VENTAVIS® (iloprost) Inhalation Solution

HIGHLIGHTS OF PRESCRIBING INFORMATION
These highlights do not include all the information needed to use VENTAVIS safely and effectively. See full prescribing information for VENTAVIS.

**INDICATIONS AND USAGE**
VENTAVIS® is a synthetic analog of prostacyclin indicated for the treatment of pulmonary arterial hypertension (PAH) (WHO Group 1) to improve a composite endpoint consisting of exercise tolerance, symptoms (NYHA Class), and lack of deterioration. Studies establishing effectiveness included predominately patients with NYHA Functional Class III-IV symptoms and etiologies of idiopathic or heritable PAH (65%) or PAH associated with connective tissue diseases (23%)(1.1).

**DOSAGE AND ADMINISTRATION**
VENTAVIS is intended to be inhaled using the I-neb® AAD® System. Patients should receive 6 to 9 doses (inhalations) per day (minimum of 2 hours between doses during waking hours) as follows:
- Starting dose: 2.5 mcg (2.1).
- Uptitrate to 5 mcg if 2.5 mcg is well tolerated (2.1).
- Maintenance dose: 5 mcg (2.1).

<table>
<thead>
<tr>
<th>Nebulizer</th>
<th>Delivered dose from ampule of:</th>
</tr>
</thead>
<tbody>
<tr>
<td>10 mcg/mL</td>
<td>2.5 or 5 mcg from one ampule</td>
</tr>
<tr>
<td>20 mcg/mL</td>
<td>5 mcg from one ampule</td>
</tr>
</tbody>
</table>

The 20 mcg/mL concentration is for patients who repeatedly experience extended treatment times (2.1).

Vital signs should be monitored while initiating VENTAVIS (2.2).

**DOSE FORMS AND STRENGTHS**
VENTAVIS® is supplied in 1 mL ampules in two concentrations: 10 mcg/mL and 20 mcg/mL.

<table>
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<td>5 mcg from one ampule</td>
</tr>
</tbody>
</table>

VENTAVIS® is intended to be inhaled using the I-neb® AAD® System. The first inhaled dose should be delivered dose from ampule of:

**CONTRAINDICATIONS**
None (4)

**WARNINGS AND PRECAUTIONS**
- Hypotension leading to syncope has been observed. VENTAVIS should not be administered in patients with systolic blood pressure below 85 mmHg (5.1).
- Pulmonary venous hypertension: Discontinue if pulmonary edema is present (5.2).
- May cause bronchospasm: Patients with a history of hyperreactive airway disease may be more sensitive (5.3).

**ADVERSE REACTIONS**
Most common (≥3% placebo adjusted) adverse reactions are vasodilation (flushing), cough increased, headache, trismus, insomnia, nausea, hypotension, vomiting, alkaline phosphatase increased, flu syndrome, back pain, tongue pain, palpitations, syncope, GGT increased, muscle cramps, hemoptysis, and pneumonia (6.1).

To report SUSPECTED ADVERSE REACTIONS, contact Actelion at 1-866-228-3546 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

**DRUG INTERACTIONS**
- VENTAVIS has the potential to increase the hypotensive effect of vasodilators and antihypertensive agents (7.2).
- There is a potential for increased risk of bleeding, particularly in patients maintained on anticoagulants (7.3).

**USE IN SPECIFIC POPULATIONS**
- Hepatic impairment: In patients with Child-Pugh Class B or C hepatic impairment, consider increasing the dosing interval (e.g., 3-4 hours between doses depending on the patient’s response at the end of the dose interval) (2.3, 8.6).

See 17 for PATIENT COUNSELING INFORMATION and preparation instructions

Revised: 11/2013

VENTAVIS® (iloprost) Inhalation Solution

VENTAVIS® is supplied in 1 mL ampules in two concentrations: 10 mcg/mL and 20 mcg/mL.

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<td>2.5 or 5 mcg from one ampule</td>
</tr>
<tr>
<td>20 mcg/mL</td>
<td>5 mcg from one ampule</td>
</tr>
</tbody>
</table>

The 20 mcg/mL concentration is intended for patients who are maintained at the 5 mcg dose and who have repeatedly experienced extended treatment times which could result in incomplete dosing. Transitioning patients to the 20 mcg/mL concentration using the I-neb® AAD® System will decrease treatment times to help maintain patient compliance.

For each inhalation session, the entire contents of each opened ampule of VENTAVIS should be transferred into the I-neb® AAD® System medication chamber immediately before use [see Patient Counseling Information (17.1)]. After each inhalation session, any solution remaining in the medication chamber should be discarded. Use of the remaining solution will result in unpredictable dosing. Patients should follow the manufacturer’s instructions for cleaning the I-neb® AAD® System components after each dose administration.

