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Substituted a-amino acids, their preparation and pharmaceutical compositions containing them

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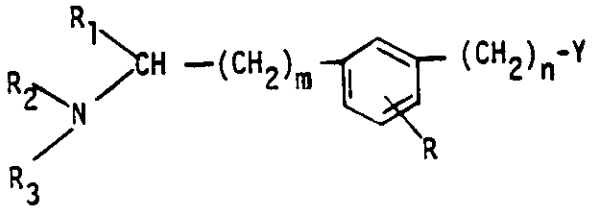
Substituted α-aminoacids, their preparation and pharmaceutical compositions containing them

The present invention relates to α-amino-α-(3-alkylphenyl)alkyl-ethanoic acids, esters or amides, in which the 3-alkyl moiety bears a phosphorus oxo acid group or an ester thereof, wherein phosphorus is attached directly to the alkyl moiety, their salts, processes for their production, pharmaceutical compositions containing them and their use as pharmaceuticals.

It is to be appreciated that the compounds of the invention may be optionally substituted. In particular the phenyl group may be further substituted. Examples of substituents in the phenyl ring are alkoxy, phenyl or phenyl substituted by e.g. halogen, alkyl or phenyl. Furthermore the α-amino group may bear substituents.

In a preferred aspect the present invention relates to compounds of formula I



I

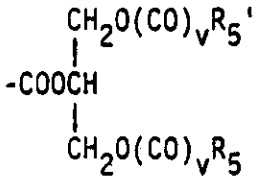
wherein

m and n are independently 1 or 2,

R_1 is carboxy, (C_{1-12})alkoxycarbonyl, benzoyl(C_{1-4})alkoxycarbonyl, phenyl(C_{2-4})alkenyloxycarbonyl, carbamoyl, monoalkyl(C_{1-6})carbamoyl or dialkyl(C_{1-6})carbamoyl,

R_2 is hydrogen or (C_{1-12})alkyl,

R_3 is hydrogen, (C_{1-12})alkyl, (C_{1-18})alkylcarbonyl, (C_{2-22})-alkenylcarbonyl, (C_{4-22})alkadienylcarbonyl, (C_{6-22})alkatrienylcarbonyl, (C_{8-22})alkatetraenylcarbonyl, (C_{1-12})alkoxycarbonyl or a group of formula II,

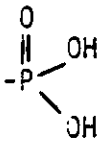


II

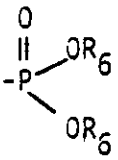
in which R_5 and R_5' are each, independently (C_{1-22})alkyl, (C_{2-22})-alkenyl, (C_{4-22})alkadienyl, (C_{6-22})alkatrienyl, (C_{8-22})alkatetraenyl and v is independently of each other 0 or 1,

R is hydrogen, halogen, hydroxy, (C_{1-12})alkyl, (C_{1-12})alkoxy, phenyl, phenyl(C_{1-8})alkoxy, phenyl(C_{1-8})alkyl; phenyl substituted by halogen, (C_{1-12})alkyl, (C_{1-12})alkoxy, amino, (C_{1-12})-alkylcarbonylamino, hydroxy or phenyl,

Y is one of the groups a), b), c) or d)



a)



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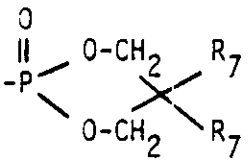
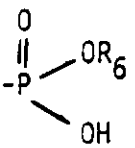


c)



d)

wherein

R_6 is (C_{1-6})alkyl and

R_7 is hydrogen or (C_{1-6})alkyl,

or a salt thereof.

Salts of the compounds of the inventior are on the one hand metal or ammonium salts of the compounds of the invention having a free carboxy or a free phosphorus oxo acid group, more particularly alkali or alkaline earth metal salts, e.g. sodium, potassium or magnesium salt, and ammonium salts derived from ammonia or organic amines. On the other hand, when a basic nitrogen atom is present the compounds of the invention can form acid addition salts of inorganic or organic acids, e.g. hydrochloric, hydrobromic or maleic acid.

It will be appreciated that the compounds of the invention contain a chiral centre at the carbon atom bearing the α-amino group and can therefore exist in racemic and optically-active forms. It is to be understood that the present invention encompasses the racemic and any optically-active form. Wherein alkenyl groups are present, stereoisomeric forms occur. These isomers are also included within the scope of the present invention.

In one group of compounds of formula I
m and n are independently 1 or 2,

R_1 is carboxy or (C_{1-12})alkoxycarbonyl,

R_2 is hydrogen or (C_{1-12})alkyl,

R_3 is hydrogen, (C_{1-12})alkyl, (C_{1-12})alkylcarbonyl, (C_{2-22})-alkenylcarbonyl, (C_{4-22})alkadienylcarbonyl, (C_{6-22})alkatrienyl-carbonyl, (C_{8-22})alkatetraenylcarbonyl, (C_{1-12})alkoxycarbonyl

or a group of formula II, in which R_5 and R_5' are each, independently (C_{1-22})alkyl, (C_{2-22})alkenyl, (C_{4-22})alkadienyl, (C_{6-22})alkatrienyl, (C_{8-22})alkatetraenyl and v is independently of each other 0 or 1,

R is hydrogen, halogen, hydroxy, (C_{1-12})alkyl, (C_{1-12})alkoxy, phenyl; phenyl(C_{1-8})alkoxy, phenyl(C_{1-8})alkyl, phenyl substituted by halogen, (C_{1-12})alkyl, (C_{1-12})alkoxy or phenyl,

Y is one of the groups a), b) or c),

wherein

R_6 is (C_{1-6})alkyl and
R_7 is hydrogen or (C_{1-6})alkyl,

or a salt thereof.

In another group of compounds of formula I, m and n are independently 1 or 2, R_1 is carboxy, (C_{1-12})alkoxycarbonyl, benzoyl(C_{1-4})alkoxycarbonyl, phenyl(C_{2-4})alkenyloxycarbonyl or carbamoyl, R_2 is hydrogen, R_3 is hydrogen or (C_{1-18})alkylcarbonyl, R is hydrogen, (C_{1-12})alkoxy, phenyl, phenyl substituted by halogen, (C_{1-12})alkyl, amino or phenyl, Y is one of the groups a), b), c) or d), wherein R_6 is (C_{1-6})alkyl and R_7 is hydrogen or (C_{1-6})alkyl, or a salt thereof.

In the above formula I, the following significances as well as combinations thereof, are preferred:

m is 1.

n is 1.

R_1 is carboxy or (C_{1-4})alkoxycarbonyl.

R_2 is hydrogen.

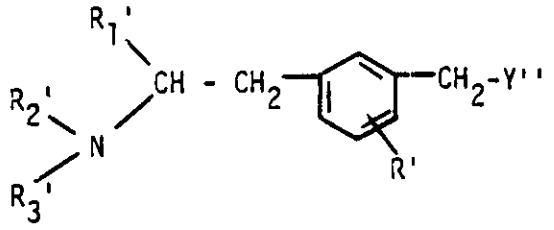
R_3 is hydrogen or (C_{1-18})alkylcarbonyl.

