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<p>(54) Title: COMBINATION OF A GABA-A ALPHA 5 INVERSE AGONIST AND AN ACETYLCHOLINESTERASE INHIBITOR</p>		
<p>(57) Abstract</p>		
<p>The present invention relates to a combination of an acetylcholinesterase inhibitor and an inverse agonist of the GABA_A α₅ receptor subtype, and the use of the combination in treating neurodegenerative conditions such as Alzheimer's Disease.</p>		

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COMBINATION OF A GABA-A ALPHA 5 INVERSE AGONIST AND
AN ACETYLCHOLINESTERASE INHIBITOR

The present invention relates to a combination of an acetylcholinesterase
5 inhibitor and an inverse agonist of the GABA_A α_5 receptor subtype, and
the use of the combination in treating neurodegenerative conditions such
as Alzheimer's Disease.

Alzheimer's Disease is a poorly understood neurodegenerative
condition mainly affecting the elderly but also younger people who are
10 generally genetically predisposed to it.

One postulated method of treatment comprises the administration
of acetylcholinesterase inhibitors which act on the cholinergic system.
However this method suffers from the disadvantages that these
compounds induce a range of side-effects including diarrhoea, salivation
15 and nausea.

The present invention provides a new and surprisingly effective
synergistic combination of an acetylcholinesterase inhibitor and an inverse
agonist of the GABA_A α_5 receptor subtype for separate, sequential or
simultaneous administration.

20 The present invention provides a greater than expected
improvement in the condition of subjects suffering from a
neurodegenerative with an associated cognitive deficit, such as
Alzheimer's Disease or Parkinson's disease, or from a cognitive deficit
which may arise from a normal process such as aging or from an abnormal
25 process such as injury, than would be expected from administration of the
active ingredients alone. Further, the combination allows for a lower
overall dose of each of the active ingredients to be administered thus
reducing side effects and decreasing any reduction in the effectiveness of
each of the active ingredients over time.

30 Acetylcholinesterase inhibitors which may be used include any
which are known to the skilled person. Examples are physostigmine,

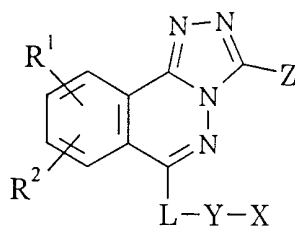
neostigmine, edrophonium and its chloride, pyridostigmine and its
bromide, eptastigmine, and its tartrate, galanthamine and its
hydrochloride and hydrobromide, metrifonate, tacrine and its
hydrochloride, eseridine and its salicylate, suronacrine and its maleate,
5 velnacrine and its maleate, amiridine and its hydrochloride, 7-
methoxytacrine, SM-10888 and its citrate, phenserine and its tartrate,
ENA-713, donepezil and its hydrochloride, TAK-147, CP-118954,
huperzine A and zifrosilone.

Any inverse agonist of the GABA_A α_5 receptor subtype may be used
10 which fulfills the criteria of WO-A-9625948. The inverse agonist may be
either binding selective for the α_5 subtype or functionally selective, or
both. Thus the inverse agonist is preferably an antagonist, or has
insignificant agonist or inverse agonist properties at the other GABA_A α
receptor subtypes when measured in oocytes as described in
15 WO-A-9625948.

Thus the inverse agonist preferably has a functional efficacy at the
 α_5 receptor subunit of less than -20% and functional efficacies at the α_1 ,
 α_2 , and α_3 receptor subunits of between -20 and +20%. By functional
efficacy is meant the percentage modulation of the EC₂₀ response produced
20 by GABA, upon coadministration of the inverse agonist, in oocytes
expressing GABA_A receptor channels containing the α receptor subunit
under test. Details of this measurement are given in WO-A-9625948.

The inverse agonist preferably binds selectively to GABA_A receptors
containing the α_5 subunit 10, 25 and particularly 50 times compared to
25 GABA_A receptors subunits containing the α_1 , α_2 or α_3 subunits. Preferably
this binding selectivity is shown over all these subunits.

A preferred class of inverse agonists, which are disclosed in
WO-A-9850385, are of formula I:



wherein:

