## **PCT**

## WORLD INTELLECTUAL PROPERTY ORGANIZATION International Bureau



International Patent Classification <sup>6</sup> :  A61K 38/31		(11) International Publication Number: WO 97/47317			
	A1	(43) International Publication Date: 18 December 1997 (18.12.97			
(21) International Application Number: PCT/EPS (22) International Filing Date: 11 June 1997 (1997) (30) Priority Data: 9612171.0 11 June 1996 (11.06.96) 9619310.7 16 September 1996 (16.09.96) (71) Applicant (for all designated States except US): NOVAG [CH/CH]; Schwarzwaldallee 215, CH-4058 Bas (72) Inventor; and (75) Inventor/Applicant (for US only): WECKBECKER, [DE/CH]; Loeliring 31, CH-4105 Biel-Benken (CH/CH) Agent: ROTH, Bernhard, M.; Novartis AG, Pate Markenabteilung, Klybeckstrasse 141, CH-4002 Bas	Gisbe	(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GH, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, MR, NE, SN, TD, TG).  Published  With international search report.			
(54) Title: COMBINATION OF A SOMATOSTATIN AN (57) Abstract  A combination of a compound of the somatostatin clapsperproliferation.		UE AND A RAPAMYCIN  a rapamycin macrolide is useful for the prevention or treatment of ce			



#### 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.

L	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
ΑZ	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	TM	Turkmenistan
BF	Burkina Faso	GR	Greece		Republic of Macedonia	TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	(E	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	zw	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's	NZ	New Zealand		
CM	Cameroon		Republic of Korea	PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Licchtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		



WO 97/47317 PCT/EP97/03036

#### COMBINATION OF A SOMATOSTATIN ANALOGUE AND A RAPAMYCIN

The present invention relates to a pharmaceutical combination and its use in the treatment of disorders associated with excess benign and malignant cell proliferation, e.g. tumors or intimal cell proliferation.

There is a continuing need for the development of drugs having increased effectiveness in inhibiting or slowing down undesired cell proliferation, particularly in the cancer field and in vasculopathies.

Accordingly, there is provided a pharmaceutical combination comprising a compound of the somatostatin class, and a rapamycin macrolide.

The somatostatin class is a known class of small peptides comprising the naturally occurring somatostatin-14 and analogues having somatostatin related activity, e.g. as disclosed by A.S. Dutta in Small Peptides, Vol.19, Elsevier (1993). By "somatostatin analogue" as used herein is meant any straight-chain or cyclic polypeptide having a structure based on that of the naturally occurring somatostatin-14 wherein one or more amino acid units have been omitted and/or replaced by one or more other amino radical(s) and/or wherein one or more functional groups have been replaced by one or more other functional groups and/or one or more groups have been replaced by one or several other isosteric groups. In general, the term covers all modified derivatives of the native somatostatin-14 which exhibit a somatostatin related activity, e.g. they bind to at least one somatostatin receptor (hSST-1, hSST-2, hSST-3, hSST-4 or hSST-5), preferably in the nMolar range, more preferably to at least the hSST-2 receptor in the nMolar range.

Cyclic, bridge cyclic and straight-chain somatostatin analogues or derivatives are known and have been described together with processes for their production e.g. in US Patent Specifications 4,310,518 and 4,235,886, in European Patent Specifications EP-A-1295; 23,192; 29,310; 29,579; 30,920; 31,303; 63,308; 70,021; 83,305; 215,171; 203,031; 214,872; 143,307; 298,732; 277,419; 389,180; 395,417; 450,480A2; in Belgian Patent Specification BE-A-900,089; and in WO 91/09056; WO 97/01579; WO 97/14715,



2

the contents thereof, in particular with respect to the compounds, being incorporated herein by reference.

Preferred somatostatin analogues are e. g. compounds of formula I

$$\begin{array}{c|cccc}
CH_2-S-Y_1 & Y_2-S-CH_2 \\
\hline
N-CH-CO-B-C-D-E-NH-CH-G & (I)
\end{array}$$

wherein

A is  $C_{1-12}$ alkyl,  $C_{7-10}$ phenylalkyl or a group of formula RCO-, whereby