**2.2 Monitoring**
Vital signs should be monitored while initiating VENTAVIS. [see Warnings and Precautions (5.1)].
VENTAVIS® (iloprost) Inhalation Solution

2.3 Use in Patients with Pre-existing Hepatic Impairment
Because iloprost elimination is reduced in patients with impaired liver function [see Special Populations (8.6)], consider increasing the dosing interval (e.g., 3-4 hours between doses depending on the patient’s response at the end of the dose interval) in patients with Child-Pugh Class B or C hepatic impairment.

2.4 Use in Patients with Pre-existing Renal Impairment
Dose adjustment is not required in patients who are not on dialysis. The effect of dialysis on iloprost is unknown [see Special Populations (8.7)].

3. DOSAGE FORMS AND STRENGTHS
1 mL ampules in two concentrations: 10 mcg/mL and 20 mcg/mL.

4. CONTRAINDICATIONS
None.

5. WARNINGS AND PRECAUTIONS
VENTAVIS solution should not be allowed to come into contact with the skin or eyes; oral ingestion of VENTAVIS solution should be avoided.

5.1 Risk of Syncope
Monitor vital signs while initiating VENTAVIS. Do not initiate VENTAVIS in patients with systolic blood pressure below 85 mmHg or diastolic blood pressure below 60 mmHg. Syncope can also occur in association with pulmonary arterial hypertension, particularly in association with physical exertion. The occurrence of exertional syncope may reflect a therapeutic gap or insufficient efficacy, and the need to adjust dose or change therapy should be considered.

5.2 Pulmonary Venous Hypertension
Should signs of pulmonary edema occur when inhaled VENTAVIS is administered in patients with pulmonary hypertension, stop treatment immediately, as this may be a sign of pulmonary venous hypertension.

5.3 Bronchospasm
VENTAVIS inhalation can induce bronchospasm. Bronchospasm may be more severe or frequent in patients with a history of hyperreactive airways. VENTAVIS has not been evaluated in patients with chronic obstructive pulmonary disease (COPD), severe asthma, or with acute pulmonary infections.

6. ADVERSE REACTIONS

6.1 Clinical Studies Experience
Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice.

Pre-marketing safety data on VENTAVIS were obtained from 215 patients with pulmonary arterial hypertension receiving iloprost in two 12-week clinical trials and two long-term extensions. Patients received inhaled VENTAVIS for periods of from 1 day to more than 3 years. The median number of weeks of exposure was 15. Forty patients completed 12 months of open-label treatment with iloprost.

The following table shows adverse events reported by at least 4 VENTAVIS patients and reported at least 3% more frequently for VENTAVIS patients than placebo patients in the 12-week placebo-controlled study.

Table 1: Adverse Events in Phase 3 Clinical Trial

<table>
<thead>
<tr>
<th>Adverse Event</th>
<th>VENTAVIS n = 101</th>
<th>Placebo n = 102</th>
<th>Placebo subtracted %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vasodilation</td>
<td>27</td>
<td>9</td>
<td>18</td>
</tr>
<tr>
<td>Cough increased</td>
<td>39</td>
<td>26</td>
<td>13</td>
</tr>
<tr>
<td>Headache</td>
<td>30</td>
<td>20</td>
<td>10</td>
</tr>
<tr>
<td>Trismus</td>
<td>12</td>
<td>3</td>
<td>9</td>
</tr>
<tr>
<td>Insomnia</td>
<td>8</td>
<td>2</td>
<td>6</td>
</tr>
<tr>
<td>Nausea</td>
<td>13</td>
<td>8</td>
<td>5</td>
</tr>
<tr>
<td>Hypotension</td>
<td>11</td>
<td>6</td>
<td>5</td>
</tr>
<tr>
<td>Vomiting</td>
<td>7</td>
<td>2</td>
<td>5</td>
</tr>
<tr>
<td>Alk phos increased</td>
<td>6</td>
<td>1</td>
<td>5</td>
</tr>
<tr>
<td>Flu syndrome</td>
<td>14</td>
<td>10</td>
<td>4</td>
</tr>
<tr>
<td>Back pain</td>
<td>7</td>
<td>3</td>
<td>4</td>
</tr>
<tr>
<td>Tongue pain</td>
<td>4</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>Palpitations</td>
<td>7</td>
<td>4</td>
<td>3</td>
</tr>
<tr>
<td>Syncope</td>
<td>8</td>
<td>5</td>
<td>3</td>
</tr>
<tr>
<td>GGT increased</td>
<td>6</td>
<td>3</td>
<td>3</td>
</tr>
<tr>
<td>Muscle cramps</td>
<td>8</td>
<td>3</td>
<td>3</td>
</tr>
<tr>
<td>Hemosphatysis</td>
<td>5</td>
<td>2</td>
<td>3</td>
</tr>
<tr>
<td>Pneumonia</td>
<td>4</td>
<td>1</td>
<td>3</td>
</tr>
</tbody>
</table>

Pre-marketing serious adverse events reported with the use of inhaled VENTAVIS and not shown in Table 1 include congestive heart failure, chest pain, supraventricular tachycardia, dyspnea, peripheral edema, and kidney failure.