- R^1 is hydrogen, halogen or CN or a group C_{1-4} alkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} alkoxy, C_{2-4} alkenyloxy or C_{2-4} alkynyloxy, each of which groups is unsubstituted or substituted with one or two halogen atoms or with a pyridyl or phenyl ring each of which rings may be unsubstituted or independently substituted by one or two halogen atoms or nitro, cyano, amino, methyl or CF_3 groups;
- R^2 is hydrogen, halogen or CN or a group C_{1-4} alkyl, C_{2-4} alkenyl, C_{2-4} alkynyl, C_{1-4} alkoxy, C_{2-4} alkenyloxy or C_{2-4} alkynyloxy each of which groups is unsubstituted or substituted with one or two halogen atoms;
- L is O, S or NR^n where R^n is H, C_{1-6} alkyl or C_{3-6} cycloalkyl;
- X is a 5-membered heteroaromatic ring containing 1, 2, 3 or 4 heteroatoms independently chosen from oxygen, nitrogen and sulphur, at most one of the heteroatoms being oxygen or sulphur, or a 6-membered heteroaromatic ring containing 1, 2 or 3 nitrogen atoms, the 5- or 6-membered heteroaromatic ring being optionally fused to a benzene ring and the heteroaromatic ring being optionally substituted by R^x and/or R^y and/or R^z , where R^x is halogen, R^3 , OR^3 , $OCOR^3$, NR^4R^5 , NR^4COR^5 , tri(C_{1-6} alkyl)silyl C_{1-6} alkoxy C_{1-4} alkyl, CN or R^9 , R^y is halogen, R^3 , OR^3 , $OCOR^3$, NR^4R^5 , NR^4COR^5 or CN and R^z is R^3 , OR^3 or $OCOR^3$, where R^3 is C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{3-6} cycloalkyl, hydroxy C_{1-6} alkyl and R^3 is optionally mono, di- or tri-fluorinated, R^4 and R^5 are each independently hydrogen, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, C_{3-6} cycloalkyl or CF_3 or R^4 and R^5 , together with the nitrogen atom to which they are attached, form a 4-7 membered heteroaliphatic ring containing the nitrogen atom as the sole heteroatom, and R^9 is benzyl or an aromatic ring containing either 6

atoms, 1, 2 or 3 of which are optionally nitrogen, or 5 atoms, 1, 2 or 3 of which are independently chosen from oxygen, nitrogen and sulphur, at most one of the atoms being oxygen or sulphur, and R⁹ is optionally substituted by one, two or three substituents independently chosen from
5 halogen atoms and C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₁₋₄alkoxy, C₂₋₄alkenyloxy and C₂₋₄alkynyloxy groups each of which groups is unsubstituted or substituted by one, two or three halogen atoms, and when X is a pyridine derivative, the pyridine derivative is optionally in the form of the N-oxide and providing that when X is a tetrazole derivative it
10 is protected by a C₁₋₄alkyl group; or X is phenyl optionally substituted by one, two or three groups independently selected from halogen, cyano, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl and C₃₋₆cycloalkyl;

Y is optionally branched C₁₋₄alkylidene optionally substituted by an oxo group or Y is a group (CH₂)_jO wherein the oxygen atom is nearest the
15 group X and j is 2, 3 or 4;

Z is a 5-membered heteroaromatic ring containing 1, 2 or 3 heteroatoms independently selected from oxygen, nitrogen and sulphur, at most one of the heteroatoms being oxygen or sulphur and providing that when two of the heteroatoms are nitrogen an oxygen or sulphur atom is
20 also present and that when one of the atoms is oxygen or sulphur then at least one nitrogen atom is present, or a 6-membered heteroaromatic ring containing 2 or 3 nitrogen atoms, Z being optionally substituted by R^v and/or R^w, where R^v is halogen, R⁶, NR⁷R⁸, NR⁷COR⁸, CN, furyl, thienyl, phenyl, benzyl, pyridyl or a 5-membered heteroaromatic ring containing at
25 least one nitrogen atom and optionally 1, 2 or 3 other heteroatoms independently selected from oxygen, nitrogen and sulphur, at most one of the other heteroatoms being oxygen or sulphur and R^w is R⁶ or CN;

R⁶ is C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₃₋₆cycloalkyl, hydroxyC₁₋₆alkyl, C₁₋₆alkoxy, C₂₋₆alkenyloxy, C₂₋₆alkynyloxy, CH₂F or CF₃;
30 and

R⁷ and R⁸ are each independently hydrogen, C₁₋₆alkyl, C₂₋₆alkenyl, C₂₋₆alkynyl, C₃₋₆cycloalkyl or CF₃ or R⁷ and R⁸, together with the nitrogen atom to which they are attached, form a 4-7 membered heteroaliphatic ring containing the nitrogen atom as the sole heteroatom;

5 or a pharmaceutically acceptable salt thereof.

As used herein, the expression "C₁₋₆alkyl" includes methyl and ethyl groups, and straight-chained and branched propyl, butyl, pentyl and hexyl groups. Particular alkyl groups are methyl, ethyl, n-propyl, isopropyl and t-butyl. Derived expressions such as "C₁₋₄alkyl", "C₂₋₄alkenyl",
10 "C₂₋₆alkenyl", "hydroxyC₁₋₆alkyl", "C₂₋₄alkyl" and "C₂₋₆alkynyl" are to be construed in an analogous manner.