- i) R is hydrogen, C<sub>1-11</sub>alkyl, phenyl or C<sub>7-10</sub>phenylalkyl, or
- ii) RCO- is
- a) a D-phenylalanine residue optionally ring-substituted by halogen, NO<sub>2</sub>, NH<sub>2</sub>, OH,
   C<sub>1-3</sub>alkyl and/or C<sub>1-3</sub>alkoxy; or
- the residue of a natural or a synthetic α-amino-acid other than defined under a)
   above, or of a corresponding D-amino acid, or
- c) a dipeptide residue in which the individual amino acid residues are the same or different and are selected from those defined under a) and/or b) above,
   the α-amino group of amino acid residues a) and b) and the N-terminal amino group of dipeptide residues c) being optionally mono- or di-C<sub>1-12</sub>alkylated or substituted by C<sub>1-8</sub>alkanoyl;

A' is hydrogen or  $C_{1-3}$ alkyl,

 $Y_1$  and  $Y_2$  represent together a direct bond or each of  $Y_1$  and  $Y_2$  is hydrogen

B is -Phe- optionally ring-substituted by halogen, NO<sub>2</sub>, NH<sub>2</sub>, OH, C<sub>1.3</sub>alkyl and /or



C<sub>1.3</sub>alkoxy (including pentafluoroalanine), naphthylalanine or pyridylalanine,

- C is (L)-Trp- or (D)-Trp- optionally α-N-methylated and optionally benzenering-substituted by halogen, NO<sub>2</sub>, NH<sub>2</sub>, OH, C<sub>1-3</sub>alkyl and/or C<sub>1-3</sub>alkoxy,
- D is Lys, 4-aminocyclohexylAla or 4-aminocyclohexylGly
- E is Thr, Ser, Val, Tyr, Ile, Leu or an aminobutyric or aminoisobutyric acid residue
- G is a group of formula

-COOR, -CH<sub>2</sub>OR<sub>10</sub>, -CON 
$$\stackrel{R_{11}}{\underset{R_{12}}{\swarrow}}$$
 or -CO-N  $\stackrel{R_{16}}{\underset{}{\swarrow}}$ 

wherein

R<sub>7</sub> is hydrogen or C<sub>1-3</sub>alkyl,

 $R_{10}$  is hydrogen or the residue of a physiologically acceptable, physiologically hydrolysable ester, e.g. formyl,  $C_{2-12}$ alkylcarbonyl, benzoyl,

R<sub>11</sub> is hydrogen, C<sub>1.3</sub>alkyl, phenyl or C<sub>7.10</sub>phenyl-alkyl,

R<sub>12</sub> is hydrogen, C<sub>1.3</sub>alkyl or a group of formula -CH(R<sub>13</sub>)-X<sub>1</sub>,

R<sub>13</sub> is CH<sub>2</sub>OH, -(CH<sub>2</sub>)<sub>2</sub>-OH, -(CH<sub>2</sub>)<sub>3</sub>-OH, -CH(CH<sub>3</sub>)OH, isobutyl, butyl, benzyl, naphthyl-methyl or indol-3-yl-methyl, and

X<sub>1</sub> is a group of formula

-COOR
$$_7$$
 -CH $_2$ OR $_{10}$  or -CO-N $<$ R $_{15}$ 

# DOCKET

## Explore Litigation Insights



Docket Alarm provides insights to develop a more informed litigation strategy and the peace of mind of knowing you're on top of things.

## **Real-Time Litigation Alerts**



Keep your litigation team up-to-date with **real-time** alerts and advanced team management tools built for the enterprise, all while greatly reducing PACER spend.

Our comprehensive service means we can handle Federal, State, and Administrative courts across the country.

## **Advanced Docket Research**



With over 230 million records, Docket Alarm's cloud-native docket research platform finds what other services can't. Coverage includes Federal, State, plus PTAB, TTAB, ITC and NLRB decisions, all in one place.

Identify arguments that have been successful in the past with full text, pinpoint searching. Link to case law cited within any court document via Fastcase.

## **Analytics At Your Fingertips**



Learn what happened the last time a particular judge, opposing counsel or company faced cases similar to yours.

Advanced out-of-the-box PTAB and TTAB analytics are always at your fingertips.

### API

Docket Alarm offers a powerful API (application programming interface) to developers that want to integrate case filings into their apps.

#### **LAW FIRMS**

Build custom dashboards for your attorneys and clients with live data direct from the court.

Automate many repetitive legal tasks like conflict checks, document management, and marketing.

#### **FINANCIAL INSTITUTIONS**

Litigation and bankruptcy checks for companies and debtors.

### **E-DISCOVERY AND LEGAL VENDORS**

Sync your system to PACER to automate legal marketing.