R is (C_{1-12})alkoxy, phenyl or phenyl substituted by halogen, (C_{1-4})alkyl or phenyl.

Y is group a) or group b), wherein R_6 is (C_{1-4})alkyl or group c), wherein R_7 is (C_{1-4})alkyl, especially group a).

Halogen is preferably chlorine or fluorine and especially chlorine.

A preferred group of compounds of formula I are compounds of formula Ia

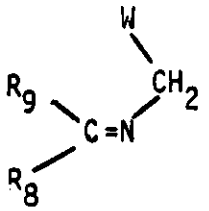


Ia

wherein R_1' is carboxy, (C_{1-4})alkoxycarbonyl, benzoyl(C_{1-4})alkoxycarbonyl, phenyl(C_{2-4})alkenyloxycarbonyl or carbamoyl, R_2' is hydrogen, R_3' is hydrogen or (C_{1-18})alkylcarbonyl, R' is (C_{1-12})-alkoxy, phenyl or phenyl substituted by halogen, (C_{1-12})alkyl, amino or phenyl, Y'' is one of the groups a), b) or c), wherein R_6 is (C_{1-6})alkyl and R_7 is hydrogen or (C_{1-6})alkyl, or a salt thereof.

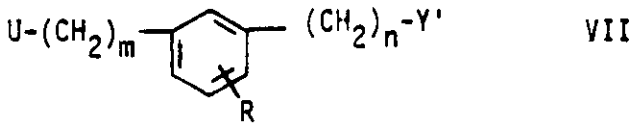
The present invention in another aspect provides a process for the production of a compound of the invention which comprises reacting a protected glycine derivative with an appropriate 1-alkyl-3-alkyl-benzene , in which one alkyl moiety bears a phosphorus oxo acid ester group wherein phosphorus is attached directly to the alkyl moiety and the other alkyl group bears a leaving group, under basic conditions and hydrolysing the resulting compound, and, if desired, converting a resulting compound of the invention into another compound of the invention, and/or, if desired, converting a resulting free compound into a salt; and/or, if desired, resolving a racemate obtained into the optical antipodes.

In particular a compound of formula I as defined above can be produced by reacting a compound of formula VI



VI

wherein R_8 is hydrogen, alkyl or phenyl, R_9 is phenyl optionally substituted by chlorine, alkyl or alkoxy, and W is -CN or $-COOR_{10}$, wherein R_{10} is an ester forming radical, with a compound of formula VII,



VII

wherein m, n and R are as defined above, U is a leaving group, and Y' is one of the groups b) or c), under basic conditions, and hydrolysing the resulting compound, and, if desired, converting a resulting compound of formula I into another compound of formula I; and/or, if desired, converting a resulting free compound into a salt; and/or, if desired, resolving a racemate obtained into the optical antipodes.

The process can be effected in conventional manner. Suitable protected glycine derivative are Schiff bases derived from glycine ester or glycinonitrile, in particular a compound of formula VI. R_{10} is e.g. alkyl or phenylalkyl. Preferably R_8 and R_9 are each phenyl. The reaction of a protected glycine derivative, in particular of a compound of formula VI with a 1-alkyl-3-alkylbenzene, in which one alkyl moiety bears a phosphorus oxo acid ester group wherein phosphorus is attached directly to the alkyl moiety and the other alkyl group bears a leaving group, in particular a compound of formula VII, wherein leaving group U is e.g. halogen, especially bromine, methylsulfonyloxy or p-methylphenylsulfonyloxy, can for example be carried out in a basic two phase system, e.g. a water-immiscible solvent such as dichlormethane and solid or aqueous sodium hydroxide using a phase-transfer catalyst, e.g. benzyl-tributylammonium chloride. Suitable temperatures range from 0°

to room temperature. Alternatively the reaction can also be carried out in an anhydrous organic solvent, such as toluene in the presence of e.g. sodium ethoxide or sodium methoxide at a temperature between 40° and 110°C. The reaction can also be carried out in a water-miscible organic solvent such as dioxane in the presence of an aqueous solution of benzyltrimethylammonium hydroxide at room temperature. The resulting alkylated Schiff base can be hydrolysed to the corresponding α-amino acid in conventional manner, e.g. with hydrochloric acid. Suitably in compounds of formula VI W is $COOR_{10}$, wherein R_{10} is alkyl, when compounds of formula I are to be prepared, wherein R_1 is esterified carboxy, e.g. alkoxycarbonyl. Using mild reaction conditions for the hydrolysis of the alkylated Schiff base, e.g. dilute hydrochloric acid at room temperature, only the imine functionality is selectively hydrolysed to yield compounds of formula I, wherein R_1 is esterified carboxy and Y is a group b) or c). Concentrated hydrochloric acid at elevated temperature leads to compounds of formula I, wherein R_1 is carboxy and Y is group a). Compounds of formula VI, wherein W is CN, are suitably used, when compounds of formula I are to be prepared, wherein R_1 is carboxy.

Compounds of formula VI, wherein W is $COOR_{10}$ are conveniently employed when compounds of formula I, wherein R_1 is carbamoyl, alkylcarbamoyl or dialkylcarbamoyl, are to be prepared. In this case, the hydrolysis of the alkylated Schiff base is preceded by conversion of the carboxylic acid ester to an amide, e.g. by reaction with ammonia, mono- or dialkylamine to yield compounds

of formula I, wherein R_1 is carbamoyl, alkylcarbamoyl or dialkylcarbamoyl. Alternatively, compounds of formula I, wherein R_1 is carbamoyl, alkylcarbamoyl or dialkylcarbamoyl, may be prepared by reacting a compound of formula I, wherein R_1 is esterified carboxy, with ammonia, mono- or dialkylamine.

The compounds of the invention may be converted to other compounds of the invention in conventional manner, e.g. by introducing substituents into the α- amino group, by converting esters to the corresponding acids or acids to esters.

The introduction of substituents into the amino group can be effected in conventional manner. For example the alkylation of the amino group may be carried out with alkyl halides or alkyl sulfates. If only one alkyl group has to be introduced suitably dialkylation is prevented by application of known methods, e.g. N-acylation, alkylation via N-acyl anion, removal of the acyl group. When the compound to be alkylated contains a free carboxy group, (i.e. R_1 is carboxy) it is preferably blocked by a protecting group e.g. benzyl removable by selective hydrogenolysis. The acylation of the amino group can be effected by reaction with the appropriate acid or a reactive derivative thereof. The urethane can be prepared by reaction with a haloformic acid ester.

The conversion of an ester to the corresponding acid can be effected by any conventional method, e.g. by hydrolysis. Using selective methods compounds of the invention can be prepared wherein either the phosphonic acid diester or the carboxylic acid ester is converted into the corresponding acid.

