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(56) Related Art  
**US 5948755 A**  
**WATASHI KOICHI et al., Hepatology, 2003, vol. 38 (5), pages 1282-1288**  
**NAKAGAWA MET AL. Biochemical and Biophysical Research Communications, 2004, vol. 313(1), pages 42-47**  
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**WO 2000/001715 A1**  
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(54) Title: USE OF [D-MEALA]3-[ETVAL]4-CYCLOSPORIN FOR THE TREATMENT OF HEPATITIS C INFECTION AND PHARMACEUTICAL COMPOSITION COMPRISING SAID [D-MEALA]3-[ETVAL]4-CYCLOSPORIN

(57) Abstract: This invention relates to the use in the treatment of HCV infection, either as single active agents or in combination with another active agent, of a cyclosporin having increased cyclophilin binding activity and essentially lacking immunosuppressive activity.

USE OF [D-MEALA]3-[ETVAL]4-CYCLOSPORIN FOR THE TREATMENT OF HEPATITIS C INFECTION  
AND PHARMACEUTICAL COMPOSITION COMPRISING SAID [D-MEALA]3-[ETVAL]4-CYCLOSPORIN

The present invention relates to the use of a cyclosporin for the treatment of  
5 hepatitis C virus (HCV) infection and to a pharmaceutical composition comprising  
said cyclosporin.

HCV was cloned and characterized about 15 years ago by Choo and  
colleagues (see *Science* 244, (1989), 359-362). HCV belongs to the family  
10 Flaviviridae and comprises an enveloped nucleocapsid and a single-stranded RNA  
genome of positive polarity (see Bartenschlager et al., *Antiviral Res.* 60, (2003),  
91-102). HCV is transmitted primarily by blood, blood products and vertical  
transmission during pregnancy. Introduction of diagnostic tests for screening blood  
products has significantly reduced the rate of new infection.

15 Still, HCV remains a serious medical problem. There are currently about  
170 million people infected with HCV. The initial course of infection is typically  
mild. However, the immune system is often incapable of clearing the virus, and  
people with persistent infections are at a high risk for liver cirrhosis and  
20 hepatocellular carcinoma (see Poynard et al., *Lancet* 349, (1997), 825-832).

There is no vaccine available, and therapeutic options are very limited (see  
Manns et al., *Indian J. Gastroenterol.* 20 (Suppl. 1), (2001), C47-51; Tan et al.,  
Nat. Rev. Drug Discov. 1, (2002), 867-881).

25 Current therapy is based on a combination of interferon alpha and ribavirin.  
This therapy produces a sustained anti-viral response in 85-90% of patients  
infected with genotypes 2 and 3, but, unfortunately, only in about 45% of patients  
infected with the prevalent genotype 1. Furthermore, side effects are significant  
30 and include myalgia, arthralgia, headache, fever, severe depression, leucopenia  
and haemolytic anaemia.

Clearly, additional therapies, with a higher antiviral activity and a better  
safety profile, are required for the treatment of HCV infection, particularly e.g. in

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the case of the prevention of HCV recurrence. In order to establish the safety profile, criteria such as low cytotoxicity and cytostatic effect and high selectivity index are particularly relevant for clinical treatment of HCV infection.

5 A novel approach for the treatment of HCV infection using cyclosporins was recently described by clinical observations (see Teraoka et al., Transplant Proc., 1988, 20 (3 suppl 3), 868-876, and Inoue et al. J Gastroenterol, 2003, 38, 567-572). Recently it was shown that Cyclosporin A (CsA) inhibited the *in vitro* intracellular replication of an HCV subgenomic replicon at clinically achievable 10 drug concentrations (see Watashi et al., Hepatology 38, 2003, 1282-1288, and Nakagawa et al., BBRC 313, 2004, 42-47). Both groups suggested that the anti-HCV effect of CsA was not associated with immunosuppressive activity based on 15 observations made with the use respectively of an immunosuppressive macrolide, i.e. the compound known under the name FK 506 and a non-immunosuppressive Cyclosporin A derivative, i.e. the compound known under the name NIM 811 or [Melle]<sup>4</sup>-CsA. Nakagawa et al. consider that expanding applications of CsA may cause substantial problems due to its well-known immunosuppressive properties and suggest that one solution to overcome this problem would be to consider the use of non-immunosuppressive cyclosporin analogs.

20

During the last 15 years, a number of medicinal chemistry studies have been conducted with the aim to identify such non-immunosuppressive cyclosporin analogs and compound NIM 811 is one of the most representative compounds having such a property.

25

NIM 811, along with 9 other Cyclosporin A derivatives, were reported by Ko et al. in patent application EP 0 484 281 for their non-immunosuppressive properties and were considered as being potentially useful in the treatment of HIV infection and the prevention of AIDS. The design of those derivatives involved the 30 modification of the amino-acids in 4- and/or 5-positions of Cyclosporin A.

By modifying amino-acids in 2- and/or 6-positions of Cyclosporin A, Sigal et al. synthesised a total of 61 cyclosporin analogs and observed that such chemical

modifications induce a decrease in the immunosuppressive activity (see Sigal *et al.*, *J. Exp. Med.*, 173, 1991, 619-628).

Further attempts for modifying amino-acid in 3-position of Cyclosporin A in order to obtain non-immunosuppressive compounds were described in particular by Barrière *et al.*, in WO 98/28328, WO98/28329, and WO98/28330.

Wenger *et al.* have designed a series of compounds that differ from Cyclosporin A in position 3, in which they contain an N-methylated, nonbulky hydrophobic or neutral amino acid other than a glycine, and in position 4, in which they contain an N-methylated or N-ethylated hydrophobic or neutral amino acid other than a leucine and they report that those compounds have a high potency to inhibit HIV-1 replication and essentially lack immunosuppressive activity (see International patent application WO 00/01715 and *Tetrahedron Lett.*, 41, (2000), 7193-6).

The aim of the present invention is to provide the clinician with a new therapy for the treatment of HCV infection, particularly e.g. in the case of the prevention of HCV recurrence. This therapy should offer a higher antiviral activity and a better safety profile in comparison to the already approved therapy or the newly proposed ones.

The present inventors surprisingly found that the administration to a patient infected with HCV of a very specific compound, i.e. [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA, meets the above requirements. They observed that, in addition to its non-immunosuppressive property, [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA has a significantly increased affinity for cyclophilins, which increased affinity is correlated with an elevated efficacy against inhibition of HCV replication.

Accordingly, one of the subject-matters of the present invention relates to the use of [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA for the manufacture of a medicinal product intended for the treatment of HCV infection or the prevention of HCV recurrence in a patient.

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[D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA has been reported by Wenger et al. in WO 00/01715 and it has been attributed the CAS Registry Number 254435-95-5. It is a cyclic undecapeptide described by the following formula:

5 -MeBmt- $\alpha$ Abu-D-MeAla-EtVal-Val-MeLeu-Ala-(D)Ala-MeLeu-MeLeu-MeVal-  
1      2      3      4      5      6      7      8      9      10      11

where MeBmt is N-methyl-(4R)-4-but-2E-en-1-yl-4-methyl-(L)threonine,  $\alpha$ Abu is L- $\alpha$ -aminobutyric acid, D-MeAla is N-methyl-D-alanine, EtVal is N-ethyl-L-valine, Val is L-valine, MeLeu is N-methyl-L-leucine, Ala is L-alanine, (D)Ala is D-alanine, and MeVal is N-methyl-L-valine. The conventional numbering of amino acid positions generally used in reference of Cyclosporin A is shown below the formula. This is achieved by using composite names comprising a first portion indicating the identity of residues that are different from those in cyclosporin A and providing their position, and a second portion labelled "CsA" indicating that all other residues are identical to those in Cyclosporin A. For example, [Melle]<sup>4</sup>-CsA is a cyclosporin that is identical to cyclosporin A except that MeLeu in position 4 is replaced by Melle (N-methyl-L-isoleucine).

