



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : A61K 38/13	A1	(11) International Publication Number: WO 97/35603 (43) International Publication Date: 2 October 1997 (02.10.97)
(21) International Application Number: PCT/US97/04627 (22) International Filing Date: 21 March 1997 (21.03.97) (30) Priority Data: 08/622,516 25 March 1996 (25.03.96) US (60) Parent Application or Grant (63) Related by Continuation US 08/622,516 (CIP) Filed on 25 March 1996 (25.03.96) (71) Applicant (for all designated States except US): SANGSTAT MEDICAL CORPORATION [US/US]; 1505-B Adams Drive, Menlo Park, CA 94025 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): FLOC'H, Robert [FR/FR]; 9, avenue d'Auray, F-44300 Nantes (FR). MERLE, Christian [FR/FR]; 7, impasse Sainte-Radégonde, F-86000 Poitiers (FR). (74) Agents: TRECARTIN, Richard, F. et al.; Flehr, Hohbach, Test, Albritton & Herbert L.L.P., Suite 3400, 4 Embarcadero Center, San Francisco, CA 94111-4187 (US).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: CYCLOSPORIN A FORMULATIONS AS NANOPARTICLES		
(57) Abstract Cyclosporin A formulations are provided as amorphous nanoparticle dispersions for physiologic absorption. The compositions have high bioavailability and patient acceptability. By providing for concentrates comprising lower alkanols and a polyoxyalkylene surfactant as a stable dispersion of cyclosporin A, upon introducing the stable dispersion into an aqueous medium, the subject formulation is produced comprising amorphous bioavailable cyclosporin nanoparticles.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

CYCLOSPORIN A FORMULATIONS AS NANOPARTICLES

BACKGROUND OF THE INVENTION

The drug cyclosporin A, despite its many shortcomings and the difficulties in formulation, variations in bioavailability, and side effects, has proven to be one of the great success stories of the drug industry. Because of cyclosporin A's

5 hydrophobicity, formulations of cyclosporin A must take into account the need for a stable dispersion of the cyclosporin A, as well as the manner of administration of the formulation. For example, if it is intended that the formulation be diluted with water prior to its being taken orally, the resulting composition must provide the cyclosporin A in a bioavailable form, where adverse effects are not enhanced,

10 preferably diminished. The cyclosporin A which will come out of solution should be dispersible, so that the dosage is repeatable. Alternatively, where the formulation is provided in a manner where the cyclosporin A formulation becomes diluted with gastric juices, such as the use of capsules, it is essential that the cyclosporin A retains its bioavailability and activity in the environment of the

15 gastric juices. In all events, the cyclosporin A must be able to be transported into the vascular system, where it can diminish the immune response.

It is therefore of interest to develop formulations which are organoleptically acceptable, provide for desirable levels of bioavailability, do not introduce adverse effects associated with cyclosporin A, and generally fulfill the requirements of

20 therapeutic formulations.

BRIEF SUMMARY OF THE INVENTION

Aqueous dispersions of cyclosporin are provided by introducing a stable dispersion of cyclosporin in a formulation comprising as co-solvents a lower alkanol and a polyoxyalkylene surfactant, and desirably a polyethylene glycol, as
5 co-solvent. Upon dilution of the stable dispersion, an aqueous dispersion is obtained comprising nanoparticles of cyclosporin in amorphous form having good bioavailability.

DETAILED DESCRIPTION

10 Methods and compositions are provided for producing an aqueous colloidal dispersion of cyclosporin nanoparticles having good bioavailability. The nanoparticles are substantially spheric, the cyclosporin is present in an amorphous form, and the average size will generally be less than about 1000 nm, greater than about 50 nm, generally in the range of about 200 - 800 nm, usually in the range of
15 about 200-600 nm. Generally, at least about 50 weight percent of the total weight of cyclosporin will be present as particles in the indicated size range. Larger particles may be present, particularly as aggregates of nanoparticles, where the average diameter will usually be less than about 50 μm , more usually less than about 25 μm , the aggregates usually not exceeding 40 weight % of the total
20 cyclosporin..