For example, compounds of formula I wherein R_1 is esterified carboxy and Y is group b) or c) can be converted to compounds of formula I, wherein R_1 is esterified carboxy and Y is group a) by silylation with e.g. bromotrimethylsilane and subsequent hydrolysis under mild conditions of the resulting bis-silyl phosphonate. By hydrolysing compounds of formula I wherein R_1 is esterified carboxy and Y is group b) or c) under mild conditions, e.g. dilute hydrochloric acid at elevated temperatures e.g. 60-70°C, compounds of formula I are obtained, wherein R_1 is carboxy and Y is a group b) or c).

The esterification can be carried out using conventional methods. When a phosphonic acid monoester [Y is a group d)] is desired the esterification can for example be effected with an alcohol in pyridine in the presence of trichloroacetonitrile at a temperature of about 100°C.When in the starting material to be esterified the amino group is unsubstituted or monosubstituted with a group that is other than a carbonyl-containing group, such amino group is suitably protected by an amino-protecting group. Conventional amino-protecting groups such as benzyloxycarbonyl or tert.-butyloxycarbonyl can be used. The deprotection can be carried out using conventional procedures, e.g. by treatment with trifluoroacetic acid. The benzyloxycarbonyl group can also be removed by hydrogenolysis.

The conversion of a carboxylic acid to an ester can be carried out using conventional methods.

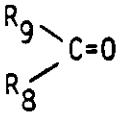
The optional formation of a salt, when the resulting compound of formula I contains a salifiable group may be carried out conventionally.

Racemates can be resolved into the optical antipodes by conventional methods, for example by e.g. separation of diastereoisomeric salts formed by a basic end product with an optically active acid, e.g. by fractional crystallisation of d- or ℓ-tartrates, d- or ℓ-di-O,O'-toluyl-tartrates or d- or ℓ-camphorsulfonates.

Compounds of formula VI used as starting material can be prepared by e.g. condensing a compound of formula VIII,

$$H_2N - CH_2 - W \qquad \text{VIII}$$

wherein W is as defined above, with a compound of formula IX,

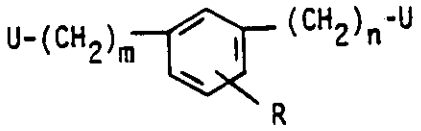


IX

wherein R_8 and R_9 are as defined above.

The reaction can be effected in known manner.

Compounds of formula VII can be prepared by reacting a compound of formula X

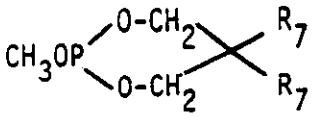


X

wherein m, n, R and U are as defined above, with a compound of formula XI or XII

$$P(OR_6)_3 \qquad \text{XI}$$

or



XII

wherein R_6 and R_7 are as defined above.

The reaction can be carried out in conventional manner.

Insofar as the production of the starting materials for the above processes is not particularly described, these may be produced in analogous manner to known compounds or to processes described herein.

In the following Examples all temperatures are given in degrees centrigrade and are uncorrected. The $[\alpha]_D^{20}$- and $[\alpha]_{365}^{20}$ - values are also uncorrected.

Example 1: (±)-α-Amino-3-(4'-chloro-5-phosphonomethyl-[1.1'-biphenyl]-3-yl)propanoic acid

To a stirred mixture of 3.7 g of the ketimine of glycinonitrile and benzophenone, 350 mg benzyltributylammonium chloride, 1.6 g sodium hydroxide, 3.2 ml water and 32 ml toluene is added dropwise at 0° over 90 minutes 4.9 g diethyl[3-bromomethyl-5-(4'-chlorophenyl)-phenyl]methylphosphonate. Stirring is then continued at room temperature for 24 hours. Thereafter the reaction mixture is diluted with water and extracted with methylene chloride. The organic layers are washed with water, dried over anhydrous sodium sulfate and evaporated. The residue is chromatographed on 200 g of silica gel (230-400 mesh) with CH_2Cl_2/acetic acid ethyl ester (3:1). The fractions with the main product are evaporated in vacuo. The residue is refluxed with 30 ml 7N hydrochloric acid for 12 hours. The mixture is extracted with toluene/ether (1:1). The aqueous layers are evaporated in vacuo, the residue dissolved in tetrahydrofuran/water, treated with propylene oxide and evaporated under vacuum. The residue is stirred in warm methanol to afford the title compound, m.p. 282-285° (decomp.).

The starting material diethyl[3-bromomethyl-5-(4'-chlorophenyl)-phenyl]methylphosphonate may be obtained as follows:

A mixture of 5.9 g 3,5-bis-bromomethyl-[4'-chloro-1.1'-biphenyl], 3.3 ml triethyl phosphite and 60 ml xylene is stirred under reflux for 90 minutes. The mixture is evaporated. The residue is chromatographed on 120 g of silica gel (230-400 mesh) with acetic acid ethyl ester. The fractions with the product are evaporated in vacuo to give the heading compound as a yellow oil.

Example 2: (±)-α-Amino-3-(3-phosphonomethyl)phenyl-propanoic acid

In manner analogous to that described in Example 1 the title compound, m.p. 271-275° (decomp.), is obtained.

Example 3: (±)-α-Amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid

To a stirred mixture of 3.7 g of the ketimine of glycinonitrile and benzophenone, 350 mg benzyltributylammonium chloride, 1.6 g sodium hydroxide, 3.2 ml water and 32 ml toluene is added dropwise at 0° over 90 minutes 4.5 g diethyl[(3-bromomethyl-5-phenyl)phenyl]-methylphosphonate. Stirring is then continued at room temperature for 24 hours. Thereafter the reaction mixture is diluted with water and extracted with methylene chloride. The organic layers are washed with water, dried over anhydrous sodium sulfate and evaporated. The residue is chromatographed on 500 g of silica gel (230-400 mesh) with acetic acid ethyl ester. The fractions with the main product are evaporated in vacuo. The residue is refluxed with 30 ml 7N hydrochloric acid for 12 hours. The mixture is extracted with toluene/ether (1:1). The aqueous layers are evaporated under vacuum, the residue dissolved in tetrahydrofuran/water, treated with propylene oxide and evaporated under vacuum. The residue is stirred in warm methanol to afford the title compound, m.p. 260-263° (decomp.).

The starting material diethyl [(3-bromomethyl-5-phenyl)phenyl] methylphosphonate may be obtained as follows:

A mixture of 11.2 g 3,5-bis-bromomethyl-[1.1'-biphenyl], 6.5 ml triethyl phosphite and 110 ml xylene is stirred under reflux for 90 minutes. The mixture is evaporated. The residue is chromatographed on 400 g of silica gel (230-400 mesh) with acetic acid ethyl ester. The fractions with the product are evaporated in vacuo to give the heading compound as a yellow oil.

Example 4: (±)-α-Amino-3-(5-octyloxy-3-phosphonomethyl)phenyl-propanoic acid

In manner analogous to that described in Example 1 but using (3-bromomethyl-5-octyloxyphenyl)-methyl-phosphonic acid diethyl-ester as starting material the title compound is prepared, m.p. 243-246° (decomp.).