The expression "C₃₋₆cycloalkyl" as used herein includes cyclic propyl, butyl, pentyl and hexyl groups such as cyclopropyl and cyclohexyl.

Suitable 5- and 6-membered heteroaromatic rings include pyridinyl,
15 pyridazinyl, pyrimidinyl, pyrazinyl, pyrrolyl, pyrazolyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, imidazolyl, oxadiazolyl and thiadiazolyl groups. A suitable 5-membered heteroaromatic ring containing four nitrogen atoms is tetrazolyl. Suitable 6-membered heteroaromatic rings containing three nitrogen atoms include 1,2,4-triazine and 1,3,5-triazine.

20 The term "halogen" as used herein includes fluorine, chlorine, bromine and iodine, of which fluorine and chlorine are preferred.

As used herein the term "C₁₋₆alkoxy" includes methoxy and ethoxy groups, and straight-chained, branched and cyclic propoxy, butoxy, pentoxy and hexoxy groups, including cyclopropylmethoxy. Derived
25 expressions such as "C₂₋₆alkenyloxy", "C₂₋₆alkynyloxy", "C₁₋₄alkoxy", "C₂₋₄alkenyloxy" and "C₂₋₄alkyloxy" should be construed in an analogous manner.

Four particular compounds which can be used are:

6-(1-methylimidazol-4-yl)methoxy-3-(5-methylisoxazol-3-yl)-1,2,4-
30 triazolo[3,4-a]phthalazine;

3-(5-methylisoxazol-3-yl)-6-(1-methyl-1,2,3-triazol-4-yl)methoxy-1,2,4-triazolo[3,4-a]phthalazine;

3-(5-methylisoxazol-3-yl)-6-(2-pyridyl)-1,2,4-triazolo[3,4-a]phthalazine;

and

5 3-(5-methylisoxazol-3-yl)-6-(1-methylimidazol-4-yl)-1,2,4-triazol-3-ylmethoxy-1,2,4-triazolo[3,4-a]phthalazine.

The second of the above compounds is particularly favoured.

The present invention also provides a pharmaceutical composition comprising an acetylcholinesterase inhibitor, an inverse agonist of the
10 GABA_A α_5 receptor subtype and a pharmaceutically acceptable carrier.

There is also provided a kit of parts comprising a first pharmaceutical composition comprising an acetylcholinesterase inhibitor and a first pharmaceutically acceptable carrier and a second pharmaceutical composition comprising an inverse agonist of the GABA_A
15 α_5 receptor subtype and a second pharmaceutically acceptable carrier for simultaneous, sequential or separate administration.

There is further provided a combination of an acetylcholinesterase inhibitor and an inverse agonist of the GABA_A α_5 receptor subtype for use in a method of treatment of the human body, particularly for the
20 treatment of a neurodegenerative disorder with associated cognitive deficit such as Alzheimer's Disease or Parkinson's disease, or of a cognitive deficit arising from a normal process such as aging or of an abnormal process such as injury. The combination is particularly beneficial in the treatment of Alzheimer's Disease.

25 There is also provided the use of a combination of an acetylcholinesterase inhibitor and an inverse agonist of the GABA_A α_5 receptor subtype in the manufacture of a medicament for the treatment of a neurodegenerative disorder such as Alzheimer's Disease or Parkinson's disease, or of a cognitive deficit arising from a normal process such as
30 aging or of an abnormal process such as injury. The treatment of Alzheimer's Disease is particularly preferred.

There is also disclosed a method of treatment of a subject suffering from a neurodegenerative disorder, such as Alzheimer's Disease or Parkinson's disease, or a cognitive deficit arising from a normal process such as aging or an abnormal process such as injury, which comprises
5 administering to that subject a therapeutically effective amount of a combination of an acetylcholinesterase inhibitor and an inverse agonist of the GABA_A α_5 receptor subtype. The treatment of Alzheimer's Disease is particularly preferred.

The pharmaceutical compositions of the present invention are
10 preferably in unit dosage forms such as tablets, pills, capsules, powders, granules, sterile parenteral solutions or suspensions, metered aerosol or liquid sprays, drops, ampoules, transdermal patches, auto-injector devices or suppositories; for oral, parenteral, intranasal, sublingual or rectal administration, or for administration by inhalation or insufflation. For
15 preparing solid compositions such as tablets, the principal active ingredient is mixed with a pharmaceutical carrier, e.g. conventional tableting ingredients such as corn starch, lactose, sucrose, sorbitol, talc, stearic acid, magnesium stearate, dicalcium phosphate or gums or surfactants such as sorbitan monooleate, polyethylene glycol, and other
20 pharmaceutical diluents, e.g. water, to form a solid preformulation composition containing a homogeneous mixture of a compound of the present invention, or a pharmaceutically acceptable salt thereof. When referring to these preformulation compositions as homogeneous, it is meant that the active ingredient is dispersed evenly throughout the
25 composition so that the composition may be readily subdivided into equally effective unit dosage forms such as tablets, pills and capsules. This solid preformulation composition is then subdivided into unit dosage forms of the type described above containing from 0.1 to about 500 mg of each active ingredient of the present invention. Typical unit dosage forms
30 contain from 1 to 100 mg, for example 1, 2, 5, 10, 25, 50 or 100 mg, of each active ingredient. The tablets or pills of the novel composition can be