20 The present invention will be explained further with following experiments and with the drawing in which:

- Fig. 1 represents a dose response histogram measured by luciferase assay in infected Huh-7-Lunet cells;  
- Fig. 2 represents a dose response histogram measured by luciferase assay in infected Huh-7.5 cells;  
- Fig. 3 represents clearance response curves of infected Huh-9-13 cells;  
- Fig. 4 represents 3D representation of dose response with combination IFN / [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA.

30 Current medical uses of Cyclosporin A relate to the ability of this compound to suppress the cell-mediated immune response by preventing production and release of several autocrine T-cell growth factors, including interleukin 2 (IL-2), from activated T cells (see Borel (1989) Transplant. Proceed. 21, 810-815; Kronke et al. (1984) Proc. Natl. Acad. Sci. USA 81, 5214-5218; Faulds et al. (1993) Drugs

45, 953-1040). Upon entry into cells, Cyclosporin A binds to cyclophilins with high affinity (see Handschumacher et al. (1984) *Science* 226, 544-547). As among different biological function, they have peptidyl-prolyl cis-trans isomerase (PPIase) activity that can be measured *in vitro* (see Fischer et al. (1989) *Nature* 337, 476-5 478; Takahashi et al. (1989) *Nature* 337, 473-475). Critical for the immunosuppressive effect of cyclosporin A is an interaction between cyclophilin-Cyclosporin A complex and calcium- and calmodulin-dependent serine/threonine phosphatase 2B (calcineurin) (see Hauske (1993) *DN&P* 6, 705-711, Friedman et al. (1991) *Cell* 66, 799-806; Liu et al. (1991) *Cell* 66, 807-815). Formation of this 10 ternary complex results in an inhibition of the phosphatase activity of calcineurin. (see Jain et al. (1993) *Nature* 365, 352-355; Rao et al. (1997) *Annu. Rev. Immunol.* 15, 707-747; Crabtree (1999) *Cell* 96, 611-614). Calcineurin promotes the selective dephosphorylation of NF-AT that then translocates to the nucleus where it associates with activator protein 1 and transactivates target genes, 15 including the IL-2 gene.

It is believed that, due to the amino-acids in 3- and 4-positions, [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA has a dramatically reduced ability to interact with calcineurin as shown by transcriptional and immunological assays as well as a 20 significantly increased affinity for cyclophilins as indicated by assays of inhibition of peptidyl-prolyl cis-trans isomerase activity.

Peptidyl-prolyl cis-trans isomerase (PPIase) activity of cyclophilins was determined using a procedure adapted from Kofron et al. (see *Biochemistry* 30, 25 6127-6134 (1991) ; *J. Am. Chem. Soc.* 114, 2670-2675 (1992)). N-succinylated Ala-Ala-Pro-Phe-para nitro-aniline (Suc-AAPF-pNA, Bachem, Bubendorf, Switzerland) was used as the substrate. The assay was based on the preferential chymotrypsin cleavage of the trans isoform of the Phe-pNA bond in the tetrapeptide Ala-Ala-Pro-Phe-pNA. This cleavage liberates the para-nitroaniline moiety that can be detected and quantitated at 390 nm ( $\epsilon = 11,814 \text{ M}^{-1}\text{cm}^{-1}$ ). Schutkowski et al. (1995) *Biochemistry* 34, 13016-13026. Cis-trans isomerization is catalysed by cyclophilin (PPIase, EC 5.2.1.8). After mixing CsA or another cyclosporin ( $10^{-9} - 2 \times 10^{-5}$  M final concentrations prepared from 1000-fold concentrated stock solutions in ethanol) with 0.1  $\mu\text{g}$  cyclophilin (Sigma) in a total 30

volume of 1.5 ml of 40 mM Hepes, pH 7.9, and incubation for 50 min on ice, the reaction mixture was transferred to a cuvette that was kept at 10°C in a Varian spectrophotometer (Varian). Subsequent to the addition of 3.75 mg of chymotrypsin (70  $\mu$ l of a solution of chymotrypsin in 10 mM HCl), the reaction was 5 initiated by addition of 10  $\mu$ l of a 3.2 mM solution of Suc-AAPF-pNA in 0.5 M LiCl/trifluoroethanol. The reaction was monitored for 3 min, and an initial rate constant was determined from the data obtained. As a control, an initial rate constant was also determined for a parallel reaction that lacked cyclophilin. Concentration-response curves were established for cyclosporin A and other 10 cyclosporins, and IC<sub>50</sub> (50% inhibitory concentration) values of different cyclosporins were expressed relative to that of cyclosporin A (1.0). A value less than 1 means that the compound has an higher cyclophilin affinity than CsA.

A NF-AT-dependent reporter assay was used initially to estimate 15 immunosuppressive activities of cyclosporins. Baumann et al. (1992) Transplant. Proc. 24, 43-48. Jurkat T cells stably transfected with a reporter construct containing a bacterial  $\beta$ -galactosidase gene under the control of a promoter of an IL-2 gene were obtained from G. Zenke, Novartis Pharma AG, Basel, Switzerland. The cells were grown in RPMI1640 medium supplemented with 20 10% heat-inactivated fetal calf serum, 100 u/ml penicillin, 100  $\mu$ g/ml streptomycin, 2 mM glutamine, 50  $\mu$ M 2-mercaptoethanol and 100 u/ml hygromycin B. The cells were stimulated by the addition of 2.4  $\mu$ M phorbol-12-myristate-13-acetate and 75  $\mu$ g/ml phytohemagglutinin in the presence or absence of cyclosporin A or another cyclosporin (10<sup>-9</sup> – 2x10<sup>-5</sup> M final concentrations prepared from 1000-fold 25 concentrated stock solutions in ethanol). Subsequent to incubation for 20 h at 37°C, cells were harvested and lysed in 50 mM Na<sub>2</sub>HPO<sub>4</sub> (pH 9.0), 10 mM KCl, 1 mM MgSO<sub>4</sub>, 1% Triton X-100, 0.5 mM 4-methylumbelliferyl- $\beta$ -D-galactoside (Sigma, Buchs, Switzerland). The  $\beta$ -galactosidase reaction was allowed to proceed for 1 h in the dark at room temperature. Fluorescent 4-methyl- 30 umbelliferone was assayed fluorometrically in the supernatant solution (excitation: 355 nm; emission: 460 nm). Concentration-response curves were established for cyclosporin A and other cyclosporins, and the IC<sub>50</sub> values of different cyclosporins were calculated relative to that of cyclosporin A (1.0). A value higher than 1 means that the compound is less immunosuppressive than CsA.

Example results are shown in Table 1 below.

Table 1: Cyclophilin binding (PPIase) and immunosuppressive (IL-2) activities of

5 CsA and other cyclosporins

Compound	PPIase	IL-2
CsA	1.0	1.0
[D-MeAla] <sup>3</sup> -[EtVal] <sup>4</sup> -CsA	0.3	7161
[Melle] <sup>4</sup> -CsA	0.5	2250

The data shown in Table 1 revealed that certain substitutions in position 4 (i.e., Val, Ile) dramatically reduced immunosuppressive activity (measured as inhibition of IL-2 expression) as well as detectably enhanced cyclophilin binding activity  
10 (measured as inhibition of PPIase activity of cyclophilin). Substitution in position 3 resulted in a substantial further increase in cyclophilin binding activity (by 2 fold or more; cf [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA. It had a higher cyclophilin binding activity and a lower residual immunosuppressive activity than [Melle]<sup>4</sup>-CsA, the best reference compound available from the literature. [Melle]<sup>4</sup>-CsA is also known as NIM811.