Example 5: (±)-α-Amino-3-(5-diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl)propanoic acid ethyl ester

To a stirred mixture of 5.0 g of the ketimine of glycine ethyl ester and benzophenone, 6.7 g diethyl[(3-bromomethyl-5-phenyl)phenyl]-methylphosphonate, 0.3 g KJ and 150 ml dioxane are added dropwise at 10° 7.1 ml aqueous benzyltrimethylammonium hydroxide (40%) over 30 minutes. Stirring is then continued at room temperature for 2 hours. Thereafter the reaction mixture is diluted with water and extracted with toluene. The organic layer is washed with water, dried (Na_2SO_4) and evaporated. The residue is stirred at room temperature with 50 ml 1N HCl and 50 ml ether for 2 hours. The

aqueous layer is separated, made alkaline with $NaHCO_3$ and extracted with CH_2Cl_2. The organic layer is dried (Na_2SO_4) and evaporated to yield the title compound as a yellow oil. M.p. of the hydrochloride 140-142°, crystallized from ethanol/diethyl ether.

Example 6: (±)-α-Amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid ethyl ester

5.5 g of the oily base of Example 5 are dissolved in 100 ml abs. CH_2Cl_2 and treated with 16.5 ml bromotrimethylsilane. The mixture is left at room temperature for 24 hours. After evaporating to dryness the residue is dissolved in 150 ml H_2O/ tetrahydrofuran (1:1) and treated with propylene oxide, whereby the title compound crystallizes, m.p. 290-293° (decomp.).

Example 7: (±)-α-Amino-3-(4'-chloro-5-(diethoxyphosphinyl) methyl-[1.1'-biphenyl]-3-yl)propanoic acid amide

To a stirred mixture of 5.0 g of the ketimine of glycine methyl ester and benzophenone, 8.0 g diethyl [3-bromomethyl-5-(4'-chlorophenyl)phenyl]methylphosphonate, 0.3 g KJ and 150 ml dioxane are added dropwise at 10° 7.8 ml aqueous benzyltrimethylammonium hydroxide (40%) over 30 minutes. Stirring is then continued at room temperature for 2 hours. Thereafter the reaction mixture is diluted with water and extracted with toluene. The organic layer is washed with water, dried (Na_2SO_4) and evaporated. The residue is taken up in 250 ml CH_3OH and gaseous NH_3 is introduced at 10°. The mixture is stirred at room temperature for 66 hours and evaporated. The residue is

stirred at room temperature with 70 ml 1N HCl and 70 ml tetrahydrofuran for 1 1/2 hours. The tetrahydrofuran is evaporated and the residue extracted with toluene/diethyl ether (1:1). The aqueous layer is separated, made alkaline with Na_2CO_3 and extracted with CH_2Cl_2. The organic layer is dried (Na_2SO_4) and evaporated to yield the title compound as a foam.

^{1}H-NMR (360 MHz, $CDCl_3$): δ 1.25(t,J=6,6H), 1.6 (br.s,2H), 2.9(m,1H), 3.1(m,1H), 3.2(d,J=24,2H), 3.7(m,1H), 4.0(m,4H), 6.6(br. s,2H), 7.1-7.6(7H).

Example 8: (±)-α-Amino-3-(4'-chloro-5-phosphonomethyl-[1.1'-biphenyl] 3-yl) propanoic acid amide

4.2 g of the Example 7 compound are dissolved in 50 ml CH_2Cl_2 and treated with 17.7 ml bromotrimethylsilane. The mixture is stirred for 48 hours. After evaporation the residue is taken up with CH_3OH and evaporated. This procedure is effected 3 times. The residue is crystallized from CH_3OH/ethyl acetate (1:1), to yield the title compound, m.p. 278-280° (decomp.).

Example 9: (±)-α-Palmitoylamino-3-(5-phosphonomethyl-[1.1'-biphenyl] 3-yl) propanoic acid

A mixture of 335 mg (±)-α-amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid in 30 ml dimethylformamide and 0.76 ml N-ethyl-diisopropylamine under nitrogen are added dropwise at room temperature within 10 minutes 0.4 ml palmitic acid chloride. The mixture is stirred at room temperature for 26 hours. The solvent is evaporated

in vacuo. The oily residue is taken up in water, acidified with 2N HCl to pH 1 and extracted with diethyl ether. The extract is washed with saturated aqueous sodium chloride solution, dried (Na_2SO_4) and evaporated. The residue is recrystallized from diethyl ether/petroleum ether to yield the title compound, m.p. 130-140°. MS (FAB):574(MH^+).

Example 10: (±)-α-Amino-3-(4'-chloro-5-(diethoxyphosphinyl)-methyl-[1.1'-biphenyl]-3-yl)-propanoic acid methyl ester

In analogous manner to that described in Example 5 using the ketimine of glycine methyl ester and benzophenone and diethyl[3-bromomethyl-5-(4'-chlorophenyl)phenyl]methylphosphonate, the title compound is obtained as an oil, DC in CH_2Cl_2/CH_3OH (9:1) Rf = 0.46.

Example 11: (±)-α-Amino-3-(4'-chloro-5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid methyl ester

In manner analogous to that described in Example 6 the title compound is obtained, m.p. 300-305° (decomp.).

Example 12: (±)-α-Amino-3-(4'-chloro-5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid cinnamyl ester

a) (±)-α-Amino-3-(4'-chloro-5-(diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl) propanoic acid

4.6 g (±)-α-amino-3-(4'-chloro-5-(diethoxyphosphinyl)methyl-[1.1'-bi-

phenyl]-3-yl)propanoic acid ethyl ester, 1 equivalent of 1N sodium hydroxide and 4 parts by volume of tetrahydrofuran are stirred at room temperature for about 15 hours. The tetrahydrofuran is evaporated in vacuo. The residue is extracted with toluene/diethyl ether (1:1). The pH of the aqueous layer is adjusted to 5, whereby the heading compound crystallizes, m.p. 195-205° (decomp.).

b) (±)-α-tert.Butyloxycarbonylamino-3-(4'-chloro-5-(diethoxy-phosphinyl)methyl-[1.1'-biphenyl]-3-yl) propanoic acid

To 1.28 g of the step a) product and 2.2 ml tert.butyl alcohol are added under stirring 3.3 ml 1N aqueous NaOH solution. The mixture is stirred until a clear solution ensues and is then treated dropwise with 0.65 g di-tert.butyl carbonate. The mixture is stirred at room temperature 21 hours. The mixture is then cooled in an ice bath and treated dropwise with a solution of 0.45 g $KHSO_4$ in 3 ml water and extracted 3 times with CH_2Cl_2. The combined extracts are evaporated to dryness to give the heading compound, as a white foam, which recrystallized from diethyl ether has a m.p. 110-114°.

c) (±)-α-tert.Butyloxycarbonylamino-3-(4'-chloro-5-(diethoxy-phosphinyl)methyl-[1.1'-biphenyl]-3-yl) propanoic acid cinnamyl ester

To a solution of 1.05 g of the product of step b) in 5 ml dimethyl-formamide are added 362 mg tetramethylammonium hydroxide penta-hydrate. The mixture is stirred at room temperature 1 1/2 hours and then treated with 394 mg cinnamyl bromide. The mixture is stirred at room temperature 17 hours. The mixture is diluted with

ice/water (about 50 ml) and extracted with diethyl ether. The extracts are washed with 10 ml aqueous 1N $KHCO_3$ solution, dried (Na_2SO_4) and evaporated to give the heading compound as an oil.

d) (±)-α-Amino-3-(4'-chloro-5-(diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl)propanoic acid cinnamyl ester

5.5 g of the product of step c) and 50 ml aqueous trifluoroacetic acid (70%) are stirred at room temperature 20 hours. To the mixture are added CH_2Cl_2 and dropwise aqueous $KHCO_3$ solution. The organic phase is dried (Na_2SO_4) and evaporated. The residue is taken up in diethyl ether, filtered and evaporated to dryness, to give the heading compound as an oil.

