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WO 03/013529 A1

(54) Title: PAROXETINE GLYCYRRHIZINATE

(57) Abstract: A salt formed from paroxetine hydrochloride and ammonium glycyrrhizinate masks the bitter taste of paroxetine and has a distinctive liquorice flavour.

PAROXETINE GLYCYRRHIZINATE

The present invention relates to a novel compound, to processes for preparing it and to its use in treating medical disorders.

Pharmaceutical products with antidepressant and anti-Parkinson properties are described in US-A-3912743 and US-A-4007196. An especially important compound among those disclosed is paroxetine, the (-) *trans* isomer of 4-(4'-fluorophenyl)-3-(3',4'-methylenedioxy-phenoxy-methyl)-piperidine. This compound is used in therapy as the hydrochloride salt for the treatment and prophylaxis of *inter alia* depression, obsessive compulsive disorder (OCD) and panic.

We have now surprisingly discovered a novel salt of paroxetine with glycyrrhizinic acid which may be used as an alternative to the currently marketed hydrochloride.

According to the present invention there is provided paroxetine glycyrrhizinate as a novel compound.

A great advantage of the glycyrrhizinate salt in oral formulations is its intense flavour of sweet liquorice which provides a taste-masking effect to hide the bitterness of paroxetine. In fact, because of the intensity of the liquorice flavour, further flavourings may be desirable to modify the liquorice taste of the formulation.

In one aspect the novel salt of this invention is provided in non-crystalline form, which may be a solid or an oil. The oil is preferably absorbed on a solid carrier, especially a carrier that is usable as a component of a pharmaceutical composition.

In another aspect the novel salt of this invention is provided in crystalline form. When the crystalline form exists as more than one polymorph, each polymorph forms another aspect of this invention.

Paroxetine glycyrrhizinate may be prepared by contacting stoichiometric amounts of the acid and paroxetine free base. Preferably the base is in solution, more preferably both are in solution.

Most commonly used solvents are suitable for mobilising paroxetine free base, for example toluene, alcohols such as methanol, ethanol, propan-2-ol, esters such as ethyl acetate, ketones such as acetone and butanone, halogenated hydrocarbons such as dichloromethane, and ethers such as tetrahydrofuran and diethyl ether. The glycyrrhizinic acid is preferably added as an aqueous or ethanolic solution. The glycyrrhizinic acid may also be added in the form of a soluble salt, for example ammonium glycyrrhizinate, or the glycyrrhizinic acid salt of an amine, for example ethylamine or diethylamine.

The concentration of paroxetine base is preferably in the range 5 to 50% weight/volume, more preferably in the range 10 to 30%. The concentration of glycyrrhizinic acid is suitably in the same range. Elevated temperatures may be used to increase solubility.

The salt may be isolated in solid form by conventional means from a solution thereof obtained as above. For example, a non-crystalline salt may be prepared by precipitation from solution, spray drying and freeze drying of solutions, evaporating a solution to a glass, or vacuum drying of oils, or solidification of melts obtained from reaction of the free base and the acid.

A crystalline salt may be prepared by directly crystallising from a solvent in which the product has limited solubility, or by triturating or otherwise crystallising a non-crystalline salt. An improved yield of the salt is obtained by evaporation of some or all of the solvent or by crystallisation at elevated temperature followed by controlled cooling, preferably in stages. Careful control of precipitation temperature and seeding may be used to improve the reproducibility of the production process and the particle size distribution and form of the product. Individual polymorphs are preferably crystallized directly from a solution of the salt, although recrystallizing a solution of one polymorph using seeds of another polymorph may also be carried out.

An alternative method of preparing paroxetine glycyrrhizinate is to start with a salt of paroxetine with an organic acid, such as acetic acid or maleic acid, rather than using paroxetine free base. Use of another salt of paroxetine as a starting material is suitable for preparation of the crystalline salt or, if a volatile acid such as acetic acid is used, non-crystalline salts by methods that involve evaporation (such as freeze-drying and spray-drying).