15

Non-immunosuppressive activity of [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA

In a confirmatory analysis, immunosuppressive activities of CsA, [Melle]<sup>4</sup>-CsA and [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA were estimated using the mixed lymphocyte reaction. In this assay, cyclosporins were dissolved in ethanol (10 mg/ml). Freshly isolated CD4<sup>+</sup> PBMCs from two healthy donors were mixed subsequent to inactivation by irradiation of one of the populations (stimulator cells; S). After five days of co-culture in the presence or absence of a cyclosporin (1 µg/ml), the proliferative response of the non-inactivated cell population (responder cells; R)  
20 was determined by [<sup>3</sup>H]-thymidine incorporation.  
25

The assay was conducted reciprocally with the two cell populations, each being inactivated and stimulated in turn. Stimulation (%) of responder cells was calculated by the formula:

30

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Percent stimulation =  $100 \times (\text{sample with cyclosporin} - \text{background}) / (\text{sample without cyclosporin} - \text{background})$

Sample refers to a mixture of stimulator and responder cells. Background

5 represents a control in which only stimulator cells are mixed. Results are shown in Table 2. They were interpreted to mean that both  $[\text{Melle}]^4\text{-CsA}$  and  $[\text{D-MeAla}]^3\text{-}[\text{EtVal}]^4\text{-CsA}$  are essentially devoid of immunosuppressive activity.

Table 2: Proliferative response of  $\text{CD4}^+$  PBMCs in the presence/absence of

10 cyclosporins

Co-culture	Compound	N	% Stimulation (relative)	Standard deviation
R1 x S2	None	8	100	30
R1 x S2	CsA	4	29	3
R1 x S2	$[\text{Melle}]^4\text{-CsA}$	4	84	13
R1 x S2	$[\text{D-MeAla}]^3\text{-}[\text{EtVal}]^4\text{-CsA}$	4	75	11
R2 x S1	None	8	100	9
R2 x S1	CsA	4	9	2
R2 x S1	$[\text{Melle}]^4\text{-CsA}$	4	75	9
R2 x S1	$[\text{D-MeAla}]^3\text{-}[\text{EtVal}]^4\text{-CsA}$	4	65	7
S1 x S2	None	4	0	0.6

R1 x S2 refers to a co-culture of responder cells from donor 1 and stimulator cells from donor 2. N is the number of measurements.

15 High anti-HCV activity and low cytotoxicity/cytostatic effect of  $[\text{D-MeAla}]^3\text{-}[\text{EtVal}]^4\text{-CsA}$

As mentioned previously, infection with hepatitis virus C (HCV) is a serious health problem because persistently infected patients are at a high risk for

20 developing chronic liver diseases including cirrhosis and hepatocellular carcinoma. Current available therapy is inadequate for a large fraction of the latter population as well as is associated with significant side effects. Until recently, development of

more effective therapies was hindered by the absence of an appropriate *in vitro* model of HCV replication that allows screening of potentially active compounds prior to evaluation in human clinical trials. This obstacle was overcome by the development of genetically modified HCV minigenomes (replicons) that self-  
5 amplify in cultured hepatoma cells to high levels (Lohmann et al. *Science* 285, (1999), 110-113). This HCV replicon system has rapidly become the standard tool for studying HCV replication, pathogenesis and persistence (Bartenschlager et al. *Antiviral Res.* 60, (2003), 91-102). The HCV genome consists of a single-stranded RNA that contains a single open reading frame for a polyprotein of about 3000  
10 amino acids. Translation of this polyprotein is initiated at an internal ribosome entry site (IRES) located at the 5' end of the RNA. The HCV polyprotein is cleaved into at least ten proteins. They include capsid protein C, envelope proteins E1 and E2, possible viroporin protein p7, non-structural proteins NS2 and NS3 having serine proteinase as well as ATPase/helicase activities, NS4A, membranous  
15 web-inducing protein NS4B, NS5A and RNA-dependent RNA polymerase NS5B. The first successful replicon was a bicistronic RNA containing in a 5' to 3' direction an HCV IRES, a coding sequence for a neomycin phosphotransferase, an IRES from an encephalocarditis virus and coding sequences for HCV proteins NS3 to NS5. Subsequent to introduction into Huh-7 cells and selection using G418  
20 (geneticin), this replicon could be shown to replicate autonomously to high levels (1,000-5,000 copies/cell) (Lohmann et al., 1999). Characterization of the system revealed that replication efficiency depended on permissiveness of the host cell and, importantly, on the selection of cell culture-adaptive mutations in the HCV protein-coding sequences. Replication was found to be sensitive to interferon  
25 alpha, providing evidence for the relevance of the system for screening drugs that have *in vivo* efficacy. Variant replicons were also constructed in which the neomycin phosphotransferase-coding sequence was replaced, e.g., by a luciferase-coding sequence or by sequences coding for a luciferase-ubiquitin-neomycin phosphotransferase fusion protein. Replication of the latter variant  
30 replicons can be assayed by the convenient luciferase assay, whereas replication of the former replicon requires determinations of RNA copy number.

Watashi et al. (2003) demonstrated by Northern blot and quantitative RT-PCR (reverse transcriptase polymerase chain reaction) that HCV RNA

accumulation was inhibited by CsA but not by the immunosuppressive macrolide FK506 and the non-immunosuppressive CsA derivative PSC 833 in HCV replicon-containing MH-14 cells. Their assays that involved 7-day exposures of cells to active agents revealed that HCV RNA titer was reduced by about 200 fold in the 5 presence of 1  $\mu$ g/ml cyclosporin A. They further found that non-immuno-suppressive cyclosporin [Melle]<sup>4</sup>-CsA also inhibited HCV replication. Results indicated that [Melle]<sup>4</sup>-CsA was about equally as effective as CsA in reducing HCV RNA titer.

10 To determine whether the cyclosporin of the present invention has anti-HCV activity and, should it has such activity, how this activity compares with the activities of CsA and [Melle]<sup>4</sup>-CsA, experiments were carried out that compared inhibitory effects of CsA, [Melle]<sup>4</sup>-CsA and [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA in HCV replicon systems.

15 Assays used Huh 5-2 cells that contained a bicistronic RNA encoding a firefly luciferase-ubiquitin-neomycin phosphotransferase fusion protein and HCV proteins NS3-5. The viral sequences originated from an HCV virus of genotype 1b. Cells were cultured in RPMI 1640 medium (Gibco) supplemented with 10% fetal 20 calf serum, 2 mM glutamine (Life Technologies), 1 x non-essential amino acids (Life Technologies), 100 u/ml penicillin, 100  $\mu$ g/ml streptomycin and 250  $\mu$ g/ml G418 (Geneticin, Life Technologies) at 37°C and 5% CO<sub>2</sub>. For antiviral (replication) assays, cells were seeded at a density of 7000 cells/well in 96-well View Plates<sup>TM</sup> (Packard) in the same medium except for G418. After a 24-h 25 incubation, medium was removed, serial dilutions of test compounds in medium were added, and cells were incubated for an additional 72 h.

Antiviral effects were estimated either by luciferase assay or quantitative RT-PCR. To carry out luciferase assays, medium was removed, and cells were 30 washed with PBS. Subsequent to lysis in 50  $\mu$ l of Glo-lysis buffer (Promega) for 15 min, 50  $\mu$ l of Stead-Glo Luciferase Assay Reagent (Promega) were added to cell lysates. Luciferase activity was measured using a luminometer, and the signal from each test well was expressed as a percentage of the signal measured in wells of cultures not exposed to a test compound.

Cell density and cytostatic effects were estimated in parallel cultures in regular 96-well plates (Beckton-Dickinson) using the MTT assay (CellTiter 96<sup>R</sup> AQ<sub>ueous</sub> Non-Radioactive Cell Proliferation Assay, Promega). In this assay, 5 3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxymethoxy-phenyl)-2-(4-sulfophenyl)-2H-tetrazolium (MTS) is bioreduced to a formazan that is quantitated at 498 nm in a plate reader. Formazan production is directly correlated with number of live cells.

