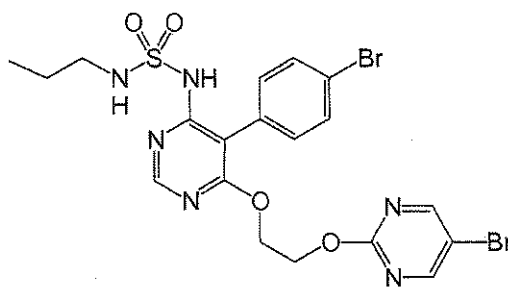


THERAPEUTIC COMPOSITIONS CONTAINING MACITENTAN

The present invention relates to a product containing macitentan, i.e. the compound of formula (I) below



(I)

or a pharmaceutically acceptable salt of this compound, in combination with at least one
5 compound having prostacyclin receptor (IP) agonist properties, or a pharmaceutically acceptable salt thereof, as well as this product for therapeutic use, simultaneously, separately or over a period of time, in the treatment of a disease wherein endothelin is involved.

PCT publication WO 02/053557 describes endothelin receptor antagonists including the
10 compound of formula (I) and the use of said endothelin receptor antagonists in the treatment of various diseases wherein endothelin is involved (i.a. heart failure, angina pectoris, pulmonary and systemic hypertension and erectile dysfunction).

Compounds having prostacyclin receptor (IP) agonist properties have been described notably in the following documents:

- 15 ❖ US 4,683,330 describe the compound treprostinil and salts and analogues thereof;
- ❖ US 4,539,333 describes epoprostenol sodium;
- ❖ US 4,692,464 describe the compound iloprost and salts and analogues thereof;
- ❖ US 4,474,802 describe the compound beraprost and salts and analogues thereof;
- ❖ US 7,205,302 describe among others 5,6-diphenylpyrazine derivatives having
20 prostacyclin receptor (IP) agonist properties, and salts and analogues thereof, an in

particular the compounds known under the code names MRE-269 and NS-304 (K. Kuwano et al., *J. Pharmacol. Exp. Ther.* (2007), **322**(3), 1181-1188).

Besides, WO 2004/017993 describes the use of the endothelin receptor antagonist bosentan together with the prostacyclin receptor agonist epoprostenol sodium for treating pulmonary arterial hypertension.

The Applicant has now found that the combination of macitentan with a compound having prostacyclin receptor (IP) agonist properties results in a strong synergistic effect in the treatment of diseases wherein endothelin is involved. Additionally, the possible side effects related to the compounds having prostacyclin receptor (IP) agonist properties (e.g. flushing or systemic hypotension) are expected to be decreased.

In one embodiment, the present disclosure relates thus to a product containing macitentan or a pharmaceutically acceptable salt thereof, and at least one (and preferably one) compound having prostacyclin receptor (IP) agonist properties, or a pharmaceutically acceptable salt thereof.

A further embodiment relates to a product containing macitentan or a pharmaceutically acceptable salt thereof, in combination with at least one (and preferably one) compound having prostacyclin receptor (IP) agonist properties, or a pharmaceutically acceptable salt thereof, for therapeutic use, simultaneously, separately or over a period of time, in the treatment of a disease wherein endothelin is involved.

In one aspect, there is provided a product containing macitentan or a pharmaceutically acceptable salt of this compound, in combination with at least one compound having prostacyclin receptor (IP) agonist properties, or a pharmaceutically acceptable salt thereof, wherein the compound having prostacyclin receptor (IP) agonist properties or pharmaceutically acceptable salt thereof is selected from iloprost and its pharmaceutically acceptable salts, beraprost and its pharmaceutically acceptable salts, 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide and its pharmaceutically acceptable salts, and {4-[(5,6-

diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}acetic acid and its pharmaceutically acceptable salts.

In another aspect, there is provided a product containing macitentan or a pharmaceutically acceptable salt of this compound, in combination with at least one compound having prostacyclin receptor (IP) agonist properties, or a pharmaceutically acceptable salt thereof, for therapeutic use, simultaneously, separately or over a period of time, in the treatment of a disease selected from hypertension, pulmonary hypertension, diabetic arteriopathy, heart failure, erectile dysfunction, angina pectoris and pulmonary fibrosis, wherein the compound having prostacyclin receptor (IP) agonist properties or pharmaceutically acceptable salt thereof is selected from iloprost and its pharmaceutically acceptable salts, beraprost and its pharmaceutically acceptable salts, 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide and its pharmaceutically acceptable salts, and {4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}acetic acid and its pharmaceutically acceptable salts.