The amount of cyclosporin A amorphous particles in the composition will be sufficient for therapeutic effect. Since the formulation may be formed by introduction into an aqueous medium prior to administration or directly into the gastric juices, the particular concentration cannot be stated, since the dilution in the
25 stomach is uncertain. For preparation in an aqueous medium prior to oral administration, generally, the cyclosporin will be present at a weight percent of about 0.01-2.5, more usually from about 0.01-0.5 weight percent. The temperature of mixing may be in the range of about 10 to 50°C, usually in the range of about 20 to 40°C. Usually the mixing will involve stirring for sufficient
30 time to provide the solution of the cyclosporin.

The colloidal amorphous suspension of the nanoparticles is sufficiently stable to allow for some standing prior to administration, frequently up to about 6 hours, more frequently up to about 3 hours.

While cyclosporin A finds primary use, any of the cyclosporins which are physiologically acceptable, e.g. A through Z, may be employed.

The amorphous cyclosporin colloidal dispersion may be produced by preparing a stable dispersion of cyclosporin in a lower alkanol and a
5 polyoxyalkylene compound, either ester or alcohol. The alkanols will be ethanol or propylene glycol, individually or in combination, particularly where ethanol will be present in the range of about 25-75 volume percent, when the combination of alkanols is employed. The particular manner in which the colloidal dispersion is produced is not critical, so long as the materials used in the stable dispersion are
10 physiologically acceptable, do not interfere with the activity of the cyclosporin, and are readily available.

Various polyalkyleneoxy compounds may be employed which may serve as surfactants and co-solvents with the lower alkanols. The polyalkyleneoxy compounds are, therefore, liquids, soluble in both water and lower alkanols, have
15 low toxicity and in conjunction with the lower alkanols are capable of maintaining a stable dispersion, usually a solution of cyclosporin A. Exemplary of polyoxyethylene surfactants are polyoxyethylene esters, such as polyoxyethylene substituted sorbitan esterified with a fatty acid of from 12-18 carbon atoms, more usually from about 16-18 carbon atoms, exemplified by polysorbate 80. The
20 number of oxyethylene groups will generally be from about from 10-30. Exemplary of polyoxyalkylene compounds as cosolvents are polyethylene glycols of an average molecular weight of less than about 2000, preferably less than about 1000, at least about 300, more usually in the range of about 300-700 particularly from about 350-500 kiloDaltons. Generally, greater than 50% by weight of the
25 polyethylene glycol will be within 50% of the average molecular weight of the polyethylene glycol.

In the formulation, the total amount of lower alkanol will generally be in the range of about 25-60 weight percent, more usually in the range of about 30-50 weight percent. The total amount of alkyleneoxy compound(s) will generally be in
30 the range of about 20-50 weight percent, more usually in the range of about 25-40 weight percent. Where combinations of polyoxyalkylene compounds are employed, the amount of the fatty acid ester will generally range from about 25-100% of the polyoxyalkylene compounds.

The weight of cyclosporin in the formulation will be sufficient to provide for a therapeutic dosage, generally in the range of about 2.5 to 25 weight percent, more usually in the range of about 5-15 weight percent.

The subject compositions may be prepared by first dissolving the cyclosporin in the lower alkanol, where a small proportion of the polyoxyalkylene compound may also be included, generally less than about 50 weight percent of the composition used for dissolving the cyclosporin. An elevated temperature may be employed, usually in the range of about 60 to 90°C. After dissolving the cyclosporin, the major proportion of the polyalkyleneoxy compound may be added and the total formulation brought to the desired ratios by the addition of the appropriate components. Generally, the cyclosporin can be dissolved in the lower alkanol (optionally including a portion of the polyalkyleneoxy compound) at a weight ratio of about 1:1.5-5, more usually 1:2-4.

The subject formulations may be used in accordance with conventional ways already described in the literature. Oral formulations have been reported in U.S. Patent Nos. 4,388,307; and 5,342,625; and UK Patent No. 2,222,770B, whose disclosures are incorporated herein by reference as describing the use of cyclosporin in oral formulations. Thus, the subject compositions may be administered as liquid solutions, capsules, or the like, taken orally in single or multiple dosages, as therapeutically required in accordance with conventional procedures. The formulations are used with patients who require that they be immunocompromised, as in the case of transplantation, autoimmune diseases and the like.