RT-PCR analysis quantitated the neomycin region of replicons using an ABI 10 PRISM 7700 sequence detector (Applied Biosystems, Foster City, CA). The forward and reverse primers used were 5'-CCGGCTACCTGCCATT-3' and 5'-CCAGATCATCCTGATCGACAAG-3', respectively. The fluorogenic probe was 5'-ACATCGCATCGAGCGAGCACGTAC-3'. As an internal control, a plasmid containing part of the neomycin phosphotransferase gene sequence was used.

15 Results from these experiments permitted calculation of EC<sub>50</sub> for the different cyclosporins, which is the effective concentration required to inhibit HCV replicon replication by 50%, and of CC<sub>50</sub>, which is the concentration required that inhibits the proliferation of exponentially growing cells by 50%, and a selectivity 20 index SI, which is the ratio between CC<sub>50</sub> and EC<sub>50</sub>.

Table 3 shows values obtained from Huh 5-2 cells using luciferase activity assays for estimation of replication efficiency and MTT assays for calibration of luciferase assays and for estimation of cytostatic effects of compounds. In 25 agreement with the above-discussed observations by Watashi et al. (2003), CsA and [Melle]<sup>4</sup>-CsA had similar anti-HCV (replication) activities.

Surprisingly, [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA was considerably more potent than CsA and [Melle]<sup>4</sup>-CsA. It was also noted that the 50% cytostatic concentration 30 (CC50) for [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA was significantly higher than the values determined for CsA and [Melle]<sup>4</sup>-CsA. Consequently, a considerably higher selectivity index was found for [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA as compared to the two other cyclosporins. Analogous experiments in which EC<sub>50</sub> values were derived from determinations of RNA titers using quantitative RT-PCR yielded similar

conclusions. SI values of 45\*, 73 and 625\* were obtained for CsA, [Melle]<sup>4</sup>-CsA and [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA, respectively. Asterisks indicate that the lower of two independently determined values are presented.

5 Table 3: EC<sub>50</sub>, CC<sub>50</sub> and SI values determined from luciferase assays of HCV RNA replication and MTT assays of cytotoxicity in Huh 5-2 cells comprising a luciferase-containing HCV minireplicon

Compound	EC <sub>50</sub> (µg/ml) +/- Std.Dev.	CC <sub>50</sub> (µg/ml) +/- Std.Dev.	Selectivity index
CsA	0.28 +/- 0.13	11.6 +/- 5.6	41
[D-MeAla] <sup>3</sup> -[EtVal] <sup>4</sup> -CsA	0.03 +/- 0.04	> 27	> 900
[Melle] <sup>4</sup> -CsA	0.22	14	64

Antiviral activity of [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA measured in infected target cells with recombinant HCV

10 The anti-HCV activity of [D-MeAla]3-[EtVal]4-CsA compared to CsA was further determined in culture systems approaching the *in vivo* situation. The method used hepatoma cells that had been infected with an infectious full length 15 chimeric HCV construct or the same virus that was modified to carry a luciferase receptor gene. After the treatment of the infected cells with the cyclosporin of the invention or CsA, the luciferase activity was measured as being directly correlated to the inhibition of the viral replication.

20 Infectious HCV viruses of full-length chimeric genome between HCV strains J6 and JFH1 (Jc1) were used to inoculate the hepatoma cells of the assays. The construct of Jc1 virus was also modified to obtain a bicistronic genome carrying a luciferase reporter gene (Jc1-Luc). Twenty-four and ninety-six hours after the transfection of RNA transcripts of the genomes by electroporation of Huh-7.5 cells, 25 cell culture supernatant was collected. Supernatants were filtered (0.45 µM) and cell culture infectious dose 50 (CCID50) per ml were determined by the limiting dilution assays according to Lindenbach et al. (Science, 309, (2005), 623-626). The CCID50 were 1.3 x 10<sup>5</sup> for Jc1 and 4.2 x 10<sup>3</sup> for Jc1-Luc.

Assays used either Huh-7-Lunet or Huh-7.5 cells (Lohmann et al., *Science* 285(5424), (1999), 110-113). Cells were grown in Dulbecco's modified Eagle's Medium (DMEM; Gibco) supplemented with 10% heat-inactivated fetal bovine serum (FCS) (Integro), 1x non-essential amino acids (Gibco), 100 IU/ml penicillin 5 (Gibco), 100 µg/ml streptomycin (Gibco) or 25 µg/ml hygromycin (Gibco) for Huh-mono cells at 37°C and 5% CO<sub>2</sub>. For antiviral (replication) assays, Huh-7-Lunet and Huh-7.5 cells were seeded at a density of 2 x 10<sup>4</sup> or 4 x 10<sup>4</sup> cells per well of a 12-well plate. Twenty four hours later, the medium was replaced by 0.5 ml of the Jc1-Luc virus stock (12-well plates) or 0.25 ml of the Jc1 virus stock (12-well 10 plates). Four hours later, the virus inoculum was replaced by medium containing different concentrations of CsA or [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA and were further incubated for an additional 72 hours.

The inhibition of viral replication were estimated by luciferase assay. To 15 carry out luciferase assay, cells were harvested, washed with PBS and lysed in luciferase lysis buffer (1% Triton X-100, 25 mM glycylglycine, 15 mM MgSO<sub>4</sub>, 4mM EGTA, and 1 mM DTT). Firefly luciferase activity was measured according to Krieger et al. (*J Virol*, 75(10), (2001), 4614-4624). Briefly, after one freeze/thaw cycle, cells were resuspended and 100 µl of cell lysate was mixed with 360 µl 20 assay buffer (25 mM glycylglycine, 15 mM MgSO<sub>4</sub>, 1 mM DTT, 2 mM ATP, 15 mM potassium phosphate buffer, pH 7.8) and 200 µl substrate solution (200 mM luciferin, 25 mM glycylglycine). Finally, luminescence was measured by using a Lumat LB9507 luminometer (Berthold) for 20 samples.

25 In these examples (Figs. 1 and 2), both [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA (white bars) and CsA (black bars) resulted in a dose-dependent antiviral activity, whereby [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA proved again superior over CsA, thus corroborating the data obtained with the subgenomic replicons. A 10-fold higher concentration of CsA was needed to result in the same replication inhibiting effect as the 30 cyclosporin of the invention.

High affinity of the cyclosporin of the invention for cyclophilin

In the above-discussed observations by Watashi et al. (2003) and Nagakawa et al. (2003), the anti-HCV effect was related to the binding capacity of cyclosporins to the cyclophilins. The effects on the PPIase activity of CsA, [Melle]<sup>4</sup>-CsA and [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA was measured for cyclophilins to determine the more potent inhibitor of PPIase activity of cyclophilin, e.g. cyclophilin A, and, consequently, of HCV replication.

Commercial human recombinant cyclophilin A (Sigma) was used in the assays. PPIase activity of cyclophilins was determined using a chymotrypsin-coupled spectrophotometric assay according to Garcia-Echverria et al. (BBRC, 191, (1993), 70-75). This method is based on the high trans selectivity of chymotrypsin for peptides of the type N-succinyl-ala-ala-pro-phe-p-nitroanilide. The peptides cleavage liberated the para-nitroaniline moiety that could be detected and quantitated at 390 nm. The hydrolysis of the cis form was limited by the rate of cis-trans isomerisation carried out by cyclophilin A. The peptide was made up in a solution of 25 nM LiCl in 2,2,2-trifluoroethanol at 470 mM to enhance the cis conformer peptides population. The assay was performed on the split beam spectrophotometer and the water bath was set at 5°C. Cyclophilin A (7500 pmol/mg total enzyme concentration; Sigma) were dissolved at 20 nM in a buffer (35 mM HEPES and 0.26 mg/ml chymotrypsin (specific activity 50 units/mg), pH 7.8 with KOH) and were incubated for 6 minutes at room temperature followed by 54 minutes in the water bath. CsA, [Melle]<sup>4</sup>-CsA or [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA were added as appropriate in these incubations using a concentration range of 2-50 nM. Then 3.5 ml of the incubated cyclophilin was added to the sample cuvette. The reference cuvette contained a reaction that had gone to completion to balance the reference beam. Peptide was added at 25 µM to initiate the reaction and the change in absorbance was monitored at 10 data points per second. As a control, rates were also determined for a parallel reaction that lacked cyclophilin. These blank rates of peptide hydrolysis (i.e. in the absence of cyclophilin) were subtracted from rates in the presence of cyclophilin A.

The initial rates obtained from the PPIase assays were analysed by first order regression analysis by using first order transformation of the traces of the time course of the change in the absorbance at 390 nm. Total enzyme concentration ( $E_t$ ), the inhibitor dissociation constant ( $K_i$ ) and the rate constant for the rate limiting reaction was calculated with the software FigSyS (2003, Biosoft) by fitting the data obtained from the regression analysis in the tight binding inhibitor multiprotein equation.

The tight binding inhibitor multiprotein equation had the following formula:

10 
$$v = k * E_t * P - k * (-b - \sqrt{b * b - 4 * c}) / 2$$

where  $b$  is defined as  $b = -(E_t * P + I + K_i)$  and  $c$  is  $c = E_t * P * I$ .

Once  $E_t$ ,  $K_i$  and  $k$  were calculated by the computer for a given set of data, a graphic representation of the data was plotted and the line fitted to the points assuming tight inhibitor binding to a single protein, defined by the following 15 equations:

$$v = K * E_t * P - K(B - \sqrt{B * B - 4 * C}) / 2$$

where  $B = E_t * P + I + K_i$  and  $C = E_t * P * I$ .

Table 4:  $E_t$ ,  $K_i$  and  $k$  values of cyclophilin A for CsA,  $[Melle]^4$ -CsA and  $[D\text{-MeAla}]^3$ -  
20  $[EtVal]^4$ -CsA determined from PPIase activity assays.

Compound	$E_t$ (pmol/mg)	$K_i$ (nM)	$k$ (s-1)
CsA	7500	$9.79 \pm 1.37$	$0.17 \pm 0.0069$
$[Melle]^4$ -CsA	7500	$2.11 \pm 0.32$	$0.17 \pm 0.0068$
$[D\text{-MeAla}]^3$ - $[EtVal]^4$ -CsA	7500	$0.34 \pm 0.12$	$0.16 \pm 0.0074$

The lowest  $K_i$  of cyclophilin A observed for the cyclosporin of the invention corroborated the high potent of antiviral activity, the specificity and selectivity index (as above-mentioned) compared to CsA and  $[Melle]^4$ -CsA. Surprisingly, the non-25 immunosuppressive  $[D\text{-MeAla}]^3$ - $[EtVal]^4$ -CsA showed an almost 6-fold higher affinity for the cyclophilin of the example compared to the other non-immunosuppressive cyclosporin  $[Melle]^4$ -CsA.

The above-described experimentation provided that  
30  $[D\text{-MeAla}]^3$ - $[EtVal]^4$ -CsA was a more effective inhibitor of HCV replication than any

other tested cyclosporin. This increased anti-HCV activity correlated with the increased cyclophilin binding activity of [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA.

#### HCV replicon clearance and rebound

5

The recurrence of HCV infection is a major problem of the disease especially even with the use of potential efficient treatment, e.g. cyclosporin and/or Interferon. To study whether the more potent anti-HCV activity of the cyclosporin of the invention as compared to CsA is reflected in the ability of the compound to 10 more efficiently cure cells producing HCV replicon from those, an *in vitro* cell assay was performed based on presence of the selective drug G418 for recombinant produced replicon.

Assays used Huh-9-13 cells, human hepatoma cells (Huh-7) (Lohmann et 15 al., Science 285(5424), (1999), 110-113) Cells were grown in the usual complete medium DMEM without G418 pressure. The cells were cultured in the presence of either CsA or [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA (both at 0.5 or 1 µg/ml) or were left untreated for 7 consecutive passages. Control was performed to guarantee that the absence of the G418 selective pressure would not influence the HCV replicon 20 content during several passages. To confirm that Huh-9-13 cells that had been treated for 7 days with [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA were indeed cleared from their replicon, G418 selection (1000 µg/ml) was restarted for 2 more passages. Only those cells that were still carrying the HCV replicon have been able to proliferate under these conditions and cells without replicon have died in the presence of 25 G418 during the rebound phase.

RT-PCR were performed on extracts of viral RNA of samples taken at different passage points. The forward and reverse primers used were 5'-CCGGCTACCTGCCATT-3' and 5'-CCAGATCATCCTGATCGACAAG-3', 30 respectively. The fluorogenic probe was 5'-ACATCGCATCGAGCGAGCACGTAC-3'. As an internal control, a plasmid containing part of the neomycin phosphotransferase gene sequence was used. Results were analysed and

expressed as a quantity of replicon RNA (ng) per 1'000 cells and used to draft a graph.

Results from these experiments (Fig. 3) showed the superior antiviral effect  
5 of [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA compared to CsA in this standard *in vitro* cell assay. Surprisingly, the cyclosporin of the invention showed virucidal effect and not only virustatic effect as the other immunosuppressive CsA. Indeed, when the [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA treated Huh-9-13 cells (circles and square in Fig. 3) were again cultured in the presence of G418 (rebound phase), the cultures died  
10 compared to CsA treated cells (diamond and triangle). Both cultures that had been treated with CsA for 7 consecutive passages were able to proliferate in the presence of G418. This confirmed that [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA was able to cure Huh-9-13 cells from their HCV replicon.

15 Drug combination

Interferon (IFN) is part of the current therapy of HCV infection. The effect of [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA / IFN- $\alpha$  2a combination was evaluated using the method of Prichard and Shipman (Antiviral Res, 1990, 14, 181-205). In brief, the  
20 theoretical additive effect is calculated from the dose-response curves of individual compounds by the equation of formula:

$$Z = X + Y(1-X),$$

where X represents the inhibition produced by [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA alone and Y represents IFN- $\alpha$  2a alone. Z represents the effect produced by the combination  
25 of [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA with IFN- $\alpha$  2a. The theoretical additive surface is subtracted from the actual experimental surface, resulting in a horizontal surface that equals the zero plane when the combination is additive, a surface that lies above the zero plane indicates a synergistic effect of the combination and a surface below the zero plane indicates antagonism. The antiviral assay was  
30 carried out essentially as described above for Huh 5-2 cells except that compounds were added in checkerboard format. For each compound three replicate plates were used to measure the dose response curve of each individual compound. The data obtained from all three plates were used to calculate the

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theoretical additive surface. Combination studies for each pair of compounds were also done in triplicate. Data were analysed for variance by the ANOVA test.

A slight synergistic activity was noted at the highest concentrations of IFN- $\alpha$  2a

5 used, but overall the combined anti-HCV activity of [D-MeAla]3-[EtVal]4-CsA with IFN- $\alpha$  2a can be considered as additive (Fig. 4).

The findings with [D-MeAla]3-[EtVal]4-CsA can be summarized as follows:

- [D-MeAla]3-[EtVal]4-CsA has a more potent anti-HCV activity and is less cytotoxic than CsA, as shown in an HCV subgenomic replicon system.

10 - This has been confirmed in an hepatoma cell culture infected with a full-length infectious chimeric genome between HCV strains J6 and JFH1.

- [D-MeAla]3-[EtVal]4-CsA is able to cure cells from their HCV replicon more efficiently than CsA

15 - These effects are related to a more pronounced cyclophilin binding affinity.