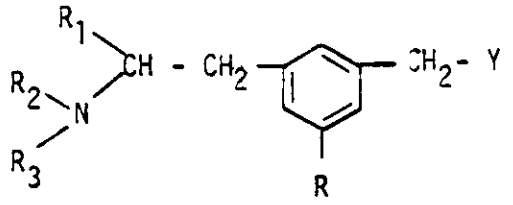
^{1}H-NMR (80MHz, $CDCl_3$): δ 1.3 (m,6H), 1.8 (br.s,2H), 2.9(m,1H), 3.1 (m,1H), 3.1 (d, J =22,2H), 3.8 (m,1H), 4.0 (m,4H), 4.8 (d, J =6, 2H), 6.3 (m,1H), 6.7 (d, J=15 , 1H), 7.0-7.6 (sh,12H).

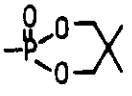
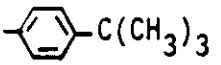
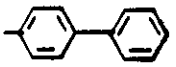
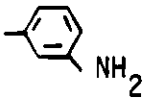
e) (±)-α-Amino-3-(4'-chloro-5-phosphonomethyl-[1.1'- biphenyl]-3-yl) propanoic acid cinnamyl ester

In manner analogous to that described in Example 6 the title compound is obtained, m.p. 253-255°.

Example 13:

In manner analogous the following compounds (racemates) are obtained:



Ex.	R_1	R_2	R_3	R	Y	m.p.	analogous to Ex.
a)	$COOC_2H_5$	H	H			amorphous [1]	5
b)	COOH	H	H		$-P(O)(OH)_2$	decomp. >330 °	1
c)	COOH	H	H		$-P(O)(OH)_2$	decomp. >295 °	1
d)	$COOCH_2CO$	H	H		$-P(O)(OH_2)$	220-225° (decomp.)	9
e)	$COOCH_3$	H	H		$-P(O)(OH)_2$	hydrobromide amorphous [2]	6
f)	COOH	H	$COCH_3$		$-P(O)(OH)_2$	amorphous [3]	8
g)	COOH	H	H		$-P(O)(OH)_2$	> 315° [4]	1

[1] Rf 0.35 (CH_2Cl_2/CH_3OH/conc. NH_3 9:1:0.1)

[2] ^{1}H-NMR (360 MHz, DMSO-d_6): δ 3.05 (d, J=20, 2H), 3.15 (2H), 3.75 (s,3H), 4.45 (br.s,1H), 7.1 (s,1H), 7.65-7.35 (7H), 8.45 (br.s,3H)

[3] Rf 0.75 (ethyl acetate/acetic acid/water 5:2:2)
^{1}H-NMR (60 MHz, CD_3OD): δ u.a. 1.8 (s,3H), 2.9 (d, J=22, 2H)

[4] ^{1}H-NMR (360 MHz, DMSO-d_6): δ u.a. 2.95 (d,J=20,2H), 3.1 (m,2H), 4.1 (m,1H).

Example 14: (+)-α-Amino-3-(5-(diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl)propanoic acid ethyl ester

An etheral solution of 29.9 g (±)-α-amino-3-(5-(diethoxyphosphinyl)-methyl-[1.1'-biphenyl]-3-yl)propanoic acid ethyl ester and an etheral solution of 27.6 g (+)-di-O,O'-p-toluyl-D-tartaric acid are mixed whereby the crude salts precipitate. The salts are filtered and crystallized from ethanol/t-butyl methyl ether (1:4). The resulting crystalls are recrystallized three times from isopropanol/tert. butyl methyl ether (1:8) to give pure (+)-α-amino-3-(5-(diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl)propanoic acid ethyl ester (+)-di-O,O'-p-toluyl-D-tartrate, m.p. 155-158°, $[\alpha]_D^{20}$ = + 88.6° (c = 1 in C_2H_5OH / 1N HCl 2:1).

The above salt is treated with saturated aqueous $KHCO_3$ solution and extracted with CH_2Cl_2 to give (+)-α-amino-3-(5-(diethoxyphosphinyl) methyl-[1.1'-biphenyl]-3-yl) propanoic acid ethyl ester as an oil. The hydrochloride has a m.p. of 150-152° (decomp.), crystallized from ethanol/diethyl ether, $[\alpha]_D^{20}$ = + 17.7° (c = 1 in 2N HCl).

Example 15: (+)-α-Amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl)-propanoic acid ethyl ester

In manner analogous to that described in Example 6 and using the compound of Example 14 as starting material, the title compound is obtained m.p. 280-285° (decomp.), $[\alpha]_D^{20}$ = + 5.0° (c = 1 in 1N HCl), $[\alpha]_{365}^{20}$ = + 31.0° (c = 1 in 1N HCl).

Example 16: (+)-α-Amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid

The compound of Example 15 and 1N hydrochloric acid are heated at 60° for 2 hours. After evaporating to dryness the residue is dissolved in tetrahydrofuran/water and treated with propylene oxide, whereby the title compound is obtained, m.p. 275-278° (decomp.). $[\alpha]_D^{20}$ = 0.0 ± 0.5° (c = 1 in 6N HCl), $[\alpha]_{365}^{20}$ = + 21.3° (c = 1 in 6N HCl).

Example 17: (-)-α-Amino-3-(5-(diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl)propanoic acid ethyl ester

In manner analogous to that described in Example 14 and using (±)-α-amino-3-(5-diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl)propanoic acid ethyl ester and (-)-di-O,O'-p-toluyl-L-tartaric acid as starting material, the title compound is obtained. The hydrochloride has a m.p. of 150-152° (decomp.), $[\alpha]_D^{20}$ = - 17.3° (c = 1 in 2N HCl).

Example 18: (-)-α-Amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl)-propanoic acid ethyl ester

In manner analogous to that described in Example 6 and using the compound of Example 17 as starting material, the title compound is obtained m.p. 277-282° (decomp.), $[\alpha]_D^{20}$ = - 4.4° (c = 1 in 1N HCl), $[\alpha]_{365}^{20}$ = - 28.1° (c = 1 in 1N HCl).

Example 19: (-)-α-Amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl)propanoic acid

In manner analogous to that described in Example 16 and using the compound of Example 18 as starting material, the title compound is obtained, m.p. 274-276° (decomp.). $[\alpha]_D^{20}$ = 0.0 ± 0.5° (c = 1 in 6N HCl), $[\alpha]_{365}^{20}$ = - 20.3° (c = 1 in 6N HCl).

The compounds of the invention exhibit pharmacological activity and are, therefore, indicated for use as pharmaceuticals e.g. for therapy. In particular, the compounds exhibit central nervous system activity as indicated in standard tests. For example, the compounds inhibit the locomotion in mice.

In this test groups of 3 male mice (18-24 g, OF-1, Sandoz Basle) receive 3.2, 10, 32, 100 and 320 mg i.p. of the test drug. 1 hour after drug administration the mice are observed individually and their locomotion compared with that of control mice concurrently treated with vehicle. The locomotion is judged to be either unaffected, definitely more or less than controls, strongly more or less than controls, or completely inhibited.

The compounds of the invention additionally exhibit anticonvulsant activity as indicated in standard tests. In a first test, the compounds inhibit the electroshock-induced convulsions in the mouse [c.f. E. Swinyard, J.Am.Pharm.Assoc.Scient.Ed. 38, 201 (1949) and J.Pharmacol.Exptl.Therap. 106, 319 (1952)]. In this test group of 3 mice (18-26 g, OF-1, Sandoz Basle) receive the test substance in a dosage of 3.2-100 mg/kg i.p. After 60 minutes a 50 mA, 200 ms long shock is applied with corneal electrodes smeared with electrolyte jelly. This supra-threshold shock produces tonic extensor convulsions of all extremities. Inhibition of the hindlimb extension is taken as a protective action. After investigation of several dose-levels an ED_{min} is estimated.

In a second test the compounds inhibit N-Methyl-D-aspartic acid (NMDA) induced convulsions in the mouse. In this test groups of 6 female mice (18-26 g, OF-1, Sandoz Basle) were pretreated with the test substance in a dosage of 0.1 - 100 mg/kg i.p. 30 minutes later

they are challenged with 400 mg/kg s.c. NMDA in the neck region and observed for 30 minutes. The latencies for the appearance of the first signs of convulsions, for the first tonic convulsions and for the occurence of death are noted. The significance of any differences is observed using the Mann-Whitney U-test [S. Siegel, Non-parametric Statistics, McGraw-Hill, New York 1956]. After investigation of several dose-levels the threshold dose is estimated. This dose represents the smallest dose at which there is a significant inhibition of convulsive symptoms.

As a result of their anticonvulsant activity the compounds of the invention are indicated for use for the treatment of epilepsy. For this use an indicated daily dosage is in the range from about 25 to about 800 mg of the compound conveniently administered in divided doses 2 to 4 times a day in unit dosage form containing for example from about 6 to about 400 mg of the compound or in sustained release form.

The compounds of the invention further interact with excitatory amino acid systems, in particular they are competitive antagonists of NMDA (N-Methyl-D-aspartic acid) receptors, as indicated by an inhibitory effect on NMDA-induced depolarizations of the isolated amphibian spinal cord [P.L. Herrling, Neuroscience 14 (1985) 417-426]. The compounds of the invention show this activity at concentrations of from about 100 nM/l to about 300 µM/l.

The compounds of the invention are also selective as indicated in that quisqualate induced depolarisations are not significantly effected in the above test wherein NMDA is replaced by quisqualic acid.

As a result of their NMDA receptor antagonism the compounds are indicated for the use i) in the treatment of disorders having an etiology comprising or associated with excess GH-secretion e.g. in the treatment of diabetes mellitus and angiopathy as well as of acromegaly and ii) in the treatment of disorders having an etiology associated with or modulated by excess LH-secretion e.g. in the treatment of prostate hypertrophy or in the treatment of menopausal syndrome. For this use an indicated daily dosage is in the range from about 1 to about 800 mg of the compound conveniently given in divided doses 2 to 4 times a day in unit dosage form containing for example from about 0.25 to about 400 mg of the compound or in sustained release form.

As a result of their NMDA receptor antagonism the compounds of the invention are further indicated for use for the treatment of anxiety, schizophrenia and depression or of CNS degenerative disorders, such as Huntington's, Alzheimer's or Parkinson's diseases. For these uses an indicated daily dosage is in the range from about 25 to about 800 mg of the compound conveniently administered in divided doses 2 to 4 times a day in unit dosage form containing for example from about 6 to about 400 mg of the compound or in sustained release form.

The compounds of the invention protect further against hypoxia-induced degeneration of rat hippocampal neurons in vitro at concentrations ranging from 1 µM to 3 mM [method of S. Rothman, J.Neurosci. 4, 1884-1891 (1984)]. The compounds are therefore useful in the treatment of cerebral hypoxic/ischaemic conditions, e.g. stroke. For this use an indicated daily dosage is in the range from about 10 to about 800 mg of the compound conveniently administered in divided doses 2 to 4 times a day in unit dosage form containing for example from about 2 to about 400 mg of the compound or in sustained release form.

Furthermore, the compounds of the invention inhibit plasma corticosterone rise, which is induced by social stress in mice. This can be shown in the following test:

One day before the experiment a group of 5 male mice (40-50 g OF-1, Sandoz, Basle) were placed in a transparent makrolon (R.T.M.) cage Typ 3, which is cut in halves by a grid. The next day each mouse was given an oral dose of 0.3-30 mg/kg of a compound of the invention. Two hours later an isolated male mouse was introduced for 15 minutes into the empty half of the cage and two trained observers recorded the behaviour of the mice in terms of acts such as dig, push-dig and rattle. Blood plasma samples were then taken from the tested mice group and assayed for corticosterone concentrations using a modified method of Paerson-Murphy B.E., J.Clin.Endocrinology 27 (1967) 973-990. The procedure was repeated with a control group of 5 mice which was given only a solvent.

As a result of their ability to inhibit plasma-corticosterone rise the compounds of the invention are indicated for use for the treatment of stress-related psychiatric disorders, e.g. where the treatment of social withdrawal, which is present in many psychiatric disorders, e.g.schizophrenia, depression, generalized anxiety or in affective disorders, e.g. adjustment disorders with social withdrawal or anxiety, and other stress-related illnesses is desired.

For this use an indicated daily dosage is in the range from about 3 to about 800 mg of the compound conveniently administered in divided doses 2 to 4 times a day in unit dosage form containing for example from about 0.75 to about 400 mg of the compound or in sustained release form.

The compounds of the invention may be administered by any conventional route, in particular enterally, preferably orally e.g. in the form of tablets or capsules, or parenterally e.g. in form of injectable solutions or suspensions.

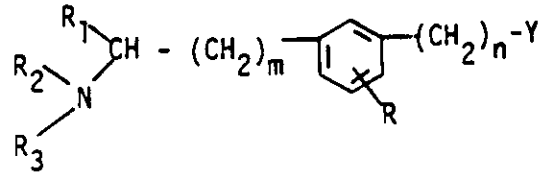
α-Amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl)propanoic acid is the preferred compound for the treatment of stress-related psychiatric disorders.

The compounds of the invention may be administered as such or as their pharmaceutically acceptable salts. Such salts exhibits the same order of activity as the compounds of the invention in free base or free acid form. The present invention also provides pharmaceutical compositions comprising a compound of the invention as such or in salt form in association with at least one pharmaceutical carrier or diluent. Such compositions may be manufactured in conventional manner.

C L A I M S :

1. A process for the production of an α-amino-α-(3-alkylphenyl)alkyl-ethanoic acid, an ester thereof or an amide thereof, in which the 3-alkyl moiety bears a phosphorus oxo acid group or an ester thereof, wherein phosphorus is attached directly to the alkyl moiety, or a salt thereof, which comprises reacting a protected glycine derivative with an appropriate 1-alkyl-3-alkyl-benzene, in which one alkyl moiety bears a phosphorus oxo acid ester group, wherein phosphorus is attached directly to the alkyl moiety and the other alkyl group bears a leaving group, under basic conditions and hydrolysing the resulting compound, and, if desired, converting a resulting compound of the invention into another compound of the invention, and/or, if desired, converting a resulting free compound into a salt; and/or, if desired, resolving a racemate obtained into the optical antipodes.

2. A process according to claim 1 for the production of a compound of formula I,



I

wherein

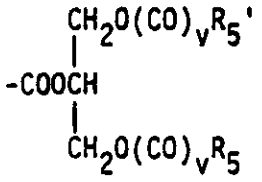
m and n are independently 1 or 2,

R_1 is carboxy, (C_{1-12})alkoxycarbonyl, benzoyl(C_{1-4})alkoxycarbonyl,

phenyl(C_{2-4})alkenyloxycarbonyl, carbamoyl, monoalkyl(C_{1-6})carbamoyl or dialkyl(C_{1-6})carbamoyl,

R_2 is hydrogen or (C_{1-12})alkyl,

R_3 is hydrogen, (C_{1-12})alkyl, (C_{1-18})alkylcarbonyl, (C_{2-22})alkenylcarbonyl, (C_{4-22})alkadienylcarbonyl, (C_{6-22})alkatrienylcarbonyl, (C_{8-22})alkatetraenylcarbonyl, (C_{1-12})alkoxycarbonyl or a group of formula II,

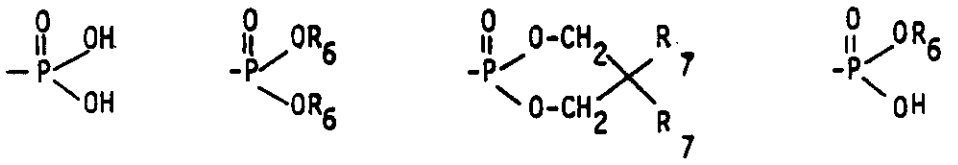


II

in which R_5 and R_5' are each, independently (C_{1-22})alkyl, (C_{2-22})alkenyl, (C_{4-22})alkadienyl, (C_{6-22})alkatrienyl, (C_{8-22})alkatetraenyl and v is independently of each other 0 or 1,

R is hydrogen, halogen, hydroxy, (C_{1-12})alkyl, (C_{1-12})alkoxy, phenyl, phenyl(C_{1-8})alkoxy, phenyl(C_{1-8})alkyl; phenyl substituted by halogen, (C_{1-12})alkyl, (C_{1-12})alkoxy, amino, (C_{1-12})alkylcarbonylamino, hydroxy or phenyl,

Y is one of the groups a), b), c) or d)



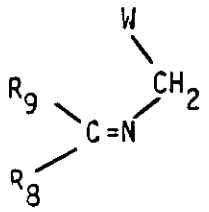
a) b) c) d)

wherein

R_6 is (C_{1-6})alkyl and

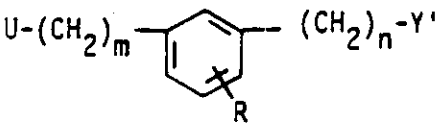
R_7 is hydrogen or (C_{1-6})alkyl,

or a salt thereof, which comprises reacting a compound of formula VI



VI

wherein R_8 is hydrogen, alkyl or phenyl, R_9 is phenyl optionally substituted by chlorine, alkyl or alkoxy, and W is -CN or $-COOR_{10}$, wherein R_{10} is an ester forming radical, with a compound of formula VII,



VII

wherein m, n and R are as defined above, U is a leaving group, and Y' is one of the groups b) or c), under basic conditions, and hydrolysing the resulting compound, and, if desired, converting a resulting compound of formula I into another compound of formula I; and/or, if desired, converting a resulting free compound into a salt; and/or, if desired, resolving a racemate obtained into the optical antipodes.

3. A process for the production of an α-amino-α-(3-alkylphenyl)-alkyl-ethanoic acid, an ester thereof or an amide thereof, in which the 3-alkyl moiety bears a phosphorus oxo acid group or an ester thereof, wherein phosphorus is attached directly to the alkyl moiety, or a salt thereof, as hereinbefore described with reference to any of the Examples.

4. An α-amino-α-(3-alkylphenyl)alkyl-ethanoic acid, an ester thereof or an amide thereof, in which the 3-alkyl moiety bears a phosphorus oxo acid group or an ester thereof, wherein phosphorus is attached directly to the alkyl moiety, or a salt thereof, whenever produced by a process of claim 1, 2 or 3.

5. An α-amino-α-(3-alkylphenyl)alkyl-ethanoic acid, an ester thereof or an amide thereof, in which the 3-alkyl moiety bears a phosphorus oxo acid group or an ester thereof, wherein phosphorus is attached directly to the alkyl moiety, or a salt thereof.

6. A compound according to claim 5, wherein phenyl is substituted by alkoxy, phenyl or phenyl substituted by halogen, alkyl or phenyl or a salt thereof.

7. A compound of formula I as defined in claim 2 or a salt thereof.

8. A compound of claim 7, wherein
m and n are independently 1 or 2,

R_1 is carboxy or (C_{1-12})alkoxycarbonyl,

R_2 is hydrogen or (C_{1-12})alkyl,

R_3 is hydrogen, (C_{1-12})alkyl, (C_{1-12})alkylcarbonyl, (C_{2-22})alkenylcarbonyl, (C_{4-22})alkadienylcarbonyl, (C_{6-22})alkatrienylcarbonyl, (C_{8-22})alkatetraenylcarbonyl, (C_{1-12})alkoxycarbonyl

or a group of formula II, in which R_5 and R_5' are each, independently (C_{1-22})alkyl, (C_{2-22})alkenyl, (C_{4-22})alkadienyl, (C_{6-22})alkatrienyl, (C_{8-22})alkatetraenyl and v is independently of each other 0 or 1,

R is hydrogen, halogen, hydroxy, (C_{1-12})alkyl, (C_{1-12})alkoxy, phenyl; phenyl(C_{1-8})alkoxy, phenyl(C_{1-8})alkyl, phenyl substituted by halogen, (C_{1-12})alkyl, (C_{1-12})alkoxy or phenyl,

Y is one of the groups a), b) or c),

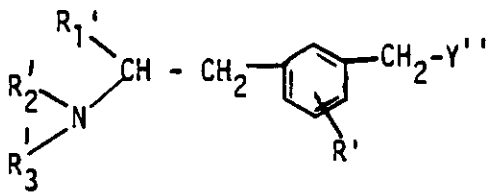
wherein

R_6 is (C_{1-6})alkyl and
R_7 is hydrogen or (C_{1-6})alkyl,

or a salt thereof.