In a small clinical trial (the STEP trial) [see Clinical Studies (14)], safety trends in patients receiving concomitant bosentan and VENTAVIS were consistent with those observed in the larger experience of the Phase 3 study in patients receiving only VENTAVIS or bosentan.

Adverse events with higher doses
In a study in healthy subjects (n=180), inhaled doses of iloprost solution were given every 2 hours, beginning with 5 mcg and increasing up to 20 mcg for a total of 6 dose inhalations (total cumulative dose of 70 mcg) or up to the highest dose tolerated in a subgroup of 40 subjects. There were 13 subjects (32%) who failed to reach the highest scheduled dose (20 mcg). Five were unable to increase the dose because of [mild to moderate] transient chest pain/discomfort/tightness, usually accompanied by headache, nausea, and dizziness. The remaining 8 subjects discontinued for other reasons.

6.2 Postmarketing Experience
The following adverse reactions have been identified during the postapproval use of VENTAVIS. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

Cases of bronchospasm and wheezing have been reported, particularly in patients with a history of hyperreactive airways [see Warnings and Precautions (5.2)]. Bleeding events most commonly reported were epistaxis and hemoptysis observed in VENTAVIS treatment [see Adverse Reactions (7.2)]. Cases of thrombocytopenia, dizziness, diarrhea, mouth and tongue irritation, nasal congestion, dysgeusia, hypersensitivity, and rash have also been reported with the use of VENTAVIS.

7. DRUG INTERACTIONS
During clinical trials, iloprost was used concurrently with anticoagulants, diuretics, cardiac glycosides, calcium channel blockers, analgesics, antipyretics, nonsteroidal anti-inflammatory drugs, corticosteroids, and other medications. Intravenous infusion of iloprost had no effect on the pharmacokinetics of digoxin. Acetylsalicylic acid did not alter the clearance (pharmacokinetics) of iloprost.

7.1 Cytochrome P450
Although clinical studies have not been conducted with VENTAVIS (inhaled iloprost), in vitro studies indicate that no relevant inhibition of cytochrome P450 450 drug metabolism would be expected.

7.2 Antithrombotic and Vasodilator
In studies in normal subjects, there was no pharmacodynamic interaction between intravenous iloprost and either nifedipine, diltiazem, or captopril. However, VENTAVIS has the potential to increase the hypotensive effect of vasodilators and antithrombotic agents.

7.3 Anticoagulants and Platelet Inhibitors
Since VENTAVIS inhibits platelet function, there is a potential for increased risk of bleeding, particularly in patients maintained on anticoagulants or platelet inhibitors.

8. USE IN SPECIFIC POPULATIONS

8.1 Pregnancy
Pregnancy Category C:
VENTAVIS has been shown to be teratogenic in rats as described below. There are no adequate and well controlled studies in pregnant women. VENTAVIS should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

In developmental toxicity studies in pregnant Han-Wistar rats, continuous intravenous administration of iloprost at a dosage of 0.01 mg/kg/day (serum levels not available) led to shortened digits of the thoracic extremity in fetuses and pups. In comparable studies in pregnant Sprague-Dawley rats which received iloprost clathrate [13% iloprost by weight] orally at dosages of up to 50 mg/kg/day (C_{oic} of 90 ng/ml), in pregnant rabbits at intravenous dosages of up to 0.5 mg/kg/day (C_{oic} of 86 ng/ml), and in pregnant monkeys at dosages of up to 0.04 mg/kg/day (serum levels of 1 ng/ml), no such digital anomalies or other gross-structural abnormalities were observed in the fetuses/pups. However, in Sprague-Dawley rats where creas of low-viable fetuses at a maternally toxic oral dosage of 250 mg/kg/day and in Han-Wistar rats was found to be embryolethal in 15 of 44 litters at an intravenous dosage of 1 mg/kg/day.