- The anti-HCV activity of the combination [D-MeAla]3-[EtVal]4-CsA / IFN- $\alpha$  2a is additive.

[D-MeAla]3-[EtVal]4-CsA can be used to treat patients infected with HCV.

20 The active compound may be administered by any conventional route. It may be administered parentally, e.g., in the form of injectable solutions or suspensions, or in the form of injectable deposit formulations. Preferably, it will be administered orally in the form of solutions or suspensions for drinking, tablets or capsules.

Pharmaceutical compositions for oral administration comprising the cyclosporin of 25 the invention are described in Examples. As is demonstrated by the examples, such pharmaceutical compositions typically comprise the cyclosporin of the invention and one or more pharmaceutically acceptable carrier substances.

Typically, these compositions are concentrated and need to be combined with an appropriate diluent, e.g., water, prior to administration. Pharmaceutical

30 compositions for parenteral administration typically also include one or more excipients. Optional excipients include an isotonic agent, a buffer or other pH-controlling agent, and a preservative. These excipients may be added for maintenance of the composition and for the attainment of preferred ranges of pH (about 6.5-7.5) and osmolarity (about 300 mosm/L).

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Additional examples of cyclosporin formulations for oral administration can be found in U.S. Pat. Nos. 5,525,590 and 5,639,724, and U.S. Pat. Appl. 2003/0104992. By the oral route, the indicated dosage of the cyclosporin of the invention for daily to twice weekly administration may be from about 1 mg/kg to about 100 mg/kg, preferably from about 1 mg/kg to about 20 mg/kg. By the intravenous route, the indicated corresponding dosage may be from about 1 mg/kg to about 50 mg/kg, preferably from about 1 mg/kg to about 25 mg/kg. An effective amount of the cyclosporin of the invention is understood to be an amount that when administered repeatedly in the course of a therapeutic regimen to a patient in need of treatment of HCV infection results in an objective clinical response such as a statistically significant reduction in serum HCV titer or a significant reduction of serum ALT activity in the patient.

Initial phase I clinical studies were carried out to assess the safety of oral doses of [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA, and to determine the pharmacokinetic profile and safety profile of the drug substance. Studies showed that doses of 50 to 1600 mg in a micro-emulsion in water were well tolerated. Mild and short-lived side effects were observed including nausea, vomiting, abdominal pain, mild headaches. These side effects were not dose-related.

Numerous factors will be taken into consideration by a clinician when determining trial doses for testing efficacy of a pharmaceutical composition comprising the cyclosporin of the present invention against HCV infection. Primary among these are the toxicity and half-life of the chosen cyclosporin of the invention. Additional factors include the size of the patient, the age of the patient, the general condition of the patient (including significant systemic or major illnesses including decompensated liver disease, severe preexisting bone marrow compromise and other viral infections), the stage of HCV infection (acute vs. chronic) as indicated, e.g., by serum alanine aminotransferase (ALT) levels, the particular genotype of HCV, previous therapy of HCV infection, the presence of other drugs in the patient, and the like. A course of treatment will require repeated administration of a pharmaceutical composition of the invention. Typically, an adequate drug dose will be administered 3-7 times per week, and duration of

- 20 -

treatment may be from about 4 weeks to 6 months, preferably from about 4 weeks to about 12 months. Treatment may be followed by determinations of HCV in serum and measurement of serum ALT levels. The endpoint of treatment is a virological response, i.e., the absence of HCV at the end of a treatment course,

5 several months after initiation of treatment, or several months after completion of treatment. HCV in serum may be measured at the RNA level by methods such as quantitative RT-PCR or northern blots or at the protein level by enzyme immunoassay or enhanced chemiluminescence immunoassay of viral proteins. The endpoint may also include a determination of a serum ALT level in the normal 10 range.

A pharmaceutical composition of the present invention may comprise one or more other ingredients active against HCV infection in addition to the cyclosporin of the present invention such as, for example, another antiviral drug substance,

15 e.g., ribavirin, or an interferon alpha. The cyclosporin of the invention and such other active ingredient can be administered together as part of the same pharmaceutical composition or can be administered separately as part of an appropriate dose regimen designed to obtain the benefits of the combination therapy. The appropriate dose regimen, the amount of each dose administered, 20 and specific intervals between doses of each active agent will depend upon the specific combination of active agents employed, the condition of the patient being treated, and other factors discussed in the previous section. Such additional active ingredients will generally be administered in amounts less than or equal to those for which they are effective as single therapeutic agents. The FDA approved 25 dosages for such active agents that have received FDA approval for administration to humans are publicly available.

All patents, patent applications and publications cited herein shall be considered to have been incorporated by reference in their entirety.

30 The invention is further elaborated by the following examples. The examples are provided for purposes of illustration to a person skilled in the art, and are not intended to be limiting the scope of the invention as described in the claims. Thus, the invention should not be construed as being limited to the examples provided,

but should be construed to encompass any and all variations that become evident as a result of the teaching provided herein.

EXAMPLE 1: Synthesis of [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA

5 (Translated from a Ph.D. thesis by Jean Francois Guichou entitled "De nouveaux analogues de Cyclosporin A comme agent anti-VIH-1", Faculte des Sciences, University of Lausanne, CH-1015 Lausanne, Switzerland (2001)).

Synthesis of H-MeLeu-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-MeBmt(Oac)-

10 Abu-Sar-OMe:

4-Dimethylaminopyridine (DMAP) (41.5 mmoles; 5.8 g) was added to a solution of cyclosporin A (CsA) (8.3 mmoles; 10 g) in 100 ml acetic anhydride. The solution was stirred for 18 h at room temperature. The reaction mixture was then diluted with 600 ml ethyl acetate, and washed twice with water and four times with a saturated aqueous solution of sodium bicarbonate. The organic phase was dried over anhydrous  $\text{Na}_2\text{SO}_4$ , filtered and solvent was evaporated under reduced pressure. The yellow residue obtained was chromatographed on silica gel (eluent: 98:2 dichloromethane/methanol) and recrystallized in ether. 9.5 g of MeBmt(OAc)-CsA, a white powder, were recovered, representing a yield of 92%.

20

Trimethyloxonium tetrafluoroborate (22.5 mmoles; 3.3 g) was added to a solution of MeBmt(OAc)-Cs (7.5 mmoles; 9.4 g) in 60 ml dichloromethane. After 16 h at room temperature, 35 ml of 0.26 M sodium methanolate in methanol were added. After 1 h, 35 ml of methanol and 35 ml of 2 N sulphuric acid were added, and the reaction mixture was stirred for another 15 min, neutralized to pH 6.0 with saturated  $\text{KHCO}_3$  (28 ml) and extracted twice with ethyl acetate. The organic phase was washed 2 times with saturated  $\text{NaCl}$ , dried over anhydrous  $\text{Na}_2\text{SO}_4$  and filtered. Subsequently, solvent was evaporated under reduced pressure. The residue was chromatographed on silica gel (eluent: 5:1 ethylacetate/methanol).

25 7.3 g of H-MeLeu-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-MeBmt(OAc)-Abu-Sar-OMe were obtained (yield: 76%).

HPLC tr=268.23 mn (98%)

ES/MS: m/z: 1277.5 [M+H<sup>+</sup>], 639.2 [M+2H<sup>+</sup>]

Synthesis of H-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-MeBmt(OAc)-Abu-Sar-OMe:

DMAP (2.3 mmoles; 334 mg) and phenylisothiocyanate (6.9 mmoles; 0.75 ml) were added to a solution of H-MeLeu-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-  
5 MeBmt(OAc)-Abu-Sar-OMe (4.6 mmoles; 7 g) in 48 ml tetrahydrofuran. After 2 h, solvent was evaporated, and the crude product was chromatographed on silica gel (eluants: 9:1 tert-butyl methyl ether (MTBE)/ethylacetate (1); 9:1 MTBE/methanol (2)). 5.8 g of Ph-NH-C(S)- MeLeu-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-MeBmt(OAc)-Abu-Sar-OMe were obtained (90% yield).

10 13.8 ml trifluroracetic acid were added to a solution of the latter compound (4 mmoles; 5.6 g) in 290 ml dichloromethane. After 1 h of reaction, the mixture was neutralized using  $\text{KHCO}_3$  and diluted with 500 ml dichloromethane. The organic phase was washed 2 times with saturated  $\text{NaCl}$ , dried over anhydrous  $\text{Na}_2\text{SO}_4$  and filtered. Subsequently, solvent was evaporated under reduced pressure. The residue was chromatographed on silica gel (eluants: 9:1 MTBE/ethylacetate (1); 3:1 MTBE/methanol (2)). 2.8 g H-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-MeBmt(OAc)-Abu-Sar-OMe were obtained (61% yield). HPLC  $\text{tr}=25.80$  mn (99%)

15 20 ES/MS: m/z: 1050.5  $[\text{M}+\text{H}^+]$ , 547.7  $[\text{M}+2\text{H}^+]$

Synthesis of Boc-D-MeAla-EtVal-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-MeBmt(OAc)-Abu-NMe- $\text{CH}_2\text{-CH}_2\text{-OH}$ :

Fluoro-N,N,N',-tetramethylformamidinium hexafluorophosphate (TFFH) (0.96 mmoles; 0.25 g) was added, under an inert atmosphere, to a solution of H-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-MeBmt(OAc)-Abu-Sar-OMe (0.87 mmoles; 1.00 g), DIPEA (2.78 mmoles; 0.48 ml) and Boc-D-MeAla-EtVal-OH (0.96 mmoles; 0.32 g) in 15 ml dichloromethane. After 15 min, dichloromethane was evaporated, and the residue was taken up in ethylacetate. The organic phase was washed successively with a saturated  $\text{NaHCO}_3$  solution, a 10% solution of citric acid and a saturated  $\text{NaCl}$  solution, and was then dried over anhydrous  $\text{Na}_2\text{SO}_4$  and concentrated. Chromatography on silica gel (in 98:2 ethylacetate/methanol) yielded 1.14 g (90%) Boc-D-MeAla-EtVal-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-MeBmt(OAc)-Abu-Sar-OMe.

The latter product (0.64 mmoles; 0.93 g) was taken up in 45 ml anhydrous methanol, and sodium borohydride (25.5 mmoles; 0.96 g) was added in small portions at 15-min intervals over a period of 3 h 30 min. At 4 h, the reaction 5 mixture was cooled to 0°C, hydrolysed by addition of 10% citric acid and concentrated. Residue was taken up in ethylacetate. The organic phase was washed with a 10% solution of citric acid and a saturated NaCl solution, and was then dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and concentrated. After chromatography on silica gel (in 95:5 ethylacetate/methanol) 0.63 g (81%) of Boc-D-MeAla-EtVal-Val- 10 MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-MeBmt(OAc)-Abu-NMe-CH<sub>2</sub>-CH<sub>2</sub>-OH were obtained.

ES/MS: m/z: 1434.9 [M+H<sup>+</sup>], 717.9 [M+2H<sup>+</sup>]

Synthesis of H-D-MeAla-EtVal-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal- 15 MeBmt-Abu-OH:

Methanesulfonic acid (3.18 mmoles; 2.060 ml) was added to a solution of Boc-D-MeAla-EtVal-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-MeBmt(OAc)-Abu-NMe-CH<sub>2</sub>-CH<sub>2</sub>-OH (0.425 mmoles; 610 mg) in 42.5 ml methanol, and the mixture was heated to and maintained at 50°C. Progress of the reaction was 20 monitored by HPLC and mass spectrometry. After 80 h, the mixture was cooled to 0°C, and hydrolysed by addition of 1 M NaHCO<sub>3</sub>. Methanol was eliminated, and the residue was taken up in ethylacetate. The organic phase was washed with 1 M NaHCO<sub>3</sub> and then a saturated NaCl solution, dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and concentrated. The product (557 mg), H-D-MeAla-EtVal-Val-MeLeu-Ala-D-Ala- 25 MeLeu-MeLeu-MeVal-MeBmt(OAc)-Abu-O-CH<sub>2</sub>-CH<sub>2</sub>-NHMe, was used in the next step without purification.

Product (0.42 mmoles; 557 mg) was dissolved in 20 ml methanol and combined, under an inert atmosphere, with a solution of sodium methanolate (1.26 mmoles) 30 in 1.26 ml methanol. After 18 h at room temperature, the reaction mixture was cooled to 0°C, and sodium hydroxide (4.2 mmoles; 168 mg) in 5 ml water was added dropwise. After 21 h at room temperature, the reaction mixture was again cooled to 0°C and neutralized with 1 M KHSO<sub>4</sub>. Methanol was eliminated, and the residue was dissolved in ethylacetate. The organic phase was washed with a

semi-saturated NaCl solution, dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and concentrated. The product (335 mg; 64%), H-D-MeAla-EtVal-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-MeBmt-Abu-OH, was used in the next step without purification. HPLC tr=26.27 mn (86%)

5 ES/MS: m/z: 1235.5 [M+H<sup>+</sup>], 618.2 [M+2H<sup>+</sup>]

Synthesis of [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA:

Under an inert atmosphere, a solution of H-D-MeAla-EtVal-Val-MeLeu-Ala-D-Ala-MeLeu-MeLeu-MeVal-MeBmt-Abu-OH (0.162 mmoles; 200 mg) and sym.collidine 10 (1.78 mmoles; 0.24 ml) in 50 ml dichloromethane was added dropwise to a solution of (7-azabenzotriazole-1-yloxy)tritypyrrolidinophosphonium hexafluorophosphate (PyAOP, 0.486 mmoles; 254 mg) in 3.2 liter dichloromethane. 72 h later, the reaction mixture was hydrolysed by addition of a 10% Na<sub>2</sub>CO<sub>3</sub> solution. Dichloromethane was evaporated, and residue taken up in ethylacetate. 15 The organic phase was washed successively with a 0.1 N HCl solution and a saturated solution of NaCl, dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and concentrated. Crude product was purified on silica gel, yielding 110 mg (59%) [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA HPLC tr=30.54 mn (100%)  
ES/MS: m/z: 1217.6 [M+H<sup>+</sup>], 609.3 [M+2H<sup>+</sup>]

20

EXAMPLE 2: Oral formulations of cyclosporins of the invention.

Amounts are expressed as % w/w.

Example A:

25 Cyclosporin of the invention 10  
Glycofurol 75 35.95  
Miglycol 812 18  
Cremophor RH40 35.95  
Alpha-Tocopherol 0.1

30

Example B:

Cyclosporin of the invention 10  
Tetraglycol 2  
Captex 800 2

- 25 -

Nikkol HCO-40	85.9
Butylhydroxytoluene (BHT)	0.1

Example C:

5	Cyclosporin of the invention	10
	Glycofurool 75	39.95
	Miglycol 812	14
	Cremophor RH40	36
	Butylhydroxyanisole (BHA)	0.05-0.1

10

## Example D:

	Cyclosporin of the invention	10
	Tetraglycol	10
5	Myritol	5
	Cremophor RH40	74.9
	Alpha-Tocopherol	0.1

## Example E:

10	Cyclosporin of the invention	10
	Ethanol	9
	Propylene glycol	8
	Cremophor RH40	41
	Glycerol monolinoleate	32

15

For individual components of formulations A-D and for methods of preparation see British Patent Appl. No. 2,222,770.