In yet another aspect, there is provided a pharmaceutical composition containing, as active principles, macitentan, or a pharmaceutically acceptable salt thereof, in combination with at least one compound having prostacyclin receptor (IP) agonist properties, or a pharmaceutically acceptable salt thereof, as well as at least one pharmaceutically acceptable excipient, wherein the compound having prostacyclin receptor (IP) agonist properties or pharmaceutically acceptable salt thereof is selected from iloprost and its pharmaceutically acceptable salts, beraprost and its pharmaceutically acceptable salts, 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide and its pharmaceutically acceptable salts, and {4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}acetic acid and its pharmaceutically acceptable salts.

In still another aspect, there is provided the use of macitentan, or a pharmaceutically acceptable salt thereof, in combination with at least one compound having prostacyclin receptor (IP) agonist properties, or a pharmaceutically acceptable salt thereof, for the manufacture of a medicament intended to treat a disease selected from hypertension, pulmonary hypertension, diabetic arteriopathy, heart failure, erectile dysfunction, angina pectoris and pulmonary fibrosis, wherein

the compound having prostacyclin receptor (IP) agonist properties or pharmaceutically acceptable salt thereof is selected from iloprost and its pharmaceutically acceptable salts, beraprost and its pharmaceutically acceptable salts, 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide and its pharmaceutically acceptable salts, and {4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}acetic acid and its pharmaceutically acceptable salts.

The following paragraphs provide definitions of the various terms used in the present patent application and are intended to apply uniformly throughout the specification and claims, unless an otherwise expressly set out definition provides a broader or narrower definition.

Macitentan is the recommended INN for the compound of formula (I) and this name will therefore be used to designate the compound of formula (I) in the present patent application.

“Simultaneously” or “simultaneous”, when referring to a therapeutic use, means in the present application that the therapeutic use concerned consists in the administration of two or more active ingredients by the same route and at the same time.

“Separately” or “separate”, when referring to a therapeutic use, means in the present application that the therapeutic use concerned consists in the administration of two or more active ingredients at approximately the same time by at least two different routes.

By therapeutic administration “over a period of time” is meant in the present application the administration of two or more ingredients at different times, and in particular an administration method according to which the entire administration of one of the active ingredients is completed before the administration of the other or others begins. In this way it is possible to administer one of the active ingredients for several months before administering the other active ingredient or ingredients. In this case, no simultaneous administration occurs. Another therapeutic administration over a period of time consists in the administration over time of the two or more active ingredients of the combination using different frequencies of administration for each of the active ingredients, whereby at certain points in time simultaneous administrations of all the active ingredients of the combination take place whereas at some other points in time only part of the active ingredients of the combination may be administered (for example, in the case of a combination of macitentan with NS-304, the therapeutic administration over a period of time could be such that macitentan will be administered once a day whereas NS-304 will be administered twice a day).

By “disease wherein endothelin is involved” is meant in particular hypertension, pulmonary hypertension (including pulmonary arterial hypertension), diabetic arteriopathy, heart failure, erectile dysfunction, angina pectoris or pulmonary fibrosis.

By “compound having prostacyclin receptor (IP) agonist properties” is meant a compound that, when submitted to the “Test for the determination of prostacyclin receptor (IP) agonist EC₅₀” described in the present patent application, has an EC₅₀ value equal to or lower than 500 nM.

Specific examples of compounds having prostacyclin receptor (IP) agonist properties include treprostinil and its pharmaceutically acceptable salts, epoprostenol and its pharmaceutically acceptable salts, iloprost and its pharmaceutically acceptable salts, beraprost and its pharmaceutically acceptable salts, 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-N-(methylsulfonyl)acetamide (NS-304) and its pharmaceutically acceptable salts, and {4-[(5,6-diphenylpyrazin-

2-yl)(isopropyl)amino]butoxy}acetic acid (MRE-269) and its pharmaceutically acceptable salts.

The term "pharmaceutically acceptable salts" refers to non-toxic, inorganic or organic acid and/or base addition salts. Reference can be made to "Salt selection for basic drugs", *Int. J.*

5 *Pharm.* (1986), **33**, 201-217.

Besides, any reference to macitentan or to a compound having prostacyclin receptor (IP) agonist properties is to be understood as referring also to the pharmaceutically acceptable salts thereof, as appropriate and expedient.

10 Preferably, the product according to this invention will be such that macitentan and the compound having prostacyclin receptor (IP) agonist properties are intended for a therapeutic use which takes place simultaneously or over a period of time.

According to one preferred variant of this invention, macitentan and the compound having prostacyclin receptor (IP) agonist properties will be intended to be administered simultaneously.

15 According to another preferred variant of this invention, macitentan and the compound having prostacyclin receptor (IP) agonist properties will be intended to be administered over a period of time.

The period of time intended for the therapeutic use of a product according to this invention will be at least one week, and preferably at least one or more months (for example six
20 months). This period of time may also be the whole life of the patient that receives the product. According to a particular mode of administration according to this invention, administration of macitentan will be alternated with administration of a compound having prostacyclin receptor (IP) agonist properties, and the interval between such administration will not exceed two or three days (and more preferably not exceed one day). According to
25 another particular mode of administration according to this invention, in the case of a combination of macitentan with NS-304, the therapeutic administration over a period of time could be such that macitentan will be administered once a day whereas NS-304 will be administered twice a day.

Preferably, the compound having prostacyclin receptor (IP) agonist properties will be selected from 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide (NS-304) and its pharmaceutically acceptable salts, and {4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}acetic acid (MRE-269) and its
5 pharmaceutically acceptable salts. According to one particularly preferred variant of the invention the compound having prostacyclin receptor (IP) agonist properties will be 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide (NS-304) or a pharmaceutically acceptable salt thereof.

The administration route of macitentan and that of the compound having prostacyclin
10 receptor (IP) agonist properties is preferably the same. In particular, the common administration route for macitentan and for the compound having prostacyclin receptor (IP) agonist properties will be the oral route.

Though the exact administration doses of a product according to this invention will have to be determined by the treating physician, it is expected that a dose of 0.05 to 2 mg (and
15 preferably 0.1 to 1 mg) of macitentan per kg of patient body weight and per day will be appropriate. Similarly, it is expected that a dose of 0.5 to 30 µg (and preferably 1.5 to 15 µg) of compound having prostacyclin receptor (IP) agonist properties per kg of patient body weight given twice a day will be appropriate.

Preferably, the disease intended to be treated by a product according to this invention will
20 be selected from hypertension, pulmonary hypertension, diabetic arteriopathy, heart failure, erectile dysfunction, angina pectoris and pulmonary fibrosis. More preferably, the disease intended to be treated by a product according to this invention will be selected from hypertension and pulmonary hypertension. In particular, the disease intended to be treated by a product according to this invention will be pulmonary hypertension (and
25 notably pulmonary arterial hypertension).

The invention also relates to a pharmaceutical composition containing, as active principles, macitentan or a pharmaceutically acceptable salt of this compound, in combination with at least one (and preferably one) compound having prostacyclin receptor (IP) agonist properties, or a pharmaceutically acceptable salt thereof, as well as at least one excipient.

The invention further relates to the use of macitentan or a pharmaceutically acceptable salt of this compound, in combination with at least one (and preferably one) compound having prostacyclin receptor (IP) agonist properties, or a pharmaceutically acceptable salt thereof, for the manufacture of a medicament intended to treat a disease wherein endothelin is involved.

Besides, preferences indicated for the product according to this invention of course apply *mutatis mutandis* to the pharmaceutical compositions and uses of this invention.

Particular embodiments of the invention are described in the following Examples, which serve to illustrate the invention in more detail without limiting its scope in any way.

10 EXAMPLES

To illustrate the usefulness of this invention, the association of macitentan, administered orally at a dose of 10 mg/kg per day, with NS-304, administered orally at a dose of 1 mg/kg twice a day, can be studied in two different hypertension models, namely the pulmonary hypertension monocrotaline rat model and the spontaneously hypertensive rat model. Other associations may of course be tested similarly. The protocols that may be used are detailed in the part entitled "Pharmacological properties of the invention compounds" hereafter.

PHARMACOLOGICAL PROPERTIES OF THE INVENTION COMPOUNDS

Experimental methods:

20 The experimental methods described hereafter can be used to show the pharmacological properties of the invention compounds.

Monocrotaline model of pulmonary hypertension in rats

Male Wistar rats are purchased from Harlan (Netherlands) and maintained under conditions in accordance with local guidelines (Basel-Landschaft cantonal veterinary office). All rats are housed in climate-controlled conditions with a 12:12 hour light:dark

cycle, and had free access to chow and water. A telemetry system is implanted under anaesthesia by inhalation of 2.5% isoflurane (in 70% O₂ + 30% N₂O). Under aseptic conditions, a pressure radio-frequency transmitter is implanted into the peritoneal cavity, and a sensing catheter is inserted in the pulmonary artery. The transmitter is sutured to the abdominal musculature and the skin is closed. A receiver platform transforms the radio signal into digitized input, that is sent to a dedicated personal computer (Compaq, Deskpro). Pulmonary arterial blood pressure measurements are calibrated by using an input from an ambient pressure reference. Telemetry units are obtained from Data Sciences (St. Paul, MN, USA). Monocrotaline (MCT; Sigma Chemicals, St Louis, MO, USA) is administered as a single subcutaneous (sc) injection (60 mg/kg) in a volume of 3 ml/kg, and control age-matched rats receive an equal volume of saline.

Variant 1: chronic effect assessment:

The animals are randomly assigned to experimental groups, and treatment is initiated within 24 h after MCT injection, for a duration of 4 weeks. Macitentan and the compound having prostacyclin receptor (IP) agonist properties are administered by the oral route. The effects of macitentan, the compound having prostacyclin receptor (IP) agonist properties and their combination on pulmonary arterial blood pressure are measured by collecting data at 5-minute intervals. Hourly means of pulmonary arterial pressure are calculated for each rat. At the end of recording, rats are sacrificed. The heart is removed and weighed, and the ratio of organ weight to body weight (BW) is calculated. The right ventricle (RV) and the left ventricle plus septum are separated and weighed; the ratio RV/BW is used as an index of right ventricular hypertrophy. The lower the ratio RV/BW, the stronger the effect of the item(s) tested for reducing right ventricular hypertrophy.

Variant 2: acute effect assessment:

Four weeks after injection of MCT, the rats become pulmonary hypertensive and the respective effects of a single oral administration of Macitentan, of the compound having prostacyclin receptor (IP) agonist properties and of their combination can be evaluated on mean pulmonary arterial pressure.

Spontaneously hypertensive rat model

The same protocol is used as for the monocrotaline model of pulmonary hypertension in rats, except that spontaneously hypertensive rats (SHR) replace the monocrotaline-treated rats. The SHR rats are purchased from Harlan (Netherlands).

5 Test for the determination of prostacyclin receptor (IP) agonist EC₅₀:

CHO cells stably expressing the human IP receptor are cultured in Ham's F-12 medium containing 10% fetal bovine serum in a humidified atmosphere of 95% air and 5% CO₂ at 37°C. Cells are seeded at 1 x 10⁵ cells/well in 24-well plates and cultured for 48 h. Following a wash and incubation with assay buffer for 1 h at 37°C, the cells are exposed to
10 various concentrations of test compound in the presence of IBMX (500 µM). After removal of supernatant, the reaction is stopped by addition of 0.2 M perchloric acid. Adherent cells are frozen for 2 h at -80°C and thawed to extract intracellular cAMP. Supernatants are collected in tubes, neutralized by 2M KHCO₃ solution, and then centrifuged at 14,000 g for 10 min at 4°C to obtain samples for measurement of cAMP
15 levels by EIA system. Protein content of cell debris adhered to culture plates is measured following solubilization in 1N NaOH solution. Levels of cAMP are expressed as pmol cAMP/ mg protein. The EC₅₀ value is determined from non-linear regression analyses of concentration-response curves, and is defined as the negative logarithm of the concentration of test compound that elicits a response that is half of the maximal effect
20 observed.

*Experimental results:***EXAMPLE 1: acute effect of macitentan, NS-304 and their combination on mean pulmonary arterial pressure in monocrotaline treated rats:**

Experiments were performed in pulmonary hypertensive male Wistar rats treated with
25 monocrotaline according to the monocrotaline model as described in the section "Monocrotaline model of pulmonary hypertension in rats" of the part entitled "Experimental methods".

25 to 30 days after the monocrotaline treatment, four groups of 6 rats were formed and studied:

- the first group was treated neither with macitentan nor with NS-304 (control group);
- 5 - the second group was treated with macitentan only (10 mg/kg per os);
- the third group was treated neither with NS-304 only (30 mg/kg per os);
- the fourth group was treated with a combination of macitentan (10 mg/kg per os) and NS-304 (30 mg/kg per os);.

10 Mean pulmonary arterial pressure was measured over time. The relationship between mean pulmonary arterial pressure and time was plotted as a graph. Area Under Curve (AUC) was calculated for each group of rats after the oral administration of the different treatment(s) (a positive area being found if the mean pulmonary arterial blood pressure increases and a negative area being found if the mean pulmonary arterial blood pressure decreases). The results thus obtained are summarized in Table 1 hereafter.

Rat group [treatment]	AUC
Group 1 [control: no treatment]	106 ± 134
Group 2 [treatment with macitentan (10 mg/kg per os)]	-381 ± 147
Group 3 [treatment with NS-304 (30 mg/kg per os)]	-99 ± 128
Group 4 [treatment with macitentan (10 mg/kg per os) and NS-304 (30 mg/kg per os)]	-758 ± 164

Table 1

- 15 These data obtained in the monocrotaline model of pulmonary arterial hypertension confirm the synergistic effect of a combination of macitentan and NS-304 in the treatment of rats having previously undergone monocrotaline administration.

Claims

1. A product containing macitentan or a pharmaceutically acceptable salt of this compound, in combination with at least one compound having prostacyclin receptor (IP) agonist properties, or a pharmaceutically acceptable salt thereof, wherein the compound having prostacyclin receptor (IP) agonist properties or pharmaceutically acceptable salt thereof is selected from iloprost and its pharmaceutically acceptable salts, beraprost and its pharmaceutically acceptable salts, 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide and its pharmaceutically acceptable salts, and {4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}acetic acid and its pharmaceutically acceptable salts.
2. A product according to claim 1, wherein the compound having prostacyclin receptor (IP) agonist properties or pharmaceutically acceptable salt thereof is selected from 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide and its pharmaceutically acceptable salts, and {4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}acetic acid and its pharmaceutically acceptable salts.
3. A product according to claim 2, wherein the compound having prostacyclin receptor (IP) agonist properties or pharmaceutically acceptable salt thereof is 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide or a pharmaceutically acceptable salt thereof.
4. A product containing macitentan or a pharmaceutically acceptable salt of this compound, in combination with at least one compound having prostacyclin receptor (IP) agonist properties, or a pharmaceutically acceptable salt thereof, for therapeutic use, simultaneously, separately or over a period of time, in the treatment of a disease selected from hypertension, pulmonary hypertension, diabetic arteriopathy, heart failure, erectile dysfunction, angina pectoris and pulmonary fibrosis, wherein the compound having prostacyclin receptor (IP) agonist properties or pharmaceutically acceptable salt thereof is selected from iloprost and its pharmaceutically acceptable salts, beraprost and its pharmaceutically acceptable salts, 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide and its pharmaceutically acceptable salts, and {4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}acetic acid and its pharmaceutically acceptable salts.

5. A product according to claim 4, wherein the compound having prostacyclin receptor (IP) agonist properties is selected from 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide and its pharmaceutically acceptable salts, and {4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}acetic acid and its pharmaceutically acceptable salts.
6. A product according to claim 5, wherein the compound having prostacyclin receptor (IP) agonist properties is 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide or a pharmaceutically acceptable salt thereof.
7. A product according to one of claims 4 to 6, wherein the disease wherein endothelin is involved is pulmonary hypertension.
8. A pharmaceutical composition containing, as active principles, macitentan, or a pharmaceutically acceptable salt thereof, in combination with at least one compound having prostacyclin receptor (IP) agonist properties, or a pharmaceutically acceptable salt thereof, as well as at least one pharmaceutically acceptable excipient, wherein the compound having prostacyclin receptor (IP) agonist properties or pharmaceutically acceptable salt thereof is selected from iloprost and its pharmaceutically acceptable salts, beraprost and its pharmaceutically acceptable salts, 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide and its pharmaceutically acceptable salts, and {4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}acetic acid and its pharmaceutically acceptable salts.
9. A pharmaceutical composition according to claim 8, wherein the compound having prostacyclin receptor (IP) agonist properties or pharmaceutically acceptable salt thereof is

2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide or a pharmaceutically acceptable salt thereof.

10. Use of macitentan, or a pharmaceutically acceptable salt thereof, in combination with at least one compound having prostacyclin receptor (IP) agonist properties, or a
5 pharmaceutically acceptable salt thereof, for the manufacture of a medicament intended to treat a disease selected from hypertension, pulmonary hypertension, diabetic arteriopathy, heart failure, erectile dysfunction, angina pectoris and pulmonary fibrosis, wherein the compound having prostacyclin receptor (IP) agonist properties or pharmaceutically acceptable salt thereof is selected from iloprost and its pharmaceutically acceptable salts,
10 beraprost and its pharmaceutically acceptable salts, 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide and its pharmaceutically acceptable salts, and {4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}acetic acid and its pharmaceutically acceptable salts.

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11. Use according to claim 10, wherein the compound having prostacyclin receptor (IP) agonist properties or pharmaceutically acceptable salt thereof is 2-{4-[(5,6-diphenylpyrazin-2-yl)(isopropyl)amino]butoxy}-*N*-(methylsulfonyl)acetamide or a pharmaceutically acceptable salt thereof.

5 12. Use according to claim 10 or 11, wherein the disease intended to be treated is pulmonary hypertension.