8.3 Nursing Mothers
It is not known whether VENTAVIS is excreted in human milk. In studies with Han-Wistar rats, higher mortality was observed in pups of lactating dams receiving iloprost intravenously at 1 mg/kg daily. In Sprague-Dawley rats, higher mortality was also observed in nursing pups at a maternally toxic oral dose of 250 mg/kg/day of iloprost clathrate (13% iloprost by weight). In rats a passage of low levels of iloprost or metabolites in to the milk was observed (less than 1% of iloprost dose given intraperitoneally). No disturbance of post-natal development and reproductive performance was seen in animals exposed during lactation. Because many drugs are excreted in human milk and because of the potential for serious adverse reactions in nursing infants from VENTAVIS, a decision to discontinue nursing should be made, taking into account the importance of the drug to the mother.

8.4 Pediatric Use
Safety and efficacy in pediatric patients have not been established.

8.5 Geriatric Use
Clinical studies of VENTAVIS did not include sufficient numbers of subjects aged 65 or older to determine whether they respond differently than younger subjects. Other reported clinical experience has not identified differences in responses between elderly and younger patients. In general, dose selection for an elderly patient should be cautious, usually starting at the low end of the dosing range, reflecting the greater frequency of decreased hepatic, renal, or cardiac function and of concomitant disease or other drug therapy.

8.6 Hepatic Impairment
VENTAVIS has not been evaluated in subjects with impaired hepatic function. However, in an intravenous iloprost study in patients with liver cirrhosis, the mean clearance in Child-Pugh Class B subjects (n=5) was approximately 10 mL/min/1.73 m² (half that of healthy subjects). Following oral administration, the mean AUC_{0-8h} in Child-Pugh B Class B subjects (n=3) was 1725 pg*h/mL compared to 177 pg*h/mL in normal subjects (n=4) receiving the same oral iloprost dose. In Child-Pugh Class A subjects (n=5), the mean AUC_{0-8h} was 839 pg*h/mL. Although exposure increased with hepatic impairment, there was no effect on half-life.

8.7 Renal Impairment
VENTAVIS has not been evaluated in subjects with impaired renal function. However, in a study with intravenous infusion of iloprost in patients with end-stage renal failure requiring intermittent dialysis treatment (n=7), the mean AUC_{0-8h} was 230 pg*h/mL compared to 54 pg*h/mL in patients with renal failure (n=8) not requiring intermittent dialysis and 48 pg*h/mL in normals. The half-life was similar in both groups. The effect of dialysis on iloprost exposure has not been evaluated.
In clinical trials of VENTAVIS, no case of overdose was reported. Signs and symptoms to be anticipated are extensions of the dose-limiting pharmacological effects, including hypotension, headache, flushing, nausea, vomiting, and diarrhea. A specific antidote is not known. Interruption of the inhalation session, monitoring, and symptomatic measures are recommended.

10. OVERDOSAGE

11. DESCRIPTION

VENTAVIS (iloprost) Inhalation Solution is a clear, colorless, sterile solution containing iloprost formulated for inhalation via the I-neb® AAD® (Adaptive Aerosol Delivery) System. VENTAVIS is supplied in 1 mL single-use glass ampules containing either 10 mcg/mL or 20 mcg/mL. For the 10 mcg/mL solution, one mL of the solution contains 0.01 mg iloprost, 0.81 mg ethanol, 0.24 mg tromethamine, 9.0 mg sodium chloride, and approximately 0.51 mg hydrochloric acid (for pH adjustment to 8.1) in water for injection. For the 20 mcg/mL solution, one mL of the solution contains 0.02 mg iloprost, 1.62 mg ethanol, 0.24 mg tromethamine, 9.0 mg sodium chloride, and approximately 0.76 mg hydrochloric acid (for pH adjustment to 8.4) in water for injection. The solution contains no preservatives.

The chemical name for iloprost is (E)-(5a,5b,6R,7R,8S,9R,10aR,10bR)-[3a-[3,5-diethyl-2-(3,4-dihydro-2H-pyran-4-yl)-1H-pyrrol-1-yl] -3,4-dihydroxy-4-methyl-1-octen-6-ynyl]-Δ2(1H),Δ5-pentalenevaleric acid. Iloprost consists of a mixture of the 4R and 4S diastereoisomers at a ratio of approximately 53:47. Iloprost is an oily substance, which is sparingly soluble in water, freely soluble in methanol, ethanol, ethyl acetate, acetone, and pH 7 buffer, sparingly soluble in buffer pH 9, and soluble in methanol, ethanol, ethyl acetate, acetone, and pH 7 buffer, sparingly soluble in buffer pH 9, and very slightly soluble in distilled water, buffer pH 3, and buffer pH 5. The molecular formula of iloprost is C22H32O4. Its relative molecular weight is 360.49. The structural formula is shown below:

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COOH
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12. CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Iloprost is a synthetic analog of prostacyclin PGI2. Iloprost dilates systemic and pulmonary arterial vascular beds. It also affects platelet aggregation but the relevance of this effect to the treatment of pulmonary hypertