washed with water and dried over anhydrous sodium sulfate. The oil resulting from the evaporation of the organic phase was purified through flash-chromatography column, using a gradient CHCl_3 to CHCl_3 : MeOH 95:5 4.4 g product were obtained in the form of a thick oil.

Yield 38.6%.

$^1\text{H-NMR}$ (200 MHz; CDCl_3):

δ : 7.3 (s, 5H) 5.4 (m, 2H), 5.05 (m, 2H), 3.8 (dd, 1H), 3.55 (d, 1H), 3.15 (s, 9H), 3.05 (m, 2H), 2.75 (m, 2H), 1.4 (m, 2H), 1.2 (brs, 12H), 0.8 (brt, 3H).

TLC silica gel CHCl_3 : MeOH 9:1;

Rf=0.29.

R,S-4-trimethylammonium-3-(nonylcarbamoyl)-oxybutyrate

10% Pd/C (0.44 g) was added to benzyl ester of R,S-4-trimethylammonium-3-(nonylcarbamoyl)-oxybutyric acid perchlorate (4.4 g, 8.44 mmoles) in MeOH (115 ml) and the mixture was hydrogenated at 47 psi for 4 hours. After filtration on celite, the solution was vacuum-concentrated and passed through an Amberlyst A-21 resin, eluting with MeOH. After solvent evaporation, 2.47 g product were obtained.

Yield 88.7%.

M.p.: 151–153° C.

$^1\text{H-NMR}$ (300 MHz; D_2O):

δ : 5.4 (m, 1H), 3.75 (dd, 1H), 3.5 (d, 1H), 3.15 (s, 9H), 3.05 (t, 2H), 2.55 (dd, 1H), 2.40 (dd, 1H), 1.45 (m, 2H), 1.20 (brs, 12H), 0.8 (brt, 3H).

FAB Mass=331, $[(\text{M}+\text{H})^+]$.

Elemental analysis: responding to the expected formula $\text{C}_{17}\text{H}_{34}\text{N}_2\text{O}_4$.

K. F.=1.5% water.

TLC silica gel MeOH.

Rf=0.22.

HPLC: SPHERISORB-SCX column (5 μm , 250 \times 4 mm), T=35° C., mobile phase 50 m KH_2PO_4 : CH_3CN 40:60, pH 4.0 with H_3PO_4 , flow 0.75 ml/min, detector: RI, UV 205 nm, RT=5.33 min.

EXAMPLE 4

R,S-4-trimethylammonium-3-(nonyloxycarbonyl)oxybutyrate chloride (ST 1297)

Benzyl ester of R,S-4-trimethylammonium-3-(nonylcarbamoyl)-oxybutyric acid chloride

Dimethylaminopyridine (3.8 g, 31.2 mmoles) and nonyl chloroformate (6.45 g, 31.2 mmoles) were added to R,S-carnitine perchlorate, benzyl ester (7.33 g, 20.8 mmoles) in anhydrous DMF (50 ml) at 0° C. The temperature was left to raise to room temperature and the reaction mixture was left to stand for 3 days under stirring. CHCl_3 was added and the solution was washed with 1N perchloric acid. The organic phase was dried over anhydrous sodium sulfate and evaporated to dryness, to give 6.02 g crude product, which was purified through flash-chromatography (CHCl_3 : MeOH 85:15). 3.52 g a thick oil were obtained, which were subse-

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quently dissolved in MeOH and passed through an Amberlyst A-21 resin (activated in HCl from), eluting with MeOH. After vacuum-evaporation of the solvent, 3.1 g oily product were obtained.

Yield 32.4%.

¹H-NMR (200 MHz; CDCl₃):

δ: 7.3 (s, 5H), 5.45 (m, 1H), 5.05 (s, 2H), 4.4 (d, 1H), 4.1 (t, 2H), 3.8 (dd, 1H), 3.4 (s, 9H), 2.9 (m, 2H), 1.55 (m, 2H), 1.2 (brs, 12H), 0.8 (brt, 3H).

Mutatis mutandis, the preparation of nonyl chloroformate was carried out as disclosed in Example 2 for tetradecyl chloroformate.

R,S-4-trimethylammonium-3-(nonyloxycarbonyl)-oxybutyric acid chloride

10% Pd/C (110 mg) was added to benzyl R,S-4-trimethylammonium-3-(nonyloxycarbonyl)-oxybutyric acid chloride (1.1 g, 2.4 mmoles) in MeOH (10 ml) and the mixture was hydrogenated at 47 psi for 2 hours. After filtration on celite, the solution was vacuum-dried giving 883 mg product (yield 100%), which was further purified by precipitation from CH₃CN/Et₂O. 600 g of product were obtained.

Yield: 68%.

M.p.: 150° C. dec.

¹H-NMR (300 MHz; D₂O):

δ: 5.4 (m, 1H), 4.1 (m, 2H), 3.75 (dd, 1H), 3.55 (d, 1H), 3.1 (s, 9H), 2.7 (m, 2H), 1.5 (m, 2H), 1.20 (brs, 12H), 0.7 (brt, 3H).

FAB Mass=332, [M⁺].

Elemental analysis: responding to the expected formula

C₁₇H₃₄ClNO₅.

K.F.=1.7% water.

TLC silica gel CHCl₃:MeOH 1:1;

Rf=0.10.

HPLC: SPHERISORB-C₁ column (5 μm, 250×4.6 mm), T=30° C., mobile phase 50 mM (NH₄)H₂PO₄:CH₃CN 60:40, pH 3.0 with H₃PO₄, flow 0.75 ml/min, detector: RI, UV 205 nm, RT=5.67 min.

EXAMPLE 5

R,S-4-trimethylphosphonium-3-(nonylcarbamoyl)oxybutyrate (ST 1300)

Ethyl ester of R,S-4-trimethylphosphonium-3-hydroxybutyric acid iodide

A 1M solution of trimethylphosphine in THF (93 ml) was added to ethyl R,S-4-iodo-3-hydroxybutyrate (20 g, 77.5 mmoles) and the reaction mixture was left to stand at room temperature for 5 days under stirring. Ethyl ether was added, and the precipitate formed was separated by decantation. The precipitate was triturated with Et₂O and dried under vacuum, giving 18.5 g product.

Yield 71.3%.

M.p.: 105–107° C.

¹H-NMR (200 MHz; CDCl₃):

δ: 4.6 (m, 1H), 4.15 (q, 2H), 3.1 (m, 1H), 2.75 (m, 3H), 2.2 (d, 9H), 1.3 (t, 3H).

The ethyl ester of R,S-4-trimethylphosphonium-3-hydroxybutyric acid was prepared as described in Tetrahedron 1990, 4277–4282, starting from R,S-3-hydroxy-4-butyrolactone.

Ethyl ester of R,S-4-trimethylphosphonium-3-(nonylcarbamoyl)-oxybutyric acid iodide

Nonyl isocyanate (4.04 g, 23.86 mmoles) was added to the ethyl ester of R,S-4-trimethylphosphonium-3-

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hydroxybutyric acid iodide (4 g, 11.97 mmoles) in anhydrous DMF (80 ml) and the solution was left to stand for 7 days at 110° C. under stirring. CHCl₃ was added (300 ml) and the solution was washed with water and dried over Na₂SO₄. The residue obtained after evaporation of the solvent was taken up with acetonitrile, the formed solid was filtered off and the filtrate was purified by silica gel flash-chromatography, using CHCl₃: MeOH 8:2. 2.07 g of product in the form of a thick oil were obtained.

Yield: 34.3%.

¹H-NMR (200 MHz; CDCl₃):

δ: 5.4 (m, 2H), 4.15 (q, 2H), 3.15 (m, 4H), 2.8 (d, 2H), 2.2 (d, 9H) 1.5 (m, 2H), 1.2 (brs, 12H), 0.8 (brt, 3H).

R,S-4-trimethylphosphonium-3-(nonylcarbamoyl)oxybutyrate

Ethyl ester of R,S-4-trimethylphosphonium-3-(nonylcarbamoyl)-oxybutyric acid iodide (2.07 g, 4.11 mmoles) was dissolved into 1 N HCl (200 ml) and the solution was warmed to 70° C. for 3 hours. The residue obtained after solvent vacuum-evaporation was taken up with MeOH and passed through Amberlyst A-21 resin, eluting with MeOH. A crude product was obtained, which was purified by flash-chromatography, eluting with MeOH and giving 700 mg product.

Yield: 49%.

M.p.: 123–127° C. dec.

¹H-NMR (300 MHz; D₂O):

δ: 5.3 (m, 1H), 3.1 (m, 2H), 2.80–2.45 (m, 4H), 1.85 (d, 9H), 1.4 (m, 2H), 1.2 (brs, 12H), 0.8 (brt, 3H).

FAB Mass=348, [(M+H)⁺].

Elemental analysis: responding to the expected formula

C₁₇H₃₄NO₄P.

K.F.=3.4% water.

TLC silica gel MeOH;

Rf=0.18.

HPLC: SPHERISORB-SCX column (5 μm, 250×4 mm), T=25° C., mobile phase 50 mM KH₂PO₄:CH₃CN 40:60, pH 4.0 with H₃PO₄, flow 0.75 ml/min, detector: RI, UV 205 nm, RT=5.18 min.

The following Examples 6 and 7 are further illustrated by FIG. 1.

EXAMPLE 6

R,S-4-trimethylammonium-3-(octyloxycarbonyl)-aminobutyrate chloride (ST 1253) (2a, FIG. 1)

Step A

3 g (0.012 mmoles) aminocarnitine isobutyl ester were dissolved into 20 ml anhydrous CH₂Cl₂. 2.48 ml (0.1078 moles) triethylamine and 3.6 g (0.0178 moles) octyl chloroformate (previously prepared by reacting the alcohol with a toluene solution of phosgene) were added to the solution. The reaction mixture was left to stand for 4.5 hours at room temperature. Then the solvent was evaporated off and the resulting solid was dissolved into ethyl acetate and filtered. The solvent was vacuum-evaporated to dryness and the resulting solid was purified on silica gel, eluting with 100% CHCl₃, then with CHCl₃:MeOH 95:5 and 90:10. The product was obtained with a 50% yield.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5)/acetone 7:3;

Rf=0.8.

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HPLC: SPHERISORB-SCX column (5 μ m, 250 \times 4 mm), mobile phase 50 mM (NH₄)H₂PO₄:CH₃CN 60:40, pH 4.0, detector: RI, UV 205 nm, RT=8.6 min.

¹H-NMR (300 MHz; CD₃OD):

δ : 4.56-4.46 (m, 1H), 4.12-4.02 (m, 2H), 3.94-3.88 (m, 2H), 3.66-3.5 (s, 9H), 3.4 (s, 9H), 2.74-2.66 (m, 2H), 2-1.86 (m, 1H), 1.68-1.56 (t, 2H), 1.4-1.2 (m, 12H), 0.97-0.7 (d, 6H), 0.6-0.3 (t, 3H).

Elemental analysis: responding to the expected formula C₂₀H₄₃ N₂O₄Cl.

Step B

The ester obtained in Step A was hydrolysed on Amberlyst IRA 402 resin (OH⁻ activated form) eluting with water. Water was evaporated to dryness; the resulting solid was triturated with acetone and subsequently filtered. A white solid was obtained.

Yield 94%.

M.p.=170° C. dec.

¹H-NMR (300 MHz; CD₃OD):

δ : 4.4 (m, 1H), 4.05 (t, 2H), 3.5 (d, 2H), 3.2 (s, 9H), 2.4 (d, 2H), 1.6 (m, 2H), 1.4-1.2 (m, 12H), 0.95-0.85 (t, 3H).

FAB Mass=454, [(M+H)⁺].

Elemental analysis: responding to the expected formula C₁₆H₃₂N₂O₄

K.F.=1.74% water.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5)

Rf=0.65.

HPLC: SGE-SCX column (5 μ m, 250 \times 4 mm), mobile phase 0.05M (NH₄)H₂PO₄:CH₃CN 60:40, detector: RI, UV 205 nm, RT=9.0 min.

EXAMPLE 7

R,S-4-trimethylammonium-3-(nonyloxy carbonyl)-aminobutyrate (ST 1285) (**2b**, FIG. 1)

Step A

The product was prepared as disclosed in Example 6, step A, using nonyl chloroformate

Yield: 50%.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5)/acetone 7:3

Rf=0.71.

HPLC: SGE-SCX column (5 μ m, 250 \times 4 mm), mobile phase 50 mM (NH₄)H₂PO₄:CH₃CN 60:40, pH 4.0, detector: RI, UV 205 nm, RT=10.417 min.

¹H-NMR (300 MHz; CD₃OD):

δ : 4.54-4.44 (m, 1H), 4.1-4.02 (m, 2H), 3.96-3.86 (m, 2H), 3.6-3.5 (m, 2H), 3.2 (s, 9H), 2.72-2.66 (m, 2H), 2-1.86 (m, 1H), 1.66-1.56 (m, 2H), 1.38-1.26 (m, 14H), 0.96-0.94 (d, 6H), 0.92-0.86 (t, 3H).

Step B

The product was prepared as disclosed in Example 6, step B.

Yield 80%.

M.p.=160° C. dec.

¹H-NMR (300 MHz; CD₃OD):

δ : 4.5-4.35 (m, 1H), 4.1-4.0 (t, 2H), 3.55-3.45 (d, 2H), 3.2 (s, 9H), 2.45-2.35 (d, 2H), 1.7-1.5 (m, 2H), 1.4-1.2 (m, 14H), 0.9-0.8 (t, 3H).

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Elemental analysis: responding to the expected formula C₁₇H₃₄N₂O₄

K.F.=1.3% water.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5);

Rf=0.62.

HPLC: SGE-SCX column (5 μ m, 250 \times 4 mm), mobile phase 0.05M (NH₄)H₂PO₄:CH₃CN 60:40, detector: RI, UV 205 nm, RT=7.56 min.

The following Examples 8-9 are further illustrated by FIG. 2.

EXAMPLE 8

R,S-4-trimethylammonium-3-octyloxybutyrate (ST 1207) (**6a**, FIG. 2)

Step A

39 g (0.3 moles) octyl alcohol were dissolved in 25 ml toluene and 14.5 ml (0.107 moles) ethyl chloroacetate and 8 ml Thionyl chloride were added thereon at -15° C. At the end of the addition, the reaction mixture was left to stand for 4 hours at room temperature Ethyl acetate was then added and the solution was washed three times with 1N NaOH and subsequently with water. The organic phase was treated with anhydrous sodium sulfate, filtered and vacuum-evaporated to dryness. The product was purified on silica gel chromatographic column, eluting with gradient from hexane alone to hexane/ethyl ether 95:5. The product was obtained with 80% yield.

TLC silica gel hexane/ ethyl ether 85:15;

Rf=0.75.

¹H-NMR (300 MHz; CDCl₃):

δ : 4.2-4.09 (q, 2H), 3.80 (s, 2H), 3.4-3.5 (dd, 2H), 2.85 (s, 2H), 1.60-1.58 (m, 2H), 1.4-1.2 (m, 10H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₂₂H₃₃ ClO₄.

Step B

9 ml BF₃.Et₂O were dropped to a mixture of 26.8 g (0.066 moles) of the product obtained in the proceeding step A and 13.5 ml triethylsilane at 0° C. At the end of the addition, the reaction mixture was refluxed for 4 hours. After cooling, ether was added and the solution was washed twice with NaOH 1N, then water; the organic phase was dried over anhydrous sodium sulfate, filtered and vacuum-evaporated to dryness. An oil was obtained, which was purified on silica gel chromatographic column, eluting with gradient from hexane alone to hexane/ethyl ether 95:5. The product was obtained with a 70% yield.

TLC silica gel hexane/ethyl ether 90:10;

Rf=0.47. ¹H-NMR (300 MHz; CDCl₃):

δ : 4.2-4.09 (dd, 2H), 4.0-3.85 (m, 1H), 3.62-3.40 (m, 4H), 2.70-2.50 (dd, 2H), 1.55-1.50 (m, 2H), 1.4-1.2 (m, 10H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₁₄H₂₇ClO₃

Step C

5.2 g (0.08 moles) NaN₃ and a catalytic amount of tetrabutyl ammonium bromide were added to a solution of 11.4 g (0.041 mmoles) product obtained in the preceding step B. The reaction mixture was left for three nights at 60° C. The

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solution was vacuum-evaporated to dryness. A thick dark solution was obtained, which was purified on silica gel chromatographic column, eluting with gradient from hexane alone to hexane/ethyl ether 95:5. The product was obtained with a 83% yield.

TLC silica gel hexane/ethyl ether 95:5;

Rf=0.23.

¹H-NMR (300 MHz; CDCl₃):

δ: 4.2-4.09 (dd, 2H), 4.0-3.80 (m, 1H), 3.60-3.40 (dd, 2H), 3.40-3.20 (dd, 2H), 2.70-2.40 (dd, 2H), 1.60-1.40 (m, 2H), 1.4-1.1 (m, 10H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₁₄H₂₇N₃O₃

Step D

The product obtained in the preceding step C (15.39 g, 0.054 moles) was dissolved in 31 ml of acetic acid and the resulting solution was subjected to catalytic hydrogenation with 10% Pd/C at 60 psi for 7 hours. The reaction process was checked by TLC, until disappearance of the starting product (hexane/ethyl ether 95:5). Thereafter, formaldehyde was added (4.6 ml, 0.167 moles) followed by 10% Pd/C and the mixture was hydrogenated at 30 psi for 2 days. The catalyst was filtered off and the mixture was vacuum-dried. A pale yellow liquid was obtained, which was taken up with methylene chloride, washed with 1N NaOH, then water, then NaCl saturated solution; the organic phase was dried over anhydrous sodium sulfate, filtered and vacuum-evaporated to dryness. A thick oil was obtained. The product was obtained with a 98% yield.

TLC silica gel AcOEt/MeOH/NH₃ 90:10:3;

Rf=0.42.

¹H-NMR (300 MHz; CDCl₃):

δ: 4.2-4.09 (dd, 2H), 3.85-3.80 (m, 1H), 3.60-3.40 (dd, 2H), 2.65-2.40 (dd, 2H), 2.40-2.20 (dd, 2H), 2.20 (s, 6H), 1.60-1.40 (m, 2H), 1.4-1.1 (m, 10H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₁₆H₃₆N₃O₃

Step B

The product obtained in the preceding step D (15.21 g, 0.053 moles) was dissolved in 98 ml THF and 8 ml methyl iodide were added thereto. The reaction progress was left overnight at room temperature. The mixture was vacuum-evaporated to dryness. A thick oil was obtained. The product was obtained with a 98% yield.

TLC silica gel AcOEt/MeOH/NH₃ 90:10:3;

Rf=0.10.

¹H-NMR (300 MHz; CDCl₃):

δ: 4.5-4.3 (m, 1H), 4.2-4.09 (dd, 2H), 3.75-3.30 (m, 2H), 3.5 (s, 9H), 2.75-2.60 (dd, 2H), 1.60-1.45 (m, 2H), 1.30-1.15 (m, 10H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₁₆H₃₉INO₃

Step F

The product obtained in the preceding step E, was hydrolysed on Amberlyst IRA 402 resin (OH⁻ activated form) eluting with water. Water was evaporated to dryness; the resulting solid was treated with isopropyl alcohol three times. A white solid was obtained.

Yield=93%

M.p.=106° C. dec.

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¹H-NMR (300 MHz; MeOD):

δ: 4.30-4.15 (m, 1H), 3.70-3.60 (dd, 1H), 3.50-3.40 (m, 2H), 3.20 (s, 9H), 2.75-2.65 (dd, 1H), 2.20-2.10 (dd, 1H), 1.60-1.50(m, 2H), 1.40-1.20 (m, 10H), 0.9-0.8 (t, 3H).

Elemental analysis: responding to the expected formula C₁₅H₃₁NO₃.

K.F.=5.7% water.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5);

Rf=0.7.

HPLC: SGE-SAX column (5 μm, 250×4 mm), mobile phase 0.025M (NH₄)H₂PO₄:CH₃CN 30:70, detector: RI, UV 205 nm, flow=0.75 ml/min, RT=5.85 min.

MS-FAB+glycerol matrix=274.

EXAMPLE 9

R,S-4-trimethylammonium-3-tetradecyloxybutyrate (ST 1228) (6b, FIG. 2)

Step A

The product was prepared as in example 8, step A using tetradecyl alcohol. The product was obtained with 73% yield.

TLC silica gel hexane/ethyl ether 95:5;

Rf=0.63.

¹H-NMR (300 MHz; CDCl₃):

δ: 4.2-4.09 (q, 2H), 3.80 (s, 2H), 3.4-3.5 (dd, 2H), 2.85 (s, 2), 1.60-1.58 (m, 2H), 1.4-1.2 (m, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₃₄H₆₇ClO₄.

Step B

The product was prepared as in example 8, step B. The product 2b, shown in FIG. 2, was obtained with a 72% yield.

TLC silica gel hexane/ethyl ether 95:5;

Rf=0.4.

¹H-NMR (300 MHz; CDCl₃):

δ: 4.2-4.09 (dd, 2H), 4.0-3.85 (m, 1H), 3.62-3.40 (m, 4H), 2.70-2.50 (dd, 2H), 1.55-1.50 (m, 2H), 1.4-1.2 (m, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₂₀H₃₉O₃

Step C

The product was prepared as in example 8, step C. The product was obtained with 79% yield.

TLC silica gel hexane/ethyl ether 90:10;

Rf=0.36.

¹H-NMR (300 MHz; CDCl₃):

δ: 4.2-4.09 (dd, 2H), 4.0-3.80 (m, 1H), 3.60-3.40 (dd, 2H), 3.40-3.20 (dd, 2H), 2.70-2.40 (dd, 2H), 1.60-1.40 (m, 2H), 1.4-1.1 (m, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₂₀H₃₉N₃O₃

Step D

The product was prepared as in example 8, step D. The product was obtained with a 98% yield.

TLC silica gel AcOEt/MeOH/NH₃ 90:10:3;

Rf=0.72.

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¹H-NMR (300 MHz; CDCl₃):

δ: 4.2-4.09 (dd, 2H), 3.85-3.80 (m, 1H), 3.60-3.40 (dd, 2H), 2.65-2.42 (dd, 2H), 2.38-2.20 (dd, 2H), 2.18 (s, 6H), 1.60-1.40 (m, 2H), 1.4-1.1 (m, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₂₂H₄₅NO₃.

Step E

The product was prepared as in example 8, step E. The product was obtained with a 99% yield.

TLC silica gel AcOEt/MeOH/NH₃ 90:10:3;

Rf=0.15.

¹H-NMR (300 MHz; CDCl₃):

δ: 4.45-4.3 (m, 1H), 4.2-4.09 (dd, 2H), 3.75-3.30 (m, 2H), 3.5 (s, 9H), 2.75-2.60 (dd, 2H), 1.60-1.45 (m, 2H), 1.30-1.15 (m, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₂₃H₄₈INO₃.

Step F

The product was prepared as in example 8, step F. The product was obtained with a 99% yield.

M.p.=106° C. dec.

¹-NMR (300 MHz; DMSO-D6):

δ: 4.10-4.0 (m, 1H), 3.60-3.20 (m, 4H), 3.05 (s, 9H), 2.40-2.30 (dd, 1H), 1.80-1.70 (dd, 1H), 1.50-1.40 (m, 2H), 1.30-1.15 (m, 22H), 0.9-0.8 (t, 3H).

Elemental analysis: responding to the expected formula C₂₁H₄₃NO₃.

K.F.=6.4% water.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5);

Rf=0.6.

HPLC: SGE-SCX column (5 μm, 250×4 mm), mobile phase 0.05M (NH₄)H₂PO₄:CH₃CN 40:60, detector: RI, UV 205 nm, flow=0.75 ml/min, RT=4.38 min.

MS-FAB+glycerol matrix=358.3.

The following Examples 10–11 are further illustrated by FIG. 3a–b.

EXAMPLE 10

R,S,-1-guanidinium-2-tetradecyloxy-3-(tetrazolate-5-yl)propane (ST 1263) (10, FIG. 3b)

Step A

6.65 g (0.0179 moles) of the intermediate prepared in Example 9, step C were dissolved in 10 ml of methanol and 10 ml of 4N NaOH were added to the solution. The reaction was left to stand for 16 hours at room temperature. 20 ml 6N HCl were added to the solution, which was extracted with ethyl acetate. The organic phase was dried over anhydrous sodium sulfate, filtered and vacuum concentrated. The product was obtained as a white solid with a 95.6% yield.

TLC silica gel hexane /ethyl ether 1:1;

Rf=0.5.

M.p.=42–45° C.

¹H-NMR (300 MHz; CD₃OD):

δ: 3.9-3.8 (m, 1H), 3.56-3.48 (m, 2H), 3.42-3.26 (dd, 2H), 2.68-2.5 (m, 2H), 1.6-1.5 (m, 2H), 1.4-1.2 (s, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₁₈H₃₅N₃O₃.

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Step B

At 0° C., 4.96 ml TEA were dropped into a solution containing 2.79 g (8.19 mmoles) of the compound obtained in step A, aminopropionitrile (0.58 g, 8.2 mmoles) and DEPC (diethylphosphocyanate) (1.71 ml) in 4.2 ml of anhydrous DMF. The reaction was left to stand for 1 hour at room temperature. The solvent was evaporated and the residue was dissolved in ethyl acetate, washed twice with water, then with a NaCl saturated solution. The organic phase was dried over anhydrous sodium sulfate, filtered and vacuum concentrated. The product was obtained and purified through a silica gel column with hexane: ethyl ether (7:3/1:1/3:7).

Yield: 71%.

TLC silica gel ethyl ether 100%;

Rf=0.42.

¹H-NMR (300 MHz; CDCl₃):

δ: 6.6-6.4 (m, 1H), 3.9-3.8 (m, 1H), 3.60-3.4 (m, 5H), 3.3-3.2 (dt, 1H), 2.7-2.6 (t, 2H), 2.6-2.4 (dd, 2H), 1.6-1.5 (m, 2H), 1.4-1.2 (m, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₂₁H₃₉N₅O₂

Step C

2.99 g (0.0114 moles) triphenylphosphine and 0.2 ml water were added to a solution containing 2.99 g (7.62 mmoles) of the compound obtained in step B. The reaction was left to stand overnight at room temperature. The solvent was evaporated off and the product was obtained and purified through a silica gel column with ethyl acetate 100%, then ethyl acetate:methanol:ammonia 7:3:0.3.

Yield: 65%.

TLC silica gel ethyl acetate:methanol:ammonia 7:3:0.3;

Rf=0.26.

¹H-NMR (300 MHz; CD₃OD):

δ: 3.78-3.7 (m, 1H), 3.58-3.48 (m, 4H), 2.8-2.7 (dd, 2H), 2.7-2.6 (m, 2H), 2.5-2.3 (dd, 2H), 1.6-1.5 (m, 2H), 1.4-1.3 (m, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₂₁H₄₁N₃O₂

Step D

1.69 g (4.6 mmoles) of the compound obtained in step C were treated with 1.2 g (5.2 mmoles) (BOC)₂O and 9.2 ml 1N NaOH for 30 minutes at room temperature. The reaction mixture was poured into ethyl acetate and washed four times with 1N HCl, then water and a saturated NaCl solution. The organic phase was dried over anhydrous sodium sulfate, filtered and vacuum concentrated to dryness. The product was obtained as a white solid.

Yield: 100%.

TLC silica gel ethyl ether 100%;

Rf=0.26.

M.p.=83–84° C.

¹H-NMR (300 MHz; CDCl₃):

δ: 7.2-7.0 (m, 1H), 4.9-4.8 (m, 1H), 3.8-3.6 (m, 1H), 3.5-3.4 (dt, 4H), 3.2-3.0 (m, 2H), 2.6 (t, 2H), 2.4 (d, 2H), 1.5 (m, 2H), 1.4 (s, 9H), 1.4-1.2 (m, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₂₆H₄₉N₃O₄.

Step E

The product obtained in step D (1.19 g, 2.56 mmoles) was dissolved into 12 ml of anhydrous THF, under argon

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atmosphere, then 3.062 g of triphenylphosphine, 1.54 ml of triethylsilylazido and 4.9 ml of DEAD (diethylazodicarboxylate) were dropped at 0° C. within three days, until disappearance of the starting product. The mixture was then treated with an aqueous solution of cerium ammonium nitrate and diluted with CH₂Cl₂. The reaction was left to stand for 2 hours, the organic phase was washed with a saturated NaCl solution, dried over anhydrous sodium sulfate and vacuum-dried. The residue was purified through a silica gel column with hexane/ethyl acetate (9:1/8:2/7:3). The product was obtained with a 66% yield.

TLC silica gel hexane/AcOEt 1:1;

Rf>0.34.

¹H-NMR (300 MHz; CDCl₃):

δ: 4.95-4.8 (m, 1H), 4.7-4.5 (m, 2H), 3.9-3.8 (m, 1H), 3.50-3.40 (m, 1H), 3.40-3.31 (m, 1H), 3.3-3.2 (m, 1H), 3.22-3.0 (dd, 2H), 3.10-3.0 (m, 3H), 1.45-1.35 (m, 1H), 1.2 (m, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₂₅H₄₈N₆O₃

Step F

The product obtained in step E (0.969 g, 1.97 mmoles) was dissolved into 13.09 ml anhydrous THF, then 13.1 ml of 3N HCl were added. The reaction mixture was left to stand for 2 hours, at 50° C. under stirring. The reaction mixture was vacuum-dried, the residue was taken up with CH₂Cl₂ and treated with a 1 N NaOH solution. The organic phase was separated, dried over anhydrous sodium sulfate and vacuum-dried. The product was obtained with a 92% yield.

TLC silica gel AcOEt/MeOH/NH₃ 9:1:0.3

Rf=0.31.

¹H-NMR (300 MHz; CDCl₃):

δ: 4.78-4.58 (m, 2H), 3.8-3.7 (m, 1H), 3.5-3.4 (m, 1H), 3.30-3.24 (m, 1H), 3.24-3.18 (m, 4H), 3.05-3.0 (dd, 2H), 3.0-2.6 (dd, 2H), 1.4 (m, 2H), 1.2 (m, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₂₁H₄₀N₆O

Step G

The product obtained in step F (2.78 g, 7.1 mmoles) was dissolved into 20 ml anhydrous MeOH, then 2.34 g iminomethanesulfonic acid (prepared with well-known methods) were added within 3 days. The obtained suspension was vacuum-concentrated, then treated with 1N NaOH and left under stirring for 30 minutes. The solid was filtered, washed with water, then acetone. The title product was obtained with a 45% yield.

TLC silica gel AcOEt/MeOH/NH₃ 7:3:0.3;

Rf=0.22.

M.p.=240° C. dec.

¹H-NMR (300 MHz; CD₃OD):

δ: 3.90-3.75 (m, 1H), 3.6-3.4 (m, 2H), 3.40-3.20 (m, 2H), 3.20-3.10 (dd, 1H), 2.95-2.85 (dd, 1H), 1.4 (m, 2H), 1.2 (s, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula C₁₉H₃₉N₇O.

HPLC: Spherisorb-C1 (5 μm, 250×4.6 mm), mobile phase 0.05 M KH₂PO₄:CH₃CN 35:65, pH=3, flow 0.75 ml/min, detector: UV 205 nm, RT=5.51 min.

MS-FAB+glycerol matrix=382.

EXAMPLE 11

R,S-1-trimethylammonium-2-tetradecyloxy-3-(tetrazolato-5-yl)propane (ST 1287) (9, FIG. 3b)

Steps A-F

The compounds were prepared as in steps A-F of Example 10.

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Step H

2.79 g (7.14 mmoles) of the compound prepared in Example 10, step F were suspended in 18 ml water and 1.47 ml HCOOH and 1.57 ml H₂CO were added thereto. The reaction mixture was refluxed overnight, then was allowed to cool down and methylene chloride was added; pH was adjusted to 9 with 0.5 N NaOH. The mixture was extracted three times with methylene chloride. The organic phase was washed with 0.5 N NaOH, water and dried over anhydrous sodium sulfate, filtered and vacuum concentrated. The product was obtained as a solid with a 100% yield.

TLC silica gel AcOEt/MeOH/NH₃ 9:1:0.3;

Rf=0.58.

¹H-NMR (300 MHz; CDCl₃):

δ: 4.7-4.5 (m, 1H), 3.8-3.7 (m, 1H), 3.5-3.4 (m, 1H), 3.30-3.20 (m, 2H), 3.10 (m, 3H), 2.45-2.35 (m, 2H), 2.30 (s, 6H), 1.4-1.3 (m, 2H), 1.2-1.0 (m, 22H), 0.90-0.80 (t, 3H).

Elemental analysis: responding to the expected formula

C₂₃H₄₄N₆O.

Step I

2.99 g (7.14 mmoles) of the compound obtained in step H were dissolved in THF and 2.5 ml of CH₃I were added thereto. The reaction was left to stand for 3 hours at room temperature. The solvent was evaporated off and the solid residue was washed with hot ether, left overnight under stirring, then filtered. The product was obtained.

Yield: 100%.

TLC silica gel CHCl₃:iPrOH:MeOH:H₂O:CH₃COOH 42:7:28:10.5:10.5;

Rf=0.73.

¹H-NMR (300 MHz; CDCl₃):

δ: 4.90-4.80 (m, 2H), 4.70-4.55 (m, 1H), 4.40-4.25 (m, 1H), 3.80-3.60 (m, 2H), 3.60-3.40 (m, 3H), 3.30 (s, 9H), 3.30-3.10 (m, 2H), 1.60-1.40 (m, 2H), 1.3-1.1 (m, 22H), 0.9-0.8 (t, 3H).

Elemental analysis: responding to the expected formula C₂₄H₄₇N₆O.

MS-FAB+glycerol matrix=436.

Step L

The product obtained in step I (2.99 g, 5.33 mmoles) was dissolved in MeOH, then passed through IRA 402 resin in OH⁻ form, conditioned in MeOH. The title product was obtained as a solid, which was subsequently triturated with AcOEt.

Yield=88%.

TLC silica gel CHCl₃:iPrOH:MeOH:H₂O:CH₃COOH (42:7:28:10.5:10.5)/acetone 8:2;

Rf=0.73.

TLC silica gel CHCl₃:iPrOH:MeOH:H₂O:CH₃COOH 42:7:28:10.5:10.5;

Rf=0.73.

M.p.=180° C. dec.

¹H-NMR (300 MHz; CDCl₃):

δ: 4.30-4.20 (m, 1H), 3.90-3.70 (m, 2H), 3.60-3.55 (m, 1H), 3.50-3.30 (m, 4H), 3.25 (m, 1H), 3.0-2.9 (m, 1H), 1.60-1.40 (m, 2H), 1.3-1.1 (m, 22H), 0.9-0.8 (t, 3H).

Elemental analysis: responding to the expected formula C₂₁H₄₃N₅O.

MS-FAB+glycerol matrix =382.

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K.F.=1% water.

HPLC: Spherisorb-C1 (5 μ m, 250 \times 4.6 mm), mobile phase 0.05 M KH_2PO_4 : CH_3CN 35:65, pH=3, flow 0.75 ml/min, detector: UV 205 nm, RT=5.18 min.

The following Examples 12–14 are further illustrated by FIG. 4.

EXAMPLE 12

R,S-3-quinuclidinium-2-(tetradecyloxycarbonyl)oxy-1-propanephosphonate monobasic (ST 1260)

Step A

In anhydrous environment, -70°C ., a hexane solution of 1.6 M BuLi (14 ml, 0.022 moles) was dropped into a solution of dibenzyl phosphite (5.8 g, 0.022 mmoles) in THF. After 15 minutes, 1.8 ml (0.022 moles) of epibromhydrine, dissolved in 5 ml THF, were added. After the addition, etherated BF_3 (3.6 ml, 0.022 moles) was dropped very slowly. The reaction was left for further 3 hours at -70°C . A saturated ammonium chloride solution was added; then the temperature was left to raise to room temperature. This solution was extracted several times with AcOEt and the gathered organic phases were treated with saturated NaHCO_3 , and dried over anhydrous sodium sulfate, filtered and vacuum concentrated. An oil was obtained, which after purification on silica gel chromatography (AcOEt/Hexane 1:1), gave 1.1 g of unreacted dibenzylphosphite and 5.3 g of product of interest.

Yield=60%.

TLC silica gel AcOEt/Hexane 7:3;

Rf=0.54.

$^1\text{H-NMR}$ (300 MHz; CD_3OD):

δ : 7.4-7.2 (m, 10H), 5.1-4.9 (m, 4H), 4.2-4.0 (m, 1H), 3.5-3.3 (dd, 2H), 2.2-2.0 (m, 2H).

Elemental analysis: responding to the expected formula $\text{C}_{17}\text{H}_{20}\text{BrO}_4\text{P}$.

MS-FAB+glycerol matrix=399, 400, 401, 402.

Step B

2 g (5. mmoles) of the compound obtained in Step A were dissolved at 10% concentration and the solution cooled down to 0°C . 1.4 ml TEA (10 mmoles) and 0.62 g (5 mmoles) DMAP (dimethylaminopyridine) were dropped thereto. Immediately after, 5.2 mmoles tetradecyl chloroformate were added and the temperature was left to raise to room temperature. The reaction progress was checked on TLC and worked up at the disappearance of the starting compound. Further chloroform was added and the reaction mixture was washed with 1N HCl and water. After drying over anhydrous sodium sulfate, the solvent was evaporated off and an oil was obtained, which was purified through flash-chromatography using hexane/AcOEt 7:3 as eluant. The product was obtained.

Yield: 75%.

TLC silica gel hexane/AcOEt 7:3;

Rf=0.31.

$^1\text{H-NMR}$ (300 MHz; CDCl_3):

δ : 7.4-7.2 (m, 10H), 5.1-4.9 (m, 5), 4.1-3.9 (m, 2H), 3.6-3.4 (dd, 2H), 2.4-2.2 (m, 2H), 1.6-1.4 (m, 2H), 1.3-1.1 (m, 22H), 0.9-0.7 (t, 3H).

Elemental analysis: responding to the expected formula $\text{C}_{32}\text{H}_{48}\text{BrO}_6\text{P}$.

Step D

The product obtained in step B (6.39 g, 10 mmoles) was dissolved in 12 ml DMF, then quinuclidine was added (2.2 g,

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20 mmoles) together with TBAI (tetrabutyl ammonium iodide) in catalytic amounts (1% by weight with respect to the substrate). The reaction was carried out at a temperature of 50°C ., until the starting product disappeared. At the end of reaction, the mixture was concentrated under high vacuum, obtaining a semisolid containing the product. The latter was purified through silica gel flash-chromatography, using $\text{CHCl}_3/\text{MeOH}$ 8:3. The product was obtained.

Yield=15%.

TLC silica gel CHCl_3 :iPrOH:MeOH:H₂O:CH₃COOH (42:7:28:10.5:10.5)/acetone 8:2;

Rf=0.8.

$^1\text{H-NMR}$ (300 MHz; MeOD):

δ : 7.4-7.1 (m, 50H), 5.3-5.1 (m, 1H), 4.9-4.8 (d, 2H), 4.1-4.0 (m, 2H), 3.8-3.4 (m, 2H), 3.4-3.2 (m, 6H), 2.2-1.7 (m, 9H), 1.6-1.4 (m, 2H), 1.3-1.1 (m, 22H), 0.9-0.7 (t, 3H).

Elemental analysis: responding to the expected formula $\text{C}_{32}\text{H}_{54}\text{NO}_6\text{P}$.

MS-FAB+glycerol matrix=580.

Step E

The product obtained in Step D was dissolved in MeOH, then 10% Pd/C (5% by weight with respect to the substrate) was added; the dispersion was hydrogenated (60 psi) at room temperature for 18 hours. At the end, the dispersion was filtered through celite and concentrated to dryness. The title product was obtained without further purifications.

Yield=99%.

TLC silica gel CHCl_3 :iPrOH:MeOH:H₂O:CH₃COOH (42:7:28:10.5:10.5)/acetone 8:2;

Rf=0.57.

$^1\text{H-NMR}$ (300 MHz; D_2O):

δ : 5.5-5.3 (m, 1H), 4.2-4.1 (m, 2H), 4.0-3.4 (m, 8H), 2.2-1.7 (m, 9H), 1.60-1.40 (m, 2H), 1.3-1.1 (m, 22H), 0.9-0.7 (t, 3H).

Elemental analysis: responding to the expected formula $\text{C}_{25}\text{H}_{48}\text{NO}_6\text{P}$.

MS-FAB+glycerol matrix=490.

K.F.=7% water.

HPLC: Spherisorb-C1 (5 μ m, 250 \times 4.6 mm), mobile phase 0.075 M KH_2PO_4 : CH_3CN 60:40, flow 0.75 ml/min, detector: RI, UV 205 nm, RT=16.53 min.

EXAMPLE 13

R,S-3-trimethylammonium-2-(nonylaminocarbonyl)oxy-1-propanephosphonate monobasic (ST 1286)

Step A

The product was prepared as disclosed in step A of Example 12.

Step C

The product obtained in the previous step (4 g, 10 mmoles) was dissolved in CH_2Cl_2 (10% solution) and etherated BF_3 (1.6 ml) and nonyl isocyanate (3.38 g, 20 mmoles) were added at room temperature. The reaction was worked up after 30 minutes, firstly adding further CH_2Cl_2 , then washing the organic phase with 1N NaOH several times. The product was purified on silica gel flash-chromatography (Hexane/AcOEt 7:3).

Yield=85%.

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TLC silica gel AcOEt/Hexane 6:4;

Rf=0.28.

¹H-NMR (300 MHz; CDCl₃):

δ: 7.4-7.2 (m, 10H), 5.1-4.9. (m, 5H), 4.6-4.2 (m, 1H),
3.7-3.5 (dd, 2H), 3.2-3.0 (m, 2H), 2.4-2.2 (m, 2H), 1.5-1.3
(m, 2H), 1.3-1.1 (m, 12H), 0.9-0.7 (t, 3H).

Elemental analysis: responding to the expected formula
C₂₇H₄₀BrNO₅P.

Step F

The compound obtained in the preceding step (5.68 g, 10
mmoles) was dissolved in DMF (11 ml), together with TBAI
(tetrabutyl ammonium iodide) in catalytic amounts (1% w/w
with respect to the substrate). This solution was saturated
with gaseous trimethylamine. The reaction was carried out at
50° C., until the starting compound disappeared. At the end
of the reaction, the solution was high vacuum-concentrated,
obtaining a semisolid, containing the product. The latter was
isolated and purified through silica gel flash-
chromatography using a gradient from CH₂Cl₂ only to
CH₂Cl₂:MeOH 1:1. The product was obtained.

Yield: 25%.

TLC silica gel CHCl₃:iPrOH:MeOH:H₂O:CH₃COOH
(42:7:28:10.5:10.5)/acetone 8:2;

Rf=0.73.

¹H-NMR (300 MHz; CDCl₃):

δ: 7.5-7.2 (m, 5H), 5.5-5.4 (m, 1H), 4.9-4.8 (m, 4H),
4.0-3.6 (m, 2H), 3.2-3.1 (s, 9H), 2.2-2.1 (s, 9H), 2.0-1.8
(m, 2H), 1.5-1.4 (m, 2H), 1.4-1.2 (m, 12H), 0.9-0.7 (t, 3H).

Elemental analysis: responding to the expected formula
C₂₇H₄₂N₂O₅P.

MS-FAB+glycerol matrix=457.

Step G

The product obtained in step F was dissolved in MeOH,
then 10% Pd/C (5% by weight with respect to the substrate)
was added; the dispersion was hydrogenated (60 psi) at room
temperature for 18 hours. At the end, the dispersion was
filtered through celite and concentrated to dryness. The title
product was obtained without further purifications.

Yield=99%.

TLC silica gel CHCl₃:iPrOH:MeOH:H₂O:CH₃COOH
(42:7:28:10.5:10.5)/acetone 8:2;

Rf=0.31.

¹H-NMR (300 MHz; D₂O):

δ: 5.6-5.5 (m, 1H), 4.1-3.5 (m, 2H), 3.2-3.1 (s, 9H), 3.1-
3.0 (m, 2H), 2.2-1.7 (m, 2H), 1.5-1.4 (m, 2H), 1.4-1.2 (m,
12H), 0.9-0.7 (t, 3H).

Elemental analysis: responding to the expected formula
C₁₅H₃₅N₂O₅P.

MS-FAB+glycerol matrix=367.

K.F.=3% water.

HPLC: Spherisorb-C1 (5 μm, 250×4.6 mm), mobile
phase 0.05 M (NH₄)H₂PO₄:CH₃CN 35:65, flow 0.75
ml/min, detector: RI, UV 205 nm, RT=7.31 min.

EXAMPLE 14

R,S-3-pyridinium-2-(nonylamino-carbonyl)-oxy-1-
propanephosphonic acid chloride (ST 1268)

Step A

The product was prepared as disclosed in Step A of
Example 12.

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Step C

The product was prepared as disclosed in Step C of
Example 13.

Step H

The compound obtained in the preceding step (5.68 g, 10
mmoles) was dissolved in anhydrous pyridine (50%
solution), together with TBAI (tetrabutyl ammonium iodide)
in catalytic amounts (1% w/w with respect to the substrate).
The reaction was carried out at 50° C., until the starting
compound disappeared. At the end of the reaction, the solu-
tion was high vacuum-concentrated, obtaining a semisolid,
containing the product, which was isolated and purified
through silica gel flash-chromatography using a gradient
from CH₂Cl₂ only to CH₂Cl₂:MeOH from 9:1 to 1:1.

Yield: 20%.

TLC silica gel CHCl₃:iPrOH:MeOH:H₂O:CH₃COOH
(42:7:28:10.5:10.5)/acetone 8:2;

Rf=0.73.

¹H-NMR (300 MHz; CDCl₃):

δ: 9.4-9.3 (d, 2H), 8.2-8.1 (t, 1H), 7.9-7.8 (t, 2H), 7.3-7.1
(m, 5H), 5.3-5.1 (m, 3H), 4.9-4.8 (m, 2H), 3.0-2.9 (m, 2H),
2.2-1.6 (m, 2H), 1.4-1.2 (m, 2H), 1.3-1.1 (m, 12H), 0.9-0.7
(t, 3H).

Elemental analysis: responding to the expected formula
C₂₄H₃₈N₂O₅P.

MS-FAB+glycerol matrix=477.

Step I

The product obtained in step H (4.76 g, 10 mmoles) was
dissolved in 100 ml CH₂Cl₂ and 20 mmoles TMSI
(trimethylsilyl iodide) were added to the resulting solution.
After 30 minutes, the reaction was finished; 0.5 ml water
were added to the mixture, which was concentrated to dry-
ness. The final product was purified and isolated by RP-18
silica gel chromatography, using a gradient water/methanol
9:1 to methanol 100%. The solid was dissolved in water and
passed through IRA 402 resin (Cl⁻ activated). ST 1268 was
obtained.

Yield=80%.

M.p.=202-204° C.

TLC silica gel CHCl₃:iPrOH:MeOH:H₂O:CH₃COOH
(42:7:28:10.5:10.5)/acetone 8:2;

Rf=0.48.

¹H-NMR (300 MHz; D₂O):

δ: 9.4-9.3 (d, 2H), 8.2-8.1 (t, 1H), 7.9-7.8 (t, 2H), 5.5-5.4
(m, 1H), 5.2-4.8 (m, 2H), 3.0-2.9 (m, 2H), 2.2-2.0 (m, 2H),
1.4-1.1 (m, 14H), 0.9-0.7 (t, 3H).

Elemental analysis: responding to the expected formula
C₁₈H₃₂N₂ClO₅P.

MS-FAB+glycerol matrix=387.

K.F.=6% water.

HPLC: Spherisorb-C1 (5 μm, 250×4.6 mm), mobile
phase 0.050 M KH₂PO₄:CH₃CN 35:65, flow 0.75 ml/min, detec-
tor: RI, UV 205 nm, RT=5.61 min.

EXAMPLE 15

R-4-trimethylammonium-3-(tetradecylcarbonyl)
amino butyrate (ST 1326)

The product was prepared as disclosed in Example 1,
starting from tetradecyl isocyanate and R-aminocarnitine,

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inner salt, except the crude product was obtained by precipitation with ethyl ether, from the reaction mixture, directly washed with ethyl ether and purified on a silica gel chromatographic column.

Yield 57%.

M.p.: 160–162° C.

$[\alpha]_{20}^D = -21.1^\circ$ (c=0.5, MeOH).

¹H-NMR (300 MHz; CD₃OD):

δ: 4.52 (m, 1H), 3.60 (dd, 1H), 3.48 (d, 1H), 3.20 (s, 9H), 3.10 (t, 2H), 2.40 (m, 2H), 1.45 (m, 2H), 1.28 (brs, 22H), 0.8 (brt, 3H).

ESI Mass=400, [(M+H)⁺].

Elemental analysis: responding to the expected formula C₂₂H₄₅N₃O₃.

K.F.=2.5% water.

TLC silica gel CHCl₃:iPrOH:MeOH:H₂O:CH₃COOH 42:7:28:10.5:10.5.

Rf=0.50.

HPLC: SGE-SCX column (5 μm, 250×4 mm), T=30° C., mobile phase 0.05 M (NH₄)H₂PO₄:CH₃CN 75:25, pH=4.9 (as such), flow 0.75 ml/min, detector: RI, UV 205 nm RT=13.63 min.

EXAMPLE 16

R-4-trimethylammonium-3-(undecylcarbamoyl)-aminobutyrate (ST 1327)

The product was prepared as disclosed in Example 1, starting from undecyl isocyanate and R-aminocarnitine, inner salt, purified on a silica gel chromatographic column and further purified by precipitation from acetonitrile.

Yield 50%.

M.p.: 149–150.2° C.

$[\alpha]_{20}^D = -21.16^\circ$ (c=1, MeOH).

¹H-NMR (300 MHz; CD₃OD):

δ: 4.52 (m, 1H), 3.60 (dd, 1H), 3.48 (d, 1H), 3.20 (s, 9H), 3.10 (t, 2H), 2.40 (m, 2H), 1.45 (m, 2H), 1.28 (brs, 16H), 0.8 (brt, 3H).

ESI Mass=358, [(M+H)⁺];

Elemental analysis: responding to the expected formula C₁₉H₃₉N₃O₃.

K.F.=2.3% water.

TLC silica gel CHCl₃:iPrOH:MeOH:H₂O:CH₃COOH 42:7:28:10.5:10.5.

Rf=0.50.

HPLC: SGE-SCX column (5 μm, 250×4 mm), T=30° C., mobile phase 0.05 M (NH₄)H₂PO₄:CH₃CN 80:20, pH=4.9 (as such), flow 0.75 ml/min, detector: RI, UV 205 nm, RT=17.37 min.

EXAMPLE 17

R-4-trimethylammonium-3-(heptylcarbamoyl)-aminobutyrate (ST 1328)

The product was prepared as disclosed in Example 1, starting from heptyl isocyanate and R-aminocarnitine, inner salt, purified on a silica gel chromatographic column and further purified by precipitation from acetonitrile.

Yield 47%.

M.p.: 149–150° C.

$[\alpha]_{20}^D = -34.0^\circ$ (c=0.97, MeOH). ¹H-NMR (300 MHz; CD₃OD):

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δ: 4.52 (m, 1H), 3.60 (dd, 1H), 3.48 (d, 1H), 3.20 (s, 9H), 3.10 (t, 2H), 2.40 (m, 2H), 1.45 (m, 2H), 1.30 (brs, 8H), 0.8 (brt, 3H).

ESI Mass=302, [(M+H)⁺];

Elemental analysis: responding to the expected formula C₁₅H₃₁N₃O₃

K.F.=6.17% water

TLC silica gel CHCl₃:iPrOH:MeOH:H₂O:CH₃COOH 42:7:28:10.5:10.5.

Rf=0.50.

HPLC: SGE-SCX column (5 μm, 250×4 mm), T=30° C., mobile phase 0.05 M (NH₄)H₂PO₄:CH₃CN 85:15, pH=6 (H₃PO₄), flow 0.75 ml/min, detector: RI, UV 205 nm, RT=7.16 min.

EXAMPLE 18

R,S-4-trimethylammonium-3-(nonylthiocarbamoyl)-aminobutyrate (ST 1329)

The product was prepared as disclosed in Example 1, starting from nonyl isothiocyanate and R,S-aminocarnitine, inner salt. Chromatography was carried out with a CHCl₃/MeOH gradient from 8:2 to 2:8.

Yield 53%

M.p.: 104–107° C.

¹H-NMR (200 MHz; CD₃OD):

δ: 5.45 (brm, 1H), 3.75 (dd, 1H), 3.55(d, 1H), 3.45 (brm, 2H), 3.22 (s, 9H), 2.48 (m, 2H), 1.55 (m, 2H), 1.30 (brs, 12H), 0.90 (brt, 3H).

ESI Mass=346, [(M+H)⁺];

Elemental analysis: responding to the expected formula C₁₇H₃₅N₃O₂S

K.F.=2.6% water;

TLC silica gel CHCl₃:iPrOH:MeOH:H₂O:CH₃COOH 42:7:28:10.5:10.5.

Rf=0.74;

HPLC: SGE-SCX column (5 μm, 250×4 mm), T=30° C., mobile phase 0.05 M (NH₄)H₂PO₄:CH₃CN 85:15, pH=6.0 (H₃PO₄), flow 0.75 ml/min, detector: RI, UV 205 nm, RT=8.87 min.

EXAMPLE 19

R-4-trimethylammonium-3-(nonylcarbamoyl)-aminobutyrate (ST 1283)

The product was prepared as disclosed in Example 1, starting from nonyl isocyanate and R-aminocarnitine, inner salt.

M.p.: 146–147° C.

$[\alpha]_{20}^D = -13.4^\circ$ (c=0.5, H₂O).

Elemental analysis: responding to the expected formula C₁₇H₃₅N₃O₃

K.F.=2.8% water.

Remaining physico-chemical data were coincident with those of racemic ST1251 (Example 1).

EXAMPLE 20

S-4-trimethylammonium-3-(nonylcarbamoyl)-aminobutyrate (ST 1338)

The product was prepared as disclosed in Example 1, starting from nonyl isocyanate and S-aminocarnitine, inner salt.

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M.p.: 146–147° C.
 $[\alpha]_{20}^D = +16.7^\circ$ (c=0.43, H₂O).
¹H-NMR (300 MHz; CD₃OD):
 δ : 4.52 (m, 1H), 3.60 (dd, 1H), 3.45 (d, 1H), 3.18 (s, 9H),
 3.10 (t, 2H), 2.40 (m, 2H), 1.45 (m, 2H), 1.28 (brs, 12H),
 0.90 (brt, 3H).
 ESI Mass=330, [(M+H)⁺];
 Elemental analysis: responding to the expected formula
 C₁₇H₃₅N₃O₃
 K.F.=1.8% water.
 Remaining physico-chemical data were coincident with
 those of racemic ST1251 (Example 1).

EXAMPLE 21

S-4-trimethylammonium-3-(tetradecylcarbamoyl)-
aminobutyrate (ST 1340)

The product was prepared as disclosed in Example 1,
 starting from tetradecyl isocyanate and S-aminocarnitine,
 inner salt, except the crude product was obtained by precipi-
 tation with ethyl ether, from the reaction mixture, directly
 washed with ethyl ether and purified on a silica gel chro-
 matographic column.

Yield=57%;
 M.p.: 166–167° C.
 $[\alpha]_{20}^D = +20.7^\circ$ (c=0.5, MeOH).
 Elemental analysis: responding to the expected formula
 C₂₂H₄₅N₃O₃
 K.F.=1.7% water.
 Remaining physico-chemical data were coincident with
 those of racemic ST1326 (Example 15).

EXAMPLE 22

isobutyl R,S-4-trimethylammonium-3-
tetradecylamino-aminobutyrate (ST 1252)R,S-4-trimethylammonium-3-tetradecylamino-
aminobutyrate isobutyl Ester Acetate

Isobutyl ester of racemic aminocarnitine (5 g, 0.0198
 moles) and tetradecanal (4.6 g, 0.0217 moles) were dis-
 solved into 250 ml methanol. Glacial acetic acid (1.13 ml,
 0.198 moles) and 1 g of 10% Pd/C were added. The mixture
 was hydrogenated at 30 psi overnight. After filtration on
 celite, the solution was vacuum-concentrated. A pale yellow
 oil was obtained, which was purified through a silica gel
 column, eluting firstly with AcOEt, then AcOEt/MeOH 9:1.
 4 g of product were obtained.

Yield=47%;
 TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol
 7/water 10.5/acetic acid 10.5)/methyl acetate 7:3
 Rf=0.74.
¹H-NMR (300 MHz; CD₃OD):
 δ : 3.92-3.90 (d, 2H), 3.64-3.58 (m, 1H), 3.50-3.30 (m,
 2H), 2.80-2.50 (m, 4H), 2.0-1.9 (m, 1H), 2.6-2.4 (m, 2H),
 1.3 (s, 22H), 0.98-0.82 (m, 9H).

R,S-4-trimethylammonium-3-tetradecylamino-
aminobutyrate

The isobutyl ester of R,S-4-trimethylammonium-3-
 tetradecylamino-aminobutyric acid, acetate salt, (3.3 g) was
 hydrolyses on Amberlyst IRA 402 resin (OH⁻ activated
 form) and eluted with water. Water was evaporated to dry-
 ness under reduced pressure; the resulting white solid was

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washed with methanol, filtered and vacuum-dried. 1.95 g of
 product were obtained.

Yield 70%

M.p.=160° C. dec.

¹H-NMR (300 MHz; CD₃OD):

δ : 4.4 (m, 1H), 3.40-3.35 (m, 3H), 3.2 (s, 9H), 2.80-2.72
 (m, 1H), 2.56-2.42 (m, 2H), 2.27-2.16 (m, 1H), 1.55-1.40
 (m, 2H), 1.3 (s, 22H), 0.92-0.85 (t, 3H).

Elemental analysis: responding to the expected formula
 C₂₁H₄₄N₂O₂

K.F.=1.93% water.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropanol alcohol
 7/water 10.5/acetic acid 10.5)

Rf=0.5.

HPLC: SGE-SCX column (5 μ m, 250 \times 4 mm), mobile
 phase 0.05M (NH₄)H₂PO₄:CH₃CN 60:40, pH=4, flow=0.75
 ml/min; detector: RI, UV 205 nm, RT=30.017 min.

EXAMPLE 23

R,S-4-trimethylammonium-3-octylaminobutyrate
(ST 1254)R,S-4-trimethylammonium-3-
octylaminoaminobutyrate Isobutyl Ester Acetate

Isobutyl ester of racemic aminocarnitine chloride, (5 g,
 0.0198 moles) and octanaldehyde (2.79 g, 0.0217 moles)
 were dissolved into 250 ml methanol. Glacial acetic acid
 (1.13 ml, 0.198 moles) and 1 g 10% Pd/C were added. The
 mixture was hydrogenated at 30 psi overnight. After filtra-
 tion on celite, the solution was vacuum-concentrated. 8.5 g
 product was obtained, subsequently purified through a silica
 gel column, eluting firstly with AcOEt, then AcOEt/MeOH
 (9:1, 8.5:1.5). 3 g of product was obtained.

Yield=40%;

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol
 7/water 10.5/acetic acid 10.5)

Rf=0.54.

¹H-NMR (300 MHz; CD₃OD):

δ : 3.92-3.90 (d, 2H), 3.64-3.58 (m, 1H), 3.50-3.30 (m,
 2H), 2.80-2.50 (m, 4H), 2.0-1.9 (m, 1H), 2.6-2.4 (m, 2H),
 1.3 (s, 10H), 0.98-0.82 (m, 9H).

R,S-4-trimethylammonium-3- octylaminobutyrate

The isobutyl ester of R,S-4-trimethylammonium-3-
 tetradecylamino-aminobutyric acid, acetate salt, (2.8 g,
 0.00719) was hydrolysed on Amberlyst IRA 402 resin (OH⁻
 activated form) and eluted with water. Water was evaporated
 to dryness under reduced pressure; the resulting white solid
 was washed with methanol, filtered and vacuum-dried. 1.8 g
 of product were obtained.

Yield 70%

M.p.=140° C. dec.

¹H-NMR (300 MHz; CD₃OD):

δ : 3.42-3.30 (m, 3H), 3.2 (s, 9H), 2.85-2.70 (m, 1H), 2.60-
 2.40 (m, 2H), 2.30-2.20 (m, 1H), 1.55-1.40 (m, 2H), 1.3 (s,
 10H), 0.92-0.85 (t, 3H).

Elemental analysis: responding to the expected formula
 C₁₅H₃₂N₂O₂

K.F.=2.8% water.

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TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5)

Rf=0.32.

HPLC: SGE-SCX column (5 μm, 250×4 mm), mobile phase 0.05M (NH₄)H₂PO₄:CH₃CN 40:60, pH=4, flow=0.75 ml/min; detector: RI, UV 205 nm, RT=43.20 min.

EXAMPLE 24

R,S-4-trimethylammonium-3-(decansulfonyl) aminobutyrate (ST 1364)

Aminocarnitine Isobutyl Ester Chloride Hydrochloride

Isobutyl ester of aminocarnitine, inner salt (3 g, 18.72 mmoles), was dissolved in isobutanol (120 ml) and ice-bath cooled. Gaseous HCl was bubbled into the solution until complete saturation and clearing of the mixture. The solution was refluxed (bath temperature 130° C.) overnight. The solvent was vacuum-evaporated and the residue was triturated with Et₂O. 5.1 g of white solid were obtained.

Yield=95%;

¹H-NMR (200 MHz; D₂O):

δ: 4.3 (m, 1H), 4.0 (d, 2H), 3.8 (d, 2H), 3.2 (s, 9H), 3.1 (m, 2H), 2.0 (m, 1H), 0.9 (d, 6H).

Elemental analysis: responding to the expected formula C₁₁H₂₆Cl₂N₂O₂.

K.F.=1% water.

R,S-4-trimethylammonium-3-(decansulfonyl) aminobutyrate

The isobutyl ester of R,S-aminocarnitine chloride, hydrochloride (1 g, 3.46 mmoles) in anhydrous dichloromethane (5 ml) was added with triethylamine (2.65 ml, 19 mmoles) and decanesulfonyl chloride (2.1 g, 8.65 mmoles) suspended in 3 ml anhydrous dichloromethane, at 0° C. The mixture was left under stirring for 3 days at room temperature. The solvent was evaporated to dryness, the residue was taken up with ethyl acetate and the white precipitate of triethylamine hydrochloride was separated by from the solution by vacuum-filtration. The ethyl acetate solution was vacuum-dried to give 2.8 g of a yellow oil. 71 ml 1N NaOH were added to hydrolyze the isobutyl ester, leaving the suspension under stirring overnight at room temperature. The suspension was evaporated and vacuum-dried, and the solid residue was completely dried under oil-vacuum, taken up with methanol and purified through silica gel chromatographic column, using methanol as eluant. 555 mg of product were obtained.

Yield 44%

M.p.=158° C. dec.

¹H-NMR (300 MHz; CD₃OD):

δ: 4.3 (m, 1H), 3.45 (m, 2H), 3.25 (s, 9H), 3.15 (m, 2H), 2.45 (d, 2H), 1.8 (m, 2H), 1.45 (m, 2H), 1.4 (brs, 12H), 0.9 (brt, 3H).

Elemental analysis: responding to the expected formula C₁₇H₃₆N₂O₄S

Mass ESI =365 [(M+H)⁺], 387[(M+Na)⁺]

K.F.=3% water.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5)

Rf=0.62.

HPLC: Spherisorb-C₁ column (5 μm, 250×4.6 mm), mobile phase 0.05M K₂H₂PO₄:CH₃CN 35:65, pH as such,

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flow=0.73 ml/min; temperature=30° C., detector: RI, UV 205 nm, RT=7.0 min.

EXAMPLE 25

R,S-4-trimethylammonium-3-(nonylsulfamoyl) aminobutyrate (ST 1362)

The isobutyl ester of R,S-aminocarnitine chloride, hydrochloride (2 g, 6.9 mmoles) in anhydrous dichloromethane (40 ml) was added with triethylamine (3.8 ml, 27.6 mmoles) and dropped with SO₂Cl₂ in dichloromethane (1.7 ml in 10 ml final solution) at 0° C. The mixture was left under stirring for 3 days at room temperature, triethylamine (1.9 ml, 13.8 mmoles) and nonylamine (2.5 ml, 13.8 mmoles) were added and the reaction mixture was left under stirring overnight at room temperature. The solvent was vacuum-evaporated, the residue was taken up with ethyl acetate (100 ml) and the precipitate of triethylamine hydrochloride was separated from the solution by vacuum-filtration. The ethyl acetate solution was vacuum-dried to give 4.8 g of a yellow oil, to which were added 105 ml 1N NaOH to hydrolyze the isobutyl ester. The mixture was left under stirring overnight at room temperature and vacuum-dried. The residue was completely dried under oil-vacuum. The yellow semisolid was crystallized from chloroform. 1.26 g of product were obtained.

Yield 50%

M.p.=152° dec.

¹H-NMR (300 MHz; CD₃OD):

δ: 4.1 (m, 1H), 3.48 (d, 2H), 3.25 (s, 9H), 2.95 (m, 2H), 2.5 (t, 2H), 1.55 (t, 2H), 1.45 (brs, 12H), 0.9 (brt, 3H).

Elemental analysis: responding to the expected formula C₁₆H₃₅N₃O₄S

Mass ESI=366 [(M+H)⁺], 388[(M+Na)⁺]

K.F.=5.8% water.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5)

Rf=0.34.

HPLC: Spherisorb-C₁ column (5 μm, 250×4.6 mm), mobile phase 0.05M KH₂PO₄:CH₃CN 35:65, pH as such, flow=0.75 ml/min; temperature=30° C., detector: RI, UV 205 nm, RT=6.68 min.

EXAMPLE 26

S-4-trimethylammonium-3-(dodecansulfonyl) aminobutyrate (ST 1391)

The product was prepared as disclosed in Example 24, starting from isobutyl ester of S-aminocarnitine chloride, hydrochloride and dodecansulfonyl chloride, to give 600 mg of product.

Yield 44%

M.p.=156° C. dec.

[α]_D²⁰=+6° (c=0.245%, H₂O)

¹H-NMR (300 MHz; CD₃OD):

δ: 4.3 (m, 1H), 3.45 (m, 2H), 3.25 (s, 9H), 3.15 (m, 2H), 2.45 (d, 2H), 1.8 (m, 2H), 1.45 (m, 2H), 1.4 (brs, 16H), 0.9 (brt, 3H).

Elemental analysis: responding to the expected formula C₁₉H₄₀N₂O₄S

K.F.=8.6% water.

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TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5)

Rf=0.65.

HPLC: Spherisorb-C1 column (5 μm, 250×4.6 mm), mobile phase 0.05M KH₂PO₄:CH₃CN 40:60, pH as such, flow=0.75 ml/min; temperature=30° C., detector: RI, UV 205 nm, RT=8.5 min.

EXAMPLE 27

R-4-trimethylammonium-3-(dodecansulfonyl) aminobutyrate (ST 1420)

The product was prepared as disclosed in Example 24, starting from isobutyl ester of R-aminocarnitine chloride, hydrochloride and dodecansulfonyl chloride, to give 450 mg of product.

Yield 34%

M.p.=158° dec.

[α]_D²⁰=-7° (c=0.265%, H₂O)

¹H-NMR (300 MHz; CD₃OD):

δ: 4.3 (m, 1H), 3.45 (m, 2H), 3.28 (s, 9H), 3.15 (m, 2H), 2.45 (d, 2H), 1.8 (m, 2H), 1.45 (m, 2H), 1.3 (brs, 16H), 0.9 (brt, 3H).

Elemental analysis: responding to the expected formula C₁₉H₄₀N₂O₄S

K.F.=6.9% water.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5)

Rf=0.66.

HPLC: Spherisorb-C1 column (5 μm, 250×4.6 mm), mobile phase 0.05M KH₂PO₄:CH₃CN 40:60, pH as such, flow=0.75 ml/min; temperature=30° C., detector: RI, UV 205 nm, RT=8.11 min.

EXAMPLE 28

S-4-trimethylammonium-3-(undecylsulfamoyl) aminobutyrate (ST 1427)

The product was prepared as disclosed in Example 25, starting from isobutyl ester of S-aminocarnitine chloride, hydrochloride and undecyl amine, except the crude product was purified on a silica gel chromatographic column, using a gradient CHCl₃:MeOH 9:1 to 1:9. The product was further purified on a silica gel chromatographic column, using MeOH. 0.7 g of pure product was obtained.

Yield 38%

M.p.=153° C. dec.

[α]_D²⁰=+4° (c=0.25%, H₂O, pH=2)

¹H-NMR (300 MHz; CD₃OD):

δ: 4.1 (m, 1H), 3.48 (d, 2H), 3.25 (s, 9H), 2.95 (m, 2H), 2.5 (m, 2H), 1.55 (brt, 2H), 1.45 (brs, 16H), 0.9 (brt, 3H).

Elemental analysis: responding to the expected formula C₁₈H₃₉N₃O₄S

K.F.=2.9% water.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5)

Rf=0.68.

HPLC: Spherisorb-C1 column (5 μm, 250×4.6 mm), mobile phase 0.05M KH₂PO₄:CH₃CN 60:40, pH as such, flow=0.7 ml/min; temperature=30° C., detector: RI, UV 205 nm, RT=8.384 min.

EXAMPLE 29

R-4-trimethylammonium-3-(undecylsulfamoyl) aminobutyrate (ST 1428)

The product was prepared as disclosed in Example 25, starting from isobutyl ester of S-aminocarnitine chloride,

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hydrochloride and undecyl amine, except the crude product was purified on a silica gel chromatographic column, using a gradient CHCl₃: MeOH 9:1 to 1:9. The product was further purified on a silica gel chromatographic column, using MeOH. 0.5 g of product was obtained.

Yield 32%

M.p.=158° dec.

[α]_D²⁰=-4° (c=0.25%, H₂O, pH=2)

¹H-NMR (300 MHz; CD₃OD):

δ: 4.1 (m, 1H), 3.48 (d, 2H), 3.25 (s, 9H), 2.95 (m, 2H), 2.5 (m, 2H), 1.55 (brm, 2H), 1.45 (brs, 16H), 0.9 (brt, 3H).

Elemental analysis: responding to the expected formula C₁₈H₃₉N₃O₄S

K.F.=4.77% water.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5)

Rf=0.68.

HPLC: Spherisorb-C₁ column (5 μm, 250×4.6 mm), mobile phase 0.05M KH₂PO₄:CH₃CN 60:40, pH as such, flow=0.7 ml/min; temperature=30° C., detector: RI, UV 205 nm, RT=8.379 min.

EXAMPLE 30

R-4-trimethylammonium-3-(dodecylcarbamoyl) aminobutyrate (ST 1375)

The product was prepared as disclosed in Example 1, starting from R-aminocarnitine inner salt and dodecylisocyanate. The crude product obtained after washing with diethyl ether was purified on a silica gel chromatographic column to give 4.8 g of product.

Yield 55%

M.p.=147° C. dec.

[α]_D²⁰=-24.6° (c=0.48%, MeOH)

¹H-NMR (300 MHz; CD₃OD):

δ: 4.51 (m, 1H), 3.60 (dd, 1H), 3.45 (dd, 1H), 3.2 (s, 9H), 3.1 (t, 2H), 2.4 (m, 2H), 1.45 (m, 2H), 1.3 (brs, 18H), 0.9 (t, 3H).

Elemental analysis: responding to the expected formula C₂₀H₄₁N₃O₃

K.F.=5.4% water.

TLC silica gel (CHCl₃ 42/MeOH 28/isopropyl alcohol 7/water 10.5/acetic acid 10.5)

Rf=0.6.

HPLC: Spherisorb-C1 column (5 μm, 250×4.6 mm), mobile phase 0.05M KH₂PO₄:CH₃CN 65:35, pH=5.6, flow=0.75 ml/min; temperature=30° C., detector: RI, UV 205 nm, RT=8.5 min.

EXAMPLE 31

R-4-trimethylammonium-3-(10-phenoxydecylcarbamoyl)aminobutyrate (ST 1449)

10-Phenoxydecyl isocyanate

A solution of 11-phenoxyundecanoyl chloride (31.1 g, 104.8 mmoles) in acetone (30 ml) was dropped into a solution of sodium azide (9.53 g, 146.6 mmoles) in water (30 ml), cooled in an ice bath, keeping the solution temperature between 10 and 15° C. After one hour, the solution was transferred in a separatory funnel and the lower phase (the aqueous one) was eliminated. The higher phase was transferred into a flask containing 100 ml of toluene, previously

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warmed at 65° C. After 1.5 hours, the solution was evaporated to dryness, giving 13.37 g of crude product, which could be used as such in the subsequent reaction.

¹H-NMR (300 MHz; CDCl₃):

δ: 7.2 (m, 2H), 6.9 (m, 3H), 3.9 (t, 2H), 3.6 (t, 2H), 1.4 (m, 2H), 1.3 (m, 10H).

R-4-trimethylammonium-3-(10-phenoxydecylcarbamoyl)-amino butyrate

10-phenoxydecylisocyanate (25.0 g, 91.12 mmoles) was added to a solution of aminocarnitine, inner salts (7.3 g, 45.56 mmoles) in anhydrous DMSO (350 ml) and the solution was left to stand for 60 hours at 40° C. The resulting mixture was transferred in a 3 l Erlenmeyer flask containing ethyl ether (2.5 l) and the solvent was separated by decantation of the formed precipitate, which was then taken with few chloroform, transferred into a flask and precipitated again with ethyl ether. The so obtained crude product was washed several times with ethyl ether and purified on a silica gel chromatographic column, using a gradient CHCl₃: MeOH 9:1 to CHCl₃: MeOH 3:7 gradient until elution of impurities with higher R_f, then eluting the product of interest with MeOH only. 13.5 g of pure product were obtained.

Yield 68%

¹H-NMR (300 MHz; CD₃OD);

δ: 7.2 (m, 2H), 6.9 (m, 3H), 4.5 (m, 1H), 3.9 (t, 2H), 3.6 (dd, 1H), 3.4 (dd, 1H), 3.2 (s, 9H), 3.1 (t, 2H), 2.4 (m, 2H), 1.8 (m, 2H), 1.6 (m, 2H), 1.4 (m, 2H), 1.3 (m, 10H).

FAB Mass=436, [(M+H)⁺];

Elemental analysis: responding to the expected formula C₂₄H₄₁N₃O₄

K.F.=2.3% water.

EXAMPLE 32

R-4-trimethylammonium-3-(trans-β-styrenesulfonyl)aminobutyrate (ST 1448)

R-aminocarnitine Isobutyl Ester Chloride Hydrochloride

R-aminocarnitine inner salt (3 g, 18.72 mmoles) was dissolved in isobutanol (120 ml) and ice-bath cooled. Gaseous HCl was bubbled into the solution until complete saturation and clearing of the mixture. The solution was refluxed (bath temperature 130° C.) overnight. The solvent was vacuum-evaporated and the residue was triturated with Et₂O. 5.1 g of white solid were obtained.

Yield=95%;

¹H-NMR (200 MHz; D₂O):

δ: 4.3 (m, 1H), 4.0 (d, 2H), 3.8 (d, 2H), 3.2 (s, 9H), 3.1 (m, 2H), 2.0 (m, 1H), 0.9 (d, 6H).

Elemental analysis: responding to the expected formula C₁₁H₂₆Cl₂N₂O₂.

K.F.=1% water.

R-4-trimethylammonium-3-(trans-β-styrenesulfonyl)-aminobutyrate

The isobutyl ester of R-aminocarnitine chloride, hydrochloride (1 g, 3.46 mmoles) in anhydrous dichloromethane (5 ml) was added with triethylamine (2.65 ml, 19 mmoles) and trans-β-styrenesulfonyl chloride (1.753 g, 8.65 mmoles) suspended in 3 ml anhydrous dichloromethane, at 0° C. The mixture was left under stirring for 3 days at room temperature. The solvent was evaporated to dryness, the residue was

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taken up with ethyl acetate (100 ml) and the white precipitate of triethylamine hydrochloride was separated by from the solution by vacuum-filtration. The ethyl acetate solution was vacuum-dried, then 71 ml 1N NaOH were added to hydrolyze the isobutyl ester, leaving the suspension under stirring overnight at room temperature. The suspension was evaporated and vacuum-dried, and the solid residue was completely dried under oil-vacuum, taken up with methanol and purified through silica gel chromatographic column, using methanol as eluant. 565 mg of product were obtained.

Yield 50%

¹H-NMR (300 MHz; CD₃OD):

δ: 7.8 (d, 1H), 7.5 (m, 5H), 7.3 (d, 1H), 4.3 (m, 1H), 3.4 (m, 2H), 3.2 (s, 9H), 2.4 (d, 2H).

Elemental analysis: responding to the expected formula C₁₅H₂₂N₂O₄S

EST Mass=327 [(M+H)⁺]

Pharmacological Activity

Determination of CPT Inhibiting Activity

CPT inhibition was evaluated essentially as described in Kerner, J. & Bieber, L. L. (1990) Biochemistry 29: 4326-34 on fresh mitochondrial preparation obtained from normally fed Fischer rat liver or heart. Mitochondria were isolated from liver or heart and suspended in 75 mM saccharose buffer, 1 mM EGTA, pH 7.5. 100 μl mitochondrial suspension, containing 50 μM [¹⁴C] palmitoyl-CoA (specific activity 10,000 DPM/mole) and 10 mM L-carnitine, were incubated at 37° C., in the presence of scalar concentrations of the test product (0-3 mM). Reaction time: 1 minute.

Table 1 shows the IC₅₀ determined.

The compounds of the present invention have higher inhibiting activity than the one of the reference compound SDZ-CPI-975, Example 1, disclosed in EP 0 574 355.

TABLE 1

IC ₅₀ of inhibition CPTI curve in rat liver mitochondria	
Compound	IC ₅₀ (μM/I)
SDZ-CPI-975	17.4
ST1326	0.75
ST1327	3.2

Determination of Oleate-stimulated β-hydroxybutyrate Production

β-hydroxybutyrate production is an index of CPT activity. In fact, the production of ketone bodies, final products of mitochondrial β-oxidation, is related to CPT activity.

Mitochondrial preparations, obtained according to the method by Venerando et al. (Am. J. Physiol. 266:C455-C461, 1994), were used. Hepatocytes are incubated at 37° C. in KRB bicarbonate buffer at pH 7.4, 6 mM glucose, 1% BSA in O₂/CO₂ 95/5 atmosphere at 2.5×10⁶ cells/ml. After 40 min incubation with the test compound at different concentrations, the first set of samples was taken (T_{0 min}) and oleate was added (1 mM final in KRB+BSA 1.4%). After 20 minutes, the second withdrawal was made (T_{20 min})

Table 2 shows the results. The data are the mean of three different experiments, twice carried out.

The compounds of the present invention have higher β-hydroxybutyrate inhibiting activity than the one of the ref-

erence compound SDZ-CPI-975, Example 1, disclosed in EP 0 574 355.

TABLE 2

IC ₅₀ of inhibition CPTI curve of β -hydroxybutyrate production in rat hepatocytes	
Compound	IC ₅₀ (μ M/I)
SDZ-CPI-975	3.7
ST1251	0.5
ST1253	0.9
ST1285	1.9

Glucose and β -hydroxybutyrate in Serum Fasted Rats Treated With CPT Inhibitors

Normally fed Fischer rats were starved for 24 hours and subsequently treated with the test compounds. One hour after the treatment, the animals were sacrificed and serum concentrations of glucose and β -hydroxybutyrate were determined.

Table 3 shows the results. For the compound ST1326 were used doses of 14.5 mg/2 ml/kg, for other test compounds, the doses are equivalent to ST1326 one.

TABLE 3

β -hydroxybutyrate and glucose serum concentrations in 24 hours-starved rats, after one hour from intraperitoneal treatment.							
	SDZ						
	control	CPI-975	ST1251	ST1253	ST1326	ST1327	ST1328
β -OHB							
Mean	1867	119.9	99.8	118.8	133.1	93.0	169.2
s.c.	240	12.8	8.3	20.4	12.4	8.7	26.7
p<	—	0.001	0.001	0.001	0.001	0.001	0.001
Glu							
Mean	108.8	87.6	76.9	88.2	84.2	84.9	79.5
s.c.	6.7	1.0	2.3	3.9	2.4	1.6	1.6
p<	—	0.05	0.01	0.05	0.05	0.05	0.05

Glucose and Insulin Levels in Diabetic Animals Treated With CPT Inhibitors

C57BL/6J male rats, 5-weeks old, were provided by Charles River. After 10 days of acclimatisation in standard conditions (22 \pm 2 $^{\circ}$ C.; 55 \pm 15% humidity; 15–20/h air changes; 12 hours light-dark cycle, with 700–1900 lux) and with standard diet with 4RF21 feedstock (Mucedola), glycaemia was controlled in post-absorption state (starving from 8.30 a.m. to 4.30 p.m.). Blood withdrawal was carried out cutting the tail end. Glucose was analyzed in blood acid supernatant (HClO₄ 0.375 N) with autoanalyzer Cobas Mira S with Glucose GDH Kit (Roche).

The animals were divided in two groups, 26 mice each and fed with a high-fat and a low-fat diet, respectively.

After 2 months from the start of the diet, glycaemia was tested, according to the starting method. After about 3 months from the start of the diet, glycaemia was tested, according to the starting method and plasma insulin levels were also determined (with blood withdrawal from end tail cutting) using [125I] Rat Insulin Kit (Amersham).

One 10 mice group fed with low-fat diet and two 10-mice groups fed with the high-fat diet were selected. One of the

two high fat diet was administered with ST 1327 at the dose of 45 mg/Kg in deionised H₂O (p.o., twice a day, 8.30 a.m. and 5.30 p.m.). administration volume was 10 ml/Kg, the two remaining groups were treated with vehicle only. High-fat or low-fat diets were continued during the treatment.

After 20 days of treatment, glycaemia and plasma insulin were measured. After 43 days of treatment, the animals were sacrificed by decapitation in post-absorption state (fasting 8.30 a.m.–4.30 p.m.), 8 hours after the last treatment. Blood was withdrawn and serum was separated by centrifugation and stored at –80 $^{\circ}$ C. Liver, heart and skeletal muscle (upper limbs) were also extracted, frozen in dry ice-acetone and kept at –80 $^{\circ}$ C.

High-fat diet determined an increase of body weight, glycaemia and insulin, with respect to low-fat diet.

After 20 days of treatment with ST 1327, glucose and insulin levels significantly decreased.

Table 4 shows the results.

TABLE 4

Glucose and Insulin levels in rats fed with fat-rich diet.			
Compound	High Fat diet Control	High Fat diet Treated	Low Fat diet Control
Glucose mg/dl	248.5 \pm 11.03 (10)	181.4 \pm 9.63* (9)	207.3 \pm 6.84** (9)
Insulin ng/ml	1.632 \pm 0.246 (10)	0.621 \pm 0.117** (9)	0.549 \pm 0.050* (9)

Student's t test, * and ** indicate p<0.001 and p<0.01, respectively, against high fat diet; () indicates the number of cases.

These results shows that the compounds, according to the present invention are effective in controlling glycaemia in fasting conditions. This is an important aspect in the treatment of diabetes, wherein hepatic gluconeogenesis occurs during fasting periods (i.e. nocturnal rest).

The Effect of CPT Inhibitors on Myocardial Ischemia

The compounds of the present invention are also effective in the treatment of ischemia, in particular myocardial ischemia.

To this end, male Wistar rats, weighing 200–225 g, provided by Charles-River, were kept at constant temperature of 23 $^{\circ}$ +/–1 $^{\circ}$ C., 50+/–10% relative humidity, 12 hours light-dark cycle, fed with pellet 4RF21 (Mucedola) tap water ad libitum.

The animals were anaesthetised with sodium Pentobarbital at the dose of 70 mg/Kg i.p.. Hearts were rapidly removed and put in a cold Krebs-Henseleit solution, before incannulation of aorta e subsequent perfusion according to Langendorff technique at 37 $^{\circ}$ C. with a pressure of 100 cm water.

Perfusion medium (Krebs-Henseleit) at pH 7.4 consists in: 128 mM NaCl, 4.7 mM KCl, 1 mM MgCl₂, 0.4 mM Na₂HP0₄, 20.2 mM NaHCO₃, 1.3 mM CaCl₂, 5 mM glucose. The medium was constantly oxygenated with carbogen (95% O₂, 5% CO₂).

After a 10 min “conditioning” period, hearts were perfused in a recirculant apparatus for 20 min. with the same medium containing 0.6 mM palmitate complexed with albumine (fraction V, fatty acid free), with or without the CPT inhibitor according to the present invention. By way of example ST 1364 was used at concentrations of 1 and 5 μ M.

After such a period ischemia was induced by reducing perfusion hydrostatic pressure from 100 cm to 20 cm for a period of 30 min. Reperfusion was started re-establishing the starting pressure conditions (100 cm). Hearts were controlled for 20 min. the inhibitor is present also during reperfusion phase.

Lactate dehydrogenases (LDH) release was monitored in the effluent in normal oxygenation conditions, during ischemia, with a withdrawal of medium at 30', and during reperfusion, with withdrawals at 1.5, 10, 15 and 20 minutes.

LDH release in the effluent is remarkably reduced, during reperfusion results significantly reduced in the presence of ST1364 at the dose of 5 μ M (FIG. 1). This result indicates a lower entity of cellular damage from reperfusion of the treated with respect to the controls.

Statistical analysis was carried out with Student's "t" test for non-paired data.

The number of the cases for each group is six (n=6).

The following Table 5 reports the results.

TABLE 5

	LDH release in perfusate (mU/ml/min)		
	Control	ST1364 1 μ M*	ST1364 5 μ M**
Basal	280	275	220
Ischemia 30'	200	220	200
Reperfusion 1'	640	480	410
Reperfusion 5'	660	500	380
Reperfusion 10'	670	495	380
Reperfusion 15'	700	510	320
Reperfusion 20'	720	580	325

Statistical analysis was carried out with Student's "t" test for non-paired data. *p<0.05 vs controls; **p<0.01 vs controls.

The number of the cases for each group is six (n=6).

LDH release in the effluent is remarkably reduced, during reperfusion results significantly reduced in the presence of ST1364 at the dose of 5 μ M (FIG. 1). This result indicates a lower entity of cellular damage from reperfusion of the treated with respect to the controls.

In another aspect, the present invention provides a combination of at least a compound of formula (I) with at least another active ingredient suitable for the treatment of the disease of interest.

In the treatment or prevention of diabetes, the present invention provides a compound of formula (I), optionally in combination with a suitable well-known active ingredient, such as for example a sulfonyleurea, L-carnitine, fibrates and other agonists of peroxisomal proliferator activated receptor (PPAR- α), agonists of 9-cis retinoic acid activated receptor, such as RXR, in particular α , β - and γ -isoforms, HMG-CoA reductase inhibitor, β -sitosterol inhibitor, cholesterol acyltransferase inhibitor, biguanides, cholestyramine, angiotensin II antagonist, melinamide, nicotinic acid, fibrinogen receptor antagonists, aspirin, α -glucosidase inhibitors, insulin secretagogue, insulin and glucagon-like peptides (incretins) and agonists of PPAR- γ (such as thiazolidinediones or others).

In the treatment or prevention of obesity, the present invention provides a compound of formula (I), optionally in combination with a suitable well-known active ingredient, such as for example fenfluramine, dexfenfluramine, phentiramine, a β -3-adrenergic receptor agonist.

In the treatment or prevention of high triglyceridemia, the present invention provides a compound of formula (I), optionally in combination with a suitable well-known active ingredient.

The compounds according to the present invention are also useful in the treatment or prevention of high cholesterol levels and in modulating HDL plasma levels, thus resulting beneficial in the treatment or prevention of the diseases related with these altered plasma levels. Examples of related diseases are hypertension, obesity, atherosclerosis, diabetes and related conditions. The medicaments containing at least a compound of the present invention may contain in combination at least another active ingredient effective in the treatment or prevention of the above mentioned diseases. Examples of other active ingredient are fibrates, such as clofibrate, benzafibrate and gemfibrozil and other PPAR- α agonists; inhibitors of cholesterol biosynthesis, such as HMG-CoA reductase inhibitors, such as statins, namely lovastatin, simvastatin and pravastatin; inhibitors of cholesterol absorption for example beta-sitosterol and (acyl CoA:cholesterol acyltransferase) inhibitors for example melinamide; anion exchange resins for example cholestyramine, colestipol or a dialkylaminoalkyl derivatives of a cross-linked dextran; nicotiny alcohol, nicotinic acid or a salt thereof; vitamin E; thyromimetics and L-carnitine.

The compounds of the present invention may be orally administered in the form of a pharmaceutical composition, comprising a therapeutically effective amount of at least a compound of formula (I) in admixture with a pharmaceutically acceptable vehicle and/or excipient. Examples of oral pharmaceutical compositions are hard or soft capsules, tablets, including sublingual administration, ampoules, sachets, elixirs, suspensions, syrups, and the like. Alternatively, the active ingredients according to the present invention may be incorporated directly with the food of the diet. The amount of active compound in such therapeutically useful compositions is such that an effective dosage will be obtained. The active compounds can also be administered intranasally as, for example, liquid drops or spray.

The tablets, pills, capsules, and the like may also contain a binder such as gum tragacanth, acacia, corn starch or gelatin; excipients such as dicalcium phosphate; a disintegrating agent such as corn starch, potato starch, alginic acid; a lubricant such as magnesium stearate; and a sweetening agent such as sucrose, lactose or saccharin. When a dosage unit form is a capsule, it may contain, in addition to materials of the above type, a liquid carrier such as a fatty oil.

Various other materials may be present as coatings or to modify the physical form of the dosage unit. For instance, tablets may be coated with shellac, sugar or both. A syrup or elixir may contain, in addition to the active ingredient, sucrose as a sweetening agent, methyl and propylparabens as preservatives, a dye and a flavoring such as cherry or orange flavor.

These active compounds may also be administered parenterally. Solutions or suspensions of these active compounds can be prepared in pyrogen-free water.

The pharmaceutical forms suitable for injectable use include sterile aqueous solutions or dispersions and sterile powders for the extemporaneous preparation of sterile injectable solutions or dispersions.

If desired, or deemed necessary, the pharmaceutical compositions may be in the controlled-release form. Various techniques for preparing these forms are known.

General reference for pharmaceutical compositions can be made to "Remington's Pharmaceutical Sciences Handbook", Mack Pub. N.Y. USA.

The effective dosage of active ingredient employed may vary depending on the particular compound employed, the mode of administration, the condition being treated and the severity of the condition being treated.

The compositions are formulated and administered in the same general manner as detailed below. The compounds of the present invention can be used effectively alone or in combination with one or more additional active agents depending on the desired target therapy. Combination therapy includes administration of a single pharmaceutical dosage formulation which contains a compound of formula I and one or more additional active agents, as well as administration of a compound of formula I and each active agent in its own separate pharmaceutical dosage formulation. For example, a compound of formula I and an HMG-CoA reductase inhibitor can be administered to the patient together in a single oral dosage composition such as a tablet or capsule, or each agent administered in separate oral dosage formulations. Where separate dosage formulations are used, a compound of formula I and one or more additional active agents can be administered at essentially the same time, i.e., concurrently, or sequentially; combination therapy is understood to include all these regimens.

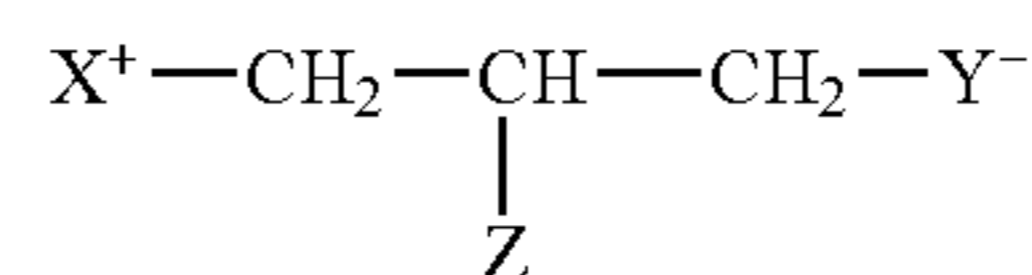
An example of combination treatment or prevention of atherosclerosis is wherein a compound of formula I is administered in combination with one or more of the following active agents: an antihyperlipidemic agent; a plasma HDL-raising agent; an antihypercholesterolemic agent such as a cholesterol biosynthesis inhibitor, for example an HMG-CoA reductase inhibitor, an HMG-CoA synthase inhibitor, a squalene epoxidase inhibitor, or a squalene synthetase inhibitor (also known as squalene synthase inhibitor); an acyl-coenzyme A: cholesterol acyltransferase (ACAT) inhibitor such as melinamide; probucol; nicotinic acid and the salts thereof and niacinamide; a cholesterol absorption inhibitor such as beta-sitosterol; a bile acid sequestrant anion exchange resin such as cholestyramine, colestipol or dialkylaminoalkyl derivatives of a cross-linked dextran; an LDL (lower density lipoprotein) receptor inducer; fibrates such as clofibrate, bezafibrate, fenofibrate, and gemfibrozil and other PPAR- α agonists, L-carnitine; vitamin B₆ and the pharmaceutically acceptable salt thereof; vitamin B₁₂; anti-oxidant vitamins such as vitamin C and E and beta carotene; a beta-blocker; an antihypertensive agent; an angiotensin converting enzyme inhibitor; and a platelet aggregation inhibitor such as fibrinogen receptor antagonists (i.e., glycoprotein IIb/IIIa fibrinogen receptor antagonists) and aspirin. The compounds of formula I can be administered in combination with more than one additional active agent.

Another example of combination therapy can be seen in treating obesity or obesity-related disorders, wherein the compounds of formula I may be effectively used in combination with for example, fenfluramine, dexfenfluramine, phentramine and β -3 adrenergic receptor agonist agents and L-carnitine.

Another example of combination therapy can be seen in treating diabetes and related disorders wherein the compounds of formula I can be effectively used in combination with for example sulfonylureas, biguanides, α -glucosidase inhibitors, other insulin secretagogues, insulin and glucagon-like peptides (incretins) and agonists of PPAR- γ (such as thiazolidinediones or others) as well as the active agents discussed above for treating atherosclerosis.

What is claimed is:

1. A compound of formula (I)



wherein: X⁺ is P⁺(R₁,R₂,R₃), wherein R₁,R₂,R₃, being the same or different, are selected in the group consisting of hydrogen, a C₁-C₉ straight or branched alkyl group, —CH=NH(NH₂), —NH₂, and —OH; or one or more R₁, R₂ and R₃, together with the phosphorus atom which they are linked to, form a saturated or unsaturated, monocyclic or bicyclic heterocyclic system; with the proviso that at least one of the R₁, R₂ and R₃ is different from hydrogen;

Z is selected from:

—OR₄,
—OCOOR₄,
—OCONHR₄,
—OCSNHR₄,
—OCSOR₄,
—NHR₄,
—NHCOR₄,
—NHCSR₄,
—NHCOOR₄,
—NHCSOR₄,
—NHCONHR₄,
—NHCSNHR₄,
—NHSOR₄,
—NHSONHR₄,
—NHSO₂R₄,
—NHSO₂NHR₄, and
—SR₄,

wherein —R₄ is a C₁-C₂₀ saturated or unsaturated, straight or branched alkyl group, optionally substituted with an A₁ group, wherein A₁ is selected from the group consisting of a halogen atom, or an aryl, heteroaryl, aryloxy or heteroaryloxy group, said aryl, heteroaryl, aryloxy or heteroaryloxy groups being optionally substituted with one or more C₁-C₂₀ saturated or unsaturated, straight or branched alkyl or alkoxy group and/or halogen atom;

Y⁻ is selected from the group consisting of —COO⁻, PO₃H⁻, —OPO₃H⁻, tetrazolate-5-yl;

with the proviso that when Z is —NHCOR₄, Y is —COO⁻, then R₄ is C₂₀ alkyl; and

with the proviso that when Z is —NHSO₂R₄, Y⁻ is —COO⁻, then R₄ is not tolyl;

their (R,S) racemic mixtures, their single R or S enantiomers, or their pharmaceutically acceptable salts.

2. A compounds according to claim 1, wherein R₁, R₂ and R₃ are methyl.

3. A compound according to claim 1, wherein R₁ and R₂ are H, R₃ is selected from the group consisting of —CH=NH(NH₂), —NH₂ and —OH.

4. A compound according to claim 1, wherein Z is selected from the group consisting of ureido (—NHCONHR₄) or carbamate (—OCONHR₄), and R₄ is a C₇-C₂₀ saturated or unsaturated, straight or branched alkyl group.

5. A compound according to claim 4, wherein R₄ is a C₉-C₁₈ saturated or unsaturated, straight or branched alkyl group.

6. [A] The compound [selected from the group consisting of] [R,S-4-trimethylammonium-3-(nonylcarbamoyl)-aminobutyrate;

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R,S-4-quinuclidinium-3-(tetradecyloxycarbonyl)-oxybutyrate;
 R,S-4-trimethylammonium-3-(nonylcarbamoyl)-oxybutyrate;
 R,S-4-trimethylammonium-3-(nonyloxycarbonyl)-oxybutyric acid chloride;]
 R,S-4-trimethylphosphonium-3-(nonylcarbamoyl)-oxybutyrate];
 R,S-4-trimethylammonium-3-(octyloxycarbonyl)-aminobutyrate;
 R,S-4-trimethylammonium-3-(nonyloxycarbonyl)-aminobutyrate;
 R,S-4-trimethylammonium-3-octyloxybutyrate;
 R,S-4-trimethylammonium-3-tetradecyloxybutyrate;
 R-4-trimethylammonium-3-(10-phenoxydecylcarbamoyl)amino-butyrate; and
 R-4-trimethylammonium-3-(trans- β -styrenesulfonyl)aminobutyrate].

7. A process for the preparation of a compound of claim 1 wherein Z is carbonate ($-\text{OCOOR}_4$), carbamate ($-\text{NHCOOR}_4$), thiocarbamate ($-\text{OCSNHR}_4$) or thiocarbonate ($-\text{OCSOR}_4$), said process comprising reacting $\text{X}^+-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{Y}-$, of the desired structure, optionally protected on the acid Y- group, respectively with an alkyl chloroformate, alkyl isocyanate, alkyl isothiocyanate or alkyl thiochloroformate, wherein the alkyl moiety is the desired R_4 alkyl group, to produce the desired compound.

8. A process for a preparation of a compound of claim 1 wherein Z is [amide] amide ($-\text{NHCOR}_4$), [thioamide] thioamide ($-\text{NHCSR}_4$), carbamate ($-\text{NHCOOR}_4$), thiocarbamate ($-\text{NHCSOR}_4$), ureido ($-\text{NHCONHR}_4$), thioureido ($-\text{NHCSNHR}_4$), sulfinamide ($-\text{NHSOR}_4$), sulfonamide ($-\text{NHSO}_2\text{R}_4$), sulfinamoylamino ($-\text{NHSONHR}_4$), and sulfamide ($-\text{NHSO}_2\text{NHR}_4$), said process comprising reacting $\text{X}^+-\text{CH}_2-\text{CH}(\text{OH})-\text{CH}_2-\text{Y}-$, of the desired structure, optionally protected on the acid Y- group, respectively with an acyl chloride, thioacyl chloride, alkyl chloroformate, alkyl thiochloroformate, alkyl isocyanate, alkyl thioisocyanate, alkyl sulfinyl chlorides, alkyl sulfonyl chlorides, SOCl_2 and alkyl amines, alkyl sulfamoyl chloride or SOCl_2 and alkyl amine, wherein the alkyl moiety is the desired R_4 alkyl group, to produce the desired compound.

9. A process for the preparation of a compound of claim 1, wherein Z is $-\text{OR}_4$ or $-\text{SR}_4$, said process comprising the steps of:

- (a) reacting a carbonyl compound of formula $\text{Hal}-\text{CH}_2-\text{CO}-\text{CH}_2-\text{COOR}'$, wherein Hal is a halogen atom and R' is the residue of a suitable ester, with respectively alcohols and thiols R_4OH or R_4SH , to give the respective ketal or thioketal;
- (b) transforming the respective ketal or thioketal into the respective ether or thioether;
- (c) substituting the Hal atom with an azido group, and
- (d) transforming the azido group into the X^+ group to produce the desired compound.

10. A process for the preparation of a compound of claim 1, wherein Z is $-\text{NHR}_4$, said process comprising reacting of

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$\text{X}^+-\text{CH}_2-\text{CH}(\text{NH}_2)-\text{CH}_2-\text{Y}-$ of the desired structure, optionally protected on the acid Y- group, with alkane carbaldehydes, wherein the alkyl moiety is a one-term lower homologue of the desired R_4 , and subsequent reduction, to produce the desired compound.

11. A pharmaceutical composition comprising a therapeutically effective amount of a compound of claim 1, in admixture with a pharmaceutically acceptable vehicle or and excipient.

12. The pharmaceutical composition [according to claim 11, wherein an active ingredient suitable] for the treatment of diabetes [is also present and is selected from the group consisting of sulfonylurea, L-carnitine, fibrate and other agonists of peroxisomal proliferator activated receptor (PPAR- α), HMG-CoA reductase inhibitor, β -sitosterol inhibitor, cholesterol acyltransferase inhibitor, biguanides, cholestyramine, angiotensin II antagonist, melinamide, nicotinic acid, fibrinogen receptor antagonists, aspirin, α -glucosidase inhibitors, insulin secretagogue, insulin and glucagon-like peptides and agonists of PPAR- γ] comprising therapeutically effective amounts of R-4-trimethylphosphonium-3-(tetradecylcarbamoyl)-amino butyrate and metformin, in admixture with a pharmaceutically acceptable vehicle or an excipient.

13. A pharmaceutical composition according to claim 11, also including an active ingredient suitable for the treatment of obesity selected from the group consisting of fenfluramine, dexfenfluramine, phentriamine, and a β -3-adrenergic receptor agonist.]

14. A pharmaceutical composition according to claim 11, also including an active ingredient suitable for the treatment of high cholesterol levels and in modulating HDL plasma levels, which is selected from the group consisting of fibrates, and other PPAR- α agonists; inhibitors of cholesterol biosynthesis, HMG-CoA reductase inhibitors, statins, inhibitors of cholesterol absorption, acyl CoA:cholesterol acyltransferase inhibitors, anion exchange resins, nicotinic alcohol, nicotinic acid or a salt thereof, vitamin E, thyromimetics and L-carnitine.]

15. A method for treating a subject having hyperactive carnitine palmitoyl-transferase comprising administering to said subject an effective amount of a compound of claim 1.]

16. A method for treating a subject having hyperglycaemia, diabetes, heart failure or ischemia comprising administering to said subject an effective amount of a compound of claim 1.

17. A method for treating a subject having obesity comprising administering to said subject an effective amount of a compound of claim 1.]

18. A method for treating a subject having high triglyceridemia comprising administering to said subject an effective amount of a compound of claim 1.]

19. A method for treating a subject a subject having hypertension comprising administering to said subject an effective amount of a compound of claim 1.]

20. A method of modulating high cholesterol levels or high density lipoprotein plasma levels in a subject in need of same, said method comprising administering to said subject an effective amount of a compound of claim 1.]

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