coated or otherwise compounded to provide a dosage form affording the advantage of prolonged action. For example, the tablet or pill can comprise an inner dosage and an outer dosage component, the latter being in the form of an envelope over the former. The two components can be
5 separated by an enteric layer which serves to resist disintegration in the stomach and permits the inner component to pass intact into the duodenum or to be delayed in release. A variety of materials can be used for such enteric layers or coatings, such materials including a number of polymeric acids and mixtures of polymeric acids with such materials as
10 shellac, cetyl alcohol and cellulose acetate.

The liquid forms in which the novel compositions of the present invention may be incorporated for administration orally or by injection include aqueous solutions, suitably flavoured syrups, aqueous or oil suspensions, and flavoured emulsions with edible oils such as cottonseed
15 oil, sesame oil, coconut oil or peanut oil, as well as elixirs and similar pharmaceutical vehicles. Suitable dispersing or suspending agents for aqueous suspensions include synthetic and natural gums such as tragacanth, acacia, alginate, dextran, sodium carboxymethylcellulose, methylcellulose, polyvinyl-pyrrolidone or gelatin.

20 For the treatment of a neurodegenerative condition, a suitable dosage level is about 0.01 to 250 mg/kg per day, preferably about 0.01 to 100 mg/kg per day, and especially about 0.01 to 5 mg/kg of body weight per day of each active ingredient. The compounds may be administered on a regimen of 1 to 4 times per day. In some cases, however, dosage outside
25 these limits may be used.

The synergistic effect of the combination of the present invention can be shown, for example, by comparing the combined dosage of the combination with dosages of the same amount of each of the active ingredients separately on subjects using the Mini-Mental State
30 Examination (MMSE) as described in Folstein and Folstein J. Psychiat.

Res., 1975, 12, 189-198 or a variant thereof as discussed in Tombaugh and McIntyre, JAGS, 1992, 40, 922-935.

CLAIMS

1. A combination of an acetylcholinesterase inhibitor and an inverse agonist of the GABA_A α_5 receptor subtype for separate, sequential or
5 simultaneous administration.
2. A combination according to claim 1 wherein the inverse agonist has a functional efficacy at the α_5 receptor subtype of less than 20%, and a functional efficacy at the α_1 , α_2 and α_3 receptor subtypes of between -20
10 and +20 %.
3. A combination according to claim 1 or 2 wherein the inverse agonist has a binding ratio of greater than 10:1 to GABA_A receptors containing the α_5 receptor subtype compared to GABA_A receptors containing the α_1 , α_2
15 or α_3 subtypes.
4. A combination according to claim 1 wherein the inverse agonist is 3-(5-methylisoxazol-3-yl)-6-(1-methyl-1,2,3-triazol-4-yl)methoxy-1,2,4-triazolo[3,4-a]phthalazine.
20
5. A combination according to any one of the preceding claims wherein the acetylcholinesterase inhibitor is selected from physostigmine bromide, eptastigmine and its tartrate, galanthamine and its hydrochloride and hydrobromide, metrifonate, tacrine and its hydrochloride, eseridine and its
25 salicylate, suronacrine and its maleate, velnacrine and its maleate, amiridine and its hydrochloride, 7-methoxytacrine, SM-10888 and its citrate, phenserine and its tartrate, ENA-713, donepezil and its hydrochloride, TAK-147, CP-118954, huperzine A and zifrosilone.

6. A pharmaceutical composition comprising a combination as defined if any one of claims 1 to 5 and a pharmaceutically acceptable carrier for simultaneous administration.
- 5 7. A kit of parts comprising a first pharmaceutical composition comprising an acetylcholinesterase inhibitor and a first pharmaceutically acceptable carrier and a second pharmaceutical composition comprising an inverse agonist of the GABA_A α_5 receptor subtype and a second pharmaceutically acceptable carrier for simultaneous, separate or
10 sequential administration.
8. A method of treatment of a subject suffering from a neurodegenerative disorder or a cognitive deficit comprising administering to that subject a therapeutically effective amount of a combination of an
15 acetylcholinesterase inhibitor and an inverse agonist of the GABA_A α_5 receptor subtype.