For convenience of the user, kits may be provided having the appropriate amount of cyclosporin, one or more dosage levels and the cosolvents, namely the lower alkanol(s) and the polyalkyleneoxy compound(s), e.g. at least one of ethanol and propylene glycol and at least one of polysorbate 80 and PEG400.

The following examples are offered by way of illustration and not by way of limitation.

30

Example 1

5 g of cyclosporin A was added to 5 mL of ethanol. The mixture was stirred to complete dissolution of cyclosporin A. To the resulting solution were

added 25 g of polysorbate 80 and the volume is completed to 50 mL by 1,2-propylene glycol. The mixture was sufficiently stirred at room temperature until a homogeneous solution was formed.

5 Example 2

5 g of cyclosporin A was added to 5 mL of ethanol. The mixture was stirred until complete dissolution of cyclosporin A. To the resulting solution were added 15 g of polysorbate 80 and the volume is completed to 50 mL by a mixture of 1,2-propylene glycol and polyethylene glycol 400. The mixture was sufficiently
10 stirred at room temperature until a homogeneous solution was formed.

Example 3

1 mL of the solution obtained in example 1 was added in 50 mL of water with a glass syringe as recommended for the oral administration of concentrated
15 emulsions or microemulsions in human. The addition of the solution was followed by a quick dissolution and a white suspension of fine particles was obtained having a blue reflect as colloidal suspensions (Tyndall effect). After centrifugation at 26,000 g during 5 hours, the sediment was washed with water and then centrifuged at 26,000 g during 24 hours. The washing and centrifugation processes were
20 repeated twice under the same conditions. After drying, an x-ray powder diagram was performed. The solid was exclusively in amorphous form.

The sediment was examined by scanning electron microscopy. The sediment was constituted of amorphous spheric nanoparticles with a diameter between 200 and 400 nm with the presence of some aggregates.

25 2 mL of the solution obtained in example 1 was added in 100 mL of water and the colloidal suspension was examined 10 minutes and 1 hour after the dilution by a diffraction/diffusion laser granulometer (Malvern SB.OD).

After 1 hour, two particle populations were observed: one representing 70% of the weight of cyclosporin A with an average diameter of 300 nm and a second
30 one representing 30% of the weight of cyclosporin A with an average diameter of 20 μ m, probably constituting aggregates of nanoparticles.

Example 4

1 mL of the solution obtained in example 1 was added to 50 mL of water and the colloidal suspension was stirred during 10 minutes.

5 The suspension was then added to 200 mL of artificial acidic gastric juice and warmed at 37°C. The homogeneous colloidal suspension was examined by diffraction/diffusion laser granulometry (Malvern SB.OD). The suspension was constituted exclusively of nanoparticles with an average diameter of 600 nm.

Example 5

10 1 mL of the solution obtained in example 1 was added directly to 200 mL of artificial acidic gastric juice.

The homogeneous suspension was warmed at 37°C and examined rapidly by diffraction/diffusion laser granulometry (Malvern SB.OD). The suspension was exclusively constituted of nanoparticles with an average diameter of 350 nm.

15

The subject cyclosporin compositions provide for excellent bioavailability in being amorphous particles, small, so as to have high surface area, and without detrimental effects other than those conventionally found with cyclosporin.

20 All publications and patent applications mentioned in this specification are herein incorporated by reference to the same extent as if each individual publication or patent application was specifically and individually indicated to be incorporated by reference.

25 The invention now being fully described, it will be apparent to one of ordinary skill in the art that many changes and modifications can be made thereto without departing from the spirit or scope of the appended claims.