## Claims

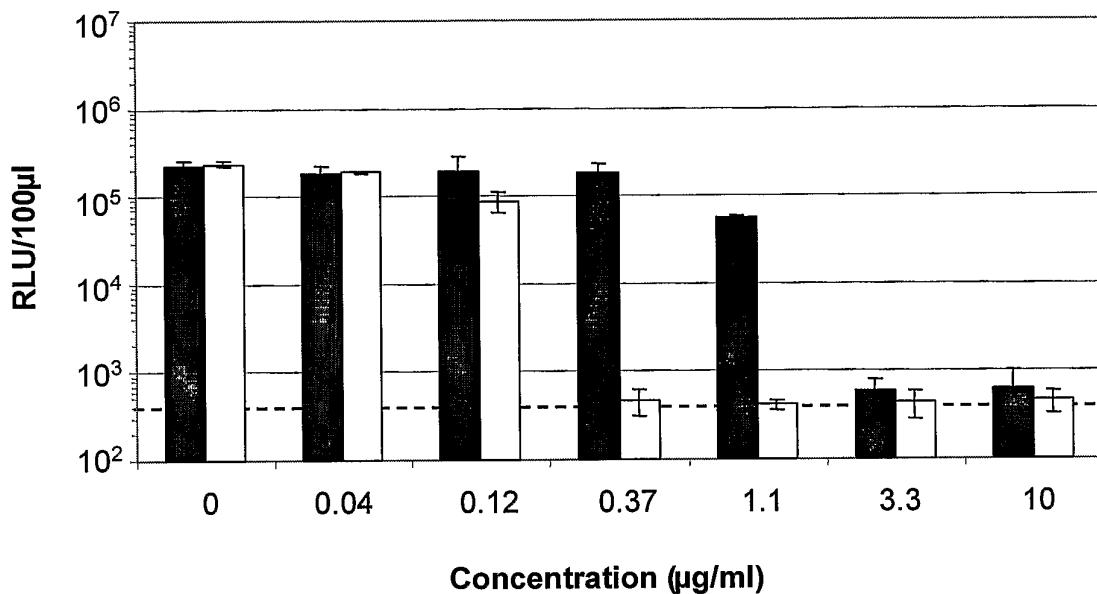
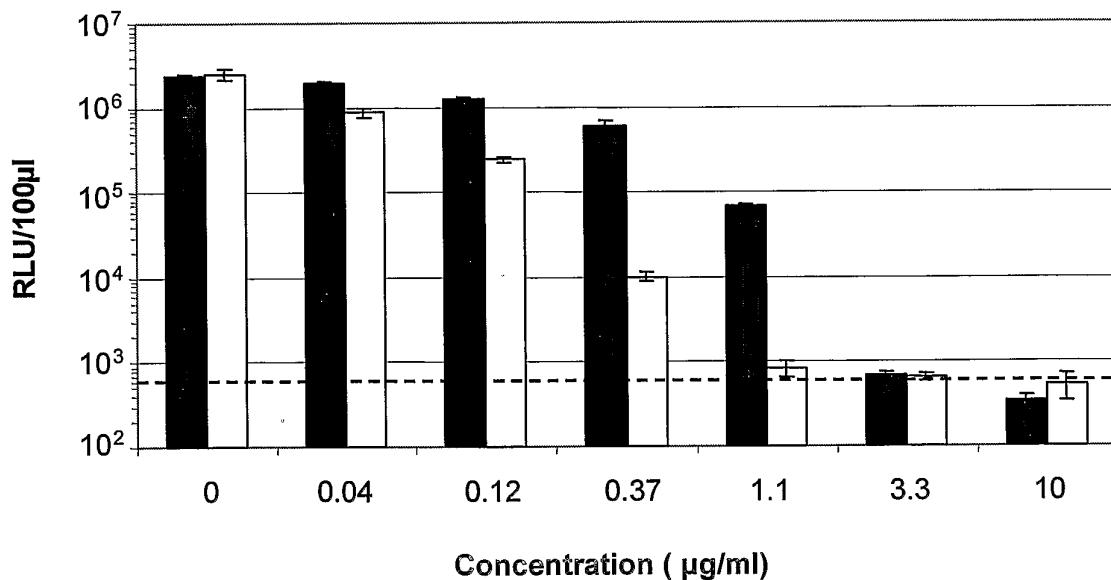
1. Use of [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA for the manufacture of a medicinal product intended for the treatment of HCV infection or the prevention of HCV recurrence in a patient.
2. Use according to claim 1 wherein said [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA is co-administered or administered separately with at least a second ingredient that is active against HCV infection as part of an appropriate dose regimen designed to obtain the benefits of the combination therapy of HCV infection in said patient.
3. A pharmaceutical composition comprising [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA as a first ingredient and at least an antiviral drug substance active against HCV as a second ingredient, wherein said second ingredient is selected in the group consisting of ribavirin and an interferon alpha.
4. The pharmaceutical composition according to claim 3 further comprising a pharmaceutically acceptable carrier and, optionally, a diluent.
5. A method for treatment of HCV infection or the prevention of HCV recurrence in a patient comprising administering to the patient an effective amount of [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA.
6. A method according to claim 5 comprising co-administering or administering separately as part of an appropriate dose regimen designed to obtain the benefits of the combination treatment effective amounts of said [D-MeAla]<sup>3</sup>-[EtVal]<sup>4</sup>-CsA and of a second ingredient that is active against HCV infection.
7. The use according to claim 1, substantially as herein described with reference to any one of the examples or figures.

**Dated 19 August, 2010**

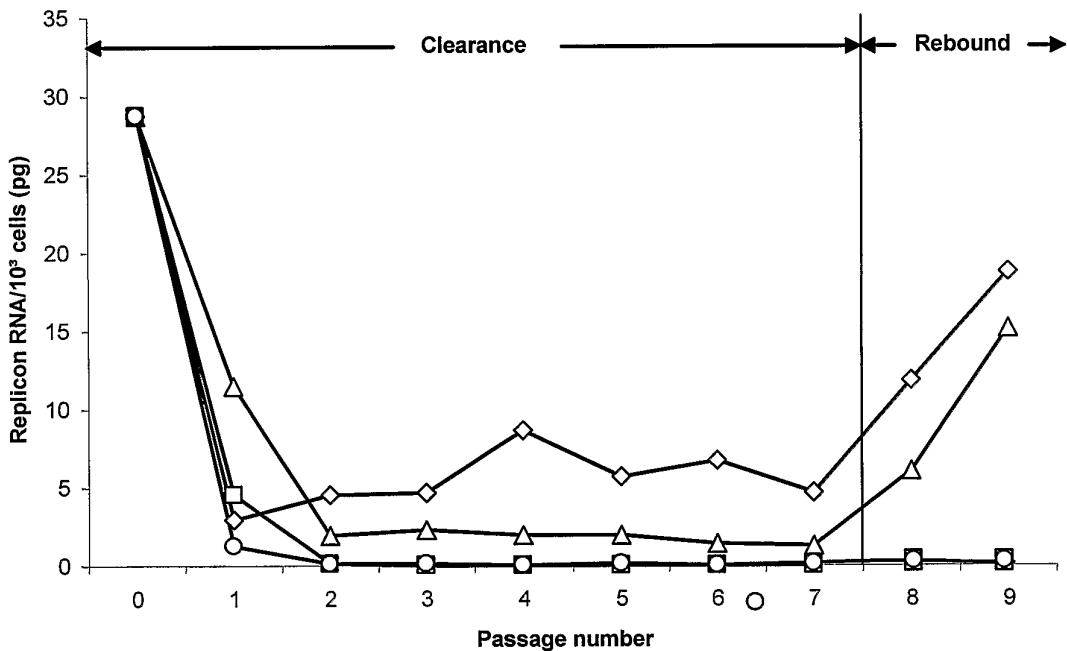
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**SPRUSON & FERGUSON**

**Fig. 1****Fig. 2**

**Fig. 3**



**Fig. 4**