9. A compound of claim 7, wherein m and n are independently 1 or 2, R_1 is carboxy, (C_{1-12})alkoxycarbonyl, benzoyl(C_{1-4})alkoxycarbonyl, phenyl(C_{2-4})alkenyloxycarbonyl or carbamoyl, R_2 is hydrogen, R_3 is hydrogen or (C_{1-18})alkylcarbonyl, R is hydrogen, (C_{1-12})alkoxy, phenyl, phenyl substituted by halogen, (C_{1-12})alkyl, amino or phenyl, Y is one of the groups a), b), c) or d), wherein R_6 is (C_{1-6})alkyl and R_7 is hydrogen or (C_{1-6})alkyl, or a salt thereof.

10. A compound according to claim 7 of the formula Ia,



Ia

wherein R_1' is carboxy, (C_{1-4})alkoxycarbonyl, benzoyl(C_{1-4})alkoxycarbonyl, phenyl(C_{2-4})alkenyloxycarbonyl or carbamoyl, R_2' is

hydrogen, R_3' is hydrogen or (C_{1-18})alkylcarbonyl, R' is (C_{1-12})-alkoxy, phenyl or phenyl substituted by halogen, (C_{1-12})alkyl , amino or phenyl, Y'' is one of the groups a), b) or c), wherein R_6 is (C_{1-6})alkyl and R_7 is hydrogen or (C_{1-6})alkyl, or a salt thereof.

11. A compound of claim 7 which is (±)-α-amino-3-(4-chloro-5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid or a salt thereof.

12. A compound of claim 7 which is a (±)-α-amino-3-(3-phosphonomethyl)phenyl-propanoic acid or a salt thereof.

13. A compound of claim 7 which is (±)-α-amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid or a salt thereof.

14. A compound of claim 7 which is (±)-α-amino-3-(5-octyloxy-3-phosphonomethyl)phenyl-propanoic acid or a salt thereof.

15. A compound of claim 7 which is (±)-α-amino-3-(5-diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl) propanoic acid ethyl ester or a salt thereof.

16. A compound of claim 7 which is (±)-α-amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid ethyl ester or a salt thereof.

17. A compound of claim 7 which is (±)-α-amino-3-(4'-chloro-5-(diethoxyphosphinyl) methyl-[1.1'-biphenyl]-3-yl)propanoic acid amide or a salt thereof.

18. A compound of claim 7 which is (±)-α-amino-3-(4'-chloro-5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid amide or a salt thereof.

19. A compound of claim 7 which is (±)-α-palmitoylamino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid or a salt thereof.

20. A compound of claim 7 which is (±)-α-amino-3-(4'-chloro-5-(diethoxyphosphinyl) methyl-[1.1'-biphenyl]-3-yl) propanoic acid methyl ester or a salt thereof.

21. A compound of claim 7 which is (±)-α-amino-3-(4'-chloro-5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid methyl ester or a salt thereof.

22. A compound of claim 7 which is (±)-α-amino-3-(4'-chloro-5-(diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl) propanoic acid or a salt thereof.

23. A compound of claim 7 which is (±)-α-tert.butyloxy-carbonylamino-3-(4'-chloro-5-(diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl) propanoic acid or a salt thereof.

24. A compound of claim 7 which is (±)-α-tert.butyloxy-carbonylamino-3-(4'-chloro-5-(diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl) propanoic acid cinnamyl ester.

25. A compound of claim 7 which is (±)-α-amino-3-(4'-chloro-5-(diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl) propanoic acid cinnamyl ester or a salt thereof.

26. A compound of claim 7 which is (±)-α-amino-3-(4'-chloro-5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid cinnamyl ester or a salt thereof.

27. A compound of claim 7 which is (±)-α-amino-3-(5-cyclic-2,2-dimethylpropylene phosphinyl)methyl-[1.1'-biphenyl] 3-yl) propanoic acid ethyl ester or a salt thereof.

28. A compound of claim 7 which is (±)-α-amino-3-(4'-tert.butyl-5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid or a salt thereof.

29. A compound of claim 7 which is (±)-α-amino-3-(4'phenyl-5-phosphonomethyl-[1.1'-biphenyl]-3-yl)propanoic acid or a salt thereof.

30. A compound of claim 7 which is (±)-α-amino-3-(4'-chloro-5-phosphonomethyl-[1,1'-biphenyl]-3-yl) propanoic acid benzoylmethyl ester or a salt thereof.

31. A compound of claim 7 which is (±)-α-amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid methyl ester or a salt thereof.

32. A compound of claim 7 which is (±)-α-acetylamino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid or a salt thereof.

33. A compound of claim 7 which is (±)-α-amino-3-(3'-amino-5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid or a salt thereof.

34. A compound of claim 7 which is (+)-α-amino-3-(5-(diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl)propanoic acid ethyl ester or a salt thereof.

35. A compound of claim 7 which is (+)-α-amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl)propanoic acid ethyl ester or a salt thereof.

36. A compound of claim 7 which is (+)-α-amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid or a salt thereof.

37. A compound of claim 7 which is (-)-α-amino-3-(5-(diethoxyphosphinyl)methyl-[1.1'-biphenyl]-3-yl)propanoic acid ethyl ester or a salt thereof.

38. A compound of claim 7 which is (-)-α-amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl)propanoic acid ethyl ester or a salt thereof.

39. A compound of claim 7 which is (-)-α-amino-3-(5-phosphonomethyl-[1.1'-biphenyl]-3-yl) propanoic acid or a salt thereof.

40. A compound according to any one of claims 4 to 39 or a pharmaceutically acceptable salt thereof for use as a pharmaceutical.

41. A compound according to claim 40 for use in the treatment of epilepsy, disorders associated with excess GH or LH secretion, anxiety, schizophrenia, depression, CNS degenerative disorders or cerebral hypoxic/ischaemic conditions.

42. A compound according to claim 40 for use in the treatment of stress-related psychiatric disorders.

43. A pharmaceutical composition which comprises a compound of claim 4 or a pharmaceutically acceptable salt thereof in association with a pharmaceutical carrier or diluent.

44. Use of a compound according to anyone of claims 4 to 39 or a pharmaceutically acceptable salt thereof in the preparation of a medicament for treating epilepsy, disorders associated with excess GH or LH secretion, anxiety, schizophrenia, depression, CNS degenerative disorders, cerebral hypoxic/ischaemic conditions or for the treatment of stress-related psychiatric disorders.

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Title SUBSTITUTED A-AMINO ACIDS, THEIR PREPARATION AND PHARMACEUTICAL COMPOSITIONS CONTAINING THEM

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