WHAT IS CLAIMED IS:

1. An aqueous dispersion of cyclosporin nanoparticles, wherein at least 50 weight percent of the cyclosporin present in the dispersion is of particles less than about 1 μm , said cyclosporin being amorphous.
5
2. A dispersion according to Claim 1, comprising in minor amounts lower alkanol and at least one polyoxyethylene surfactant.
- 10 3. A dispersion according to Claim 2, wherein said polyoxyethylene surfactant is polysorbate 80.
4. A dispersion according to Claim 2, wherein said lower alkanol is at least one of ethanol and propylene glycol.
15
5. A dispersion according to Claim 1, comprising a polyethylene glycol of less than about 2000 Daltons.
6. In a method for orally administering cyclosporin to a patient, the improvement which comprises:
20 providing said cyclosporin, wherein at least 50 weight % of said cyclosporin is as amorphous particles of less than about 1000nm.
7. A method according to Claim 6, wherein said providing comprises adding to an aqueous medium a composition comprising cyclosporin dispersed in a combination of lower alkanol consisting of at least one of ethanol and propylene glycol and a polyoxyethylene surfactant.
25
8. A method according to Claim 7, wherein said polyoxyethylene surfactant is polysorbate 80.
30
9. A method according to Claim 7, wherein said lower alkanol is present in from about 25 to 60 weight percent, said polyoxyalkylene surfactant is

present in from about 20 to 50 weight percent, and said cyclosporin is present in from about 2.5 to 25 weight percent.

10. A method according to Claim 7, wherein said composition further
5 comprises polyoxyethylene cosolvent of less than 2000 Daltons.

11. A kit comprising cyclosporin, at least one of ethanol and propylene glycol, and polysorbate 80.

10 12. A kit according to Claim 11, further comprising PEG400.

13. A method for preparing a formulation according to Claim 1
comprising:

combining at least one of ethanol and propylene glycol with cyclosporin A
15 to from a solution; and

combining said solution with a polyethyleneoxy surfactant to form a second solution, which upon dilution with water forms amorphous nanoparticles of said cyclosporin A.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US97/04627

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : A61K 38/13
US CL : 424/450; 514/11

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)

U.S. : 424/450; 514/9, 11, 938; 530/317, 321

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 practicable, search terms used)

Please See Extra Sheet.

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 4,996,193 A (HEWITT et al) 26 February 1991 (26/02/91), see column 4, lines 16-23, column 15, lines 1-4.	11, 13
X	US 5,047,396 A (ORBAN et al) 10 September 1991 (10/09/91), see column 2, lines 27-35 and 66-68, column 3, lines 1-10 and 52-59, column 4, line 46.	1, 2, 4 - 7, 9, 10, 13
X	US 5,154,930 A (POPESCU et al) 13 October 1992 (13/10/92), see column 9, lines 44-58, column 12, lines 18-31, Example 3.	1, 5, 6
X	US 5,206,219 A (A. DESAI) 27 April 1993 (27/04/93), see column 2, lines 15-16, 23-29, 51-53, column 3, line 56 - column 4, line 20, examples 1-5.	11, 12

Further documents are listed in the continuation of Box C. See patent family annex.

* Special categories of cited documents:	*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
A document defining the general state of the art which is not considered to be of particular relevance	*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
E earlier document published on or after the international filing date	*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
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)	*Z*	document member of the same patent family
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		

Date of the actual completion of the international search

23 APRIL 1997

Date of mailing of the international search report

06 JUN 1997

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Authorized officer

JEFFREY E. RUSSEL

Facsimile No. (703) 305-3230

Telephone No. (703) 308-0196

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US97/04627

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5,474,979 A (DING et al) 12 December 1995 (12/12/95), see column 3, lines 58-63, example 1, column 5, lines 29-32.	1-3,6
X,P	US 5,583,105 A (KOVACS et al) 10 December 1996 (10/12/96), see column 2, line 49 - column 3, line 6, column 3, lines 26-32.	1,2,4,6,7
X,P	US 5,540,931 A (HEWITT et al) 30 July 1996 (30/07/96), see column 6, lines 37-43, column 7, lines 3-8.	1-4, 6-8, 11

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US97/04627

B. FIELDS SEARCHED

Electronic data bases consulted (Name of data base and where practicable terms used):

APS, DIALOG

search terms: cyclosporin, ethanol, propylene glycol, polyoxyethylene sorbitan, TWEEN, dispersion, emulsion, nanoemulsion, nanoparticle, microparticle, oral, amorphous, noncrystalline