We have found it particularly effective to combine paroxetine hydrochloride with ammonium glycyrrhizinate.

The salt may be obtained as a solvate, when during isolation from solution it becomes associated with the solvent in which it is dissolved. Any such solvate forms a further aspect of this invention. Solvates may be returned to the unsolvated salt by heating, for example by oven-drying, or by treatment with a displacement solvent which does not form a solvate.

Prior to the isolation of the paroxetine glycyrrhizinate, water may be removed from the solution containing the salt by azeotropic distillation to avoid the formation of hydrates or to obtain the product in anhydrous form. In that case, suitable solvents for the solution of the salt are those which form an azeotrope with water such as toluene and propan-2-ol. It should also be appreciated that mixtures of solvents can also be used to aid the azeotropic removal of water.

Paroxetine free base may be prepared according to the procedures generally outlined in US Patent No 4,007,196 and EP-B-0223403. Glycyrrhizinic acid is commercially available as the mono-ammonium, disodium and dipotassium salts.

The compounds of this invention may be used to treat and prevent the following disorders:

Alcoholism

Anxiety

Depression

Obsessive Compulsive Disorder

Panic Disorder	Chronic Pain
Obesity	Senile Dementia
Migraine	Bulimia
Anorexia	Social Phobia
Pre-Menstrual Syndrome (PMS)	Adolescent Depression
Trichotillomania	Dysthymia
Substance Abuse	

These disorders are herein after referred to as "the Disorders".

The present invention further provides a method for treating and/or preventing any one or more of the Disorders by administering an effective and/or prophylactic amount of a salt of the invention to a sufferer in need thereof.

The present invention further provides a pharmaceutical composition for use in the treatment and/or prevention of the Disorders which comprises an admixture of a salt of the invention with a pharmaceutically acceptable carrier.

The present invention also provides the use of a salt of the invention for treating and/or preventing the Disorders.

The present invention also provides the use of a salt of the invention in the manufacture of a medicament for treating and/or preventing the Disorders.

Most suitably the present invention is applied to the treatment of depression, OCD and panic.

Compositions containing the salt of this invention may be formulated for administration by any route, and examples are oral, sub-lingual, rectal, topical, parenteral, intravenous or intramuscular administration. Preparations may, if desired, be designed to give slow release of the paroxetine salt.

The medicaments may, for example, be in the form of tablets, capsules, sachets, vials, powders, granules, lozenges, reconstitutable powders, or liquid preparations, for example solutions or suspensions, or suppositories.

- 5 The composition is usually presented as a unit dose composition containing from 1 to 200mg of paroxetine calculated from the amount of salt on a free base basis, more usually from 5 to 100mg, for example 10 to 50mg such as 10, 12.5, 15, 20, 25, 30 or 40mg by a human patient. Most preferably unit doses contain 20mg of paroxetine calculated on a free base basis. Such a composition is normally taken from 1 to 6 times daily, for
10 example 2, 3 or 4 times daily so that the total amount of active agent administered is within the range 5 to 400mg of paroxetine calculated on a free base basis. Most preferably the unit dose is taken once a day.

The compositions of the invention are usually adapted for oral administration; preferred
15 unit dosage forms include tablets or capsules.

The compositions of this invention may be formulated by conventional methods of admixture such as blending, filling and compressing.

- 20 Suitable carriers for use in this invention include a diluent, a binder, a disintegrant, a colouring agent, a flavouring agent and/or preservative. These agents may be utilized in conventional manner, for example in a manner similar to that already used for marketed anti-depressant agents.

- 25 Specific examples of pharmaceutical compositions include those described EP-B-0223403 and US 4,007,196, in which the products of the present invention may be used as the active ingredients.

The following Examples illustrate the present invention:

30

Example 1 : preparation of tablets

INGREDIENTS	20 mg Tablet	30mg Tablet
Paroxetine Glycyrrhizinate	20.00 mg (calc. as free base)	30.0 mg (calc. as free base)
Dicalcium Phosphate (DCP)	83.34 mg	125.0 mg
Microcrystalline Cellulose	50.67 mg	76.0 mg
Sodium Starch Glycollate	8.34 mg	12.5 mg
Magnesium Stearate	1.67 mg	2.5 mg

Commercial source of the ingredients

	Dicalcium Phosphate Dihydrate	-	Emcompress or Ditab*
5	Microcrystalline Cellulose	-	Avicel PH 102*
	Sodium Starch Glycollate	-	Explotab.*

* Trade names

10 **Method**

1. Pass DCP through a screen and weigh it into a Planetary mixer.
2. Add 30 mesh Paroxetine Glycyrrhizinate to the bowl.
3. Add 20 mesh Avicel and Explotab and mix all the powders for 10 minutes.
4. Add magnesium stearate and mix for 5 minutes.

15

Tablet into Pentagonal Tablets using the following punches:

30 mg Tablet	9.5 mm	Circumcircle
20 mg Tablet	8.25 mm	Circumcircle

- 20 The tablets are made satisfactorily on a single punch or a Rotary press.

Example 2 : preparation of tablets

INGREDIENTS	10 mg Tablet	20 mg Tablet	30mg Tablet
Paroxetine Glycyrrhizinate..	10 mg (calc.as free base)	20 mg (calc.as free base)	30 mg (calc.as free base)
Sodium Starch Glycollate	2.98 mg	5.95 mg	8.93 mg
Granular Dicalcium Phosphate (DITAB) or Dicafos	158.88 mg	317.75 mg	476.63 mg
Magnesium Stearate	1.75 mg	3.50 mg	5.25 mg

5

Method

1. Paroxetine Glycyrrhizinate, Sodium Starch Glycollate and Dicalcium Phosphate Dihydrate are screened and mixed together in a suitable mixer.
(Planetary, Cuble or High Energy Shear mixer.)
- 10 2. Add Magnesium Stearate and compress it into a tablet using a single punch or Rotary Tablet machine.

CLAIMS

1. A paroxetine glycyrrhizinate salt.
- 5 2. A compound according to claim 1 in non-crystalline form.
3. A compound according to claim 1 in crystalline form.
4. A process for the preparation of a compound as claimed in claim 1 or 2 by
10 precipitation from a solution of a paroxetine glycyrrhizinate, spray drying or freeze
drying a solution of a paroxetine glycyrrhizinate, evaporating a solution of a paroxetine
glycyrrhizinate to a glass, or by vacuum drying of oils of a paroxetine glycyrrhizinate, or
solidification of melts of a paroxetine glycyrrhizinate.
- 15 5. A process for the preparation of a compound as claimed in claim 1 or 3 by
crystallization or re-crystallization from a solution of a paroxetine glycyrrhizinate.
6. A process according to claim 4 or 5 in which the solution, oil or melt of a
paroxetine glycyrrhizinate is prepared by treating paroxetine free base or an organic acid
salt thereof with glycyrrhizinic acid or an ammonium or amine salt thereof.
- 20 7. A method for treating and/or preventing any one or more of the Disorders by
administering an effective and/or prophylactic amount of a paroxetine glycyrrhizinate to a
sufferer in need thereof.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 02/08926

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61K31/4525

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, BIOSIS, EMBASE, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 95 15155 A (FRANCESE FRANCO ;OLDANI DIEGO (IT); MANESCHI MASSIMO (IT); SMITHKL) 8 June 1995 (1995-06-08) abstract page 1, line 24 - line 26 page 2, line 6 - line 8 page 2, line 14 claims 1-7	1-7
Y	--- DATABASE REGISTRY 'Online! CHEMICAL ABSTRACTS SERVICE, COLUMBUS, OHIO, US; RN:1405-86-3, "Glycyrrhizin" XP002225024 the whole document --- -/--	1-7

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

° Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

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Name and mailing address of the ISA

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INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 02/08926

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5 811 436 A (COOPER DAVID ET AL) 22 September 1998 (1998-09-22) claims 1-9 column 1, line 31 - line 39 column 1, line 50 - line 54 ----	1-7
A	DE 201 00 529 U (SYNTHON B.V., NIJMEGEN, NL) 13 June 2001 (2001-06-13) page 8, paragraph 2 ----	
A	US 4 393 200 A (MIYASHITA AKIRA ET AL) 12 July 1983 (1983-07-12) column 1, paragraph 2 ----	1-7
A	US 5 763 449 A (FAWZY ABDEL A ET AL) 9 June 1998 (1998-06-09) abstract ----	1-7
P,Y	WO 02 074238 A (LAVIPHARM LAB INC) 26 September 2002 (2002-09-26) claims 1-44 abstract -----	1-7

INTERNATIONAL SEARCH REPORT

International application No.
PCT/EP 02/08926

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

Although claim 7 is directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- The additional search fees were accompanied by the applicant's protest.
- No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 02/08926

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 9515155	A	08-06-1995	IT MI932540 A1	05-06-1995
			AU 693144 B2	25-06-1998
			AU 1219895 A	19-06-1995
			CA 2177721 A1	08-06-1995
			CN 1145586 A	19-03-1997
			WO 9515155 A1	08-06-1995
			EP 0804168 A1	05-11-1997
			JP 9505818 T	10-06-1997
			NZ 277238 A	27-04-1998
			ZA 9409567 A	10-10-1995
			US 5811436	A
AU 1536895 A	21-08-1995			
BG 62843 B1	29-09-2000			
BG 100763 A	31-03-1997			
BR 9507055 A	02-09-1997			
DE 69508924 D1	12-05-1999			
DE 69508924 T2	21-10-1999			
DK 742715 T3	18-10-1999			
EP 0742715 A1	20-11-1996			
FI 963051 A	01-08-1996			
GR 3030131 T3	30-07-1999			
HK 1012288 A1	12-05-2000			
JP 9508402 T	26-08-1997			
NO 963244 A	02-08-1996			
NZ 278891 A	26-01-1998			
PL 315679 A1	25-11-1996			
RO 116342 B	30-01-2001			
RU 2136281 C1	10-09-1999			
SI 742715 T1	31-08-1999			
SK 100496 A3	04-12-1996			
AP 536 A	26-09-1996			
AP 611 A	03-09-1997			
AT 178489 T	15-04-1999			
CN 1140411 A ,B	15-01-1997			
CZ 9602293 A3	15-01-1997			
WO 9520964 A1	10-08-1995			
ES 2129806 T3	16-06-1999			
HU 75941 A2	28-05-1997			
IL 112521 A	22-09-1999			
TW 436296 B	28-05-2001			
ZA 9500776 A	01-08-1996			
DE 20100529	U	10-05-2001	DE 20100529 U1	10-05-2001
			WO 02055062 A2	18-07-2002
US 4393200	A	12-07-1983	JP 56113793 A	07-09-1981
			JP 1610300 C	15-07-1991
			JP 2022080 B	17-05-1990
			JP 56115797 A	11-09-1981
			FR 2473526 A1	17-07-1981
			GB 2071665 A ,B	23-09-1981
US 5763449	A	09-06-1998	AU 727175 B2	07-12-2000
			AU 3913297 A	25-02-1998
			EP 0938302 A1	01-09-1999
			JP 2000505093 T	25-04-2000
			US 5962461 A	05-10-1999

INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

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Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5763449	A	WO 9805312 A1	12-02-1998
WO 02074238	A	26-09-2002 WO 02074238 A2 US 2002147201 A1	26-09-2002 10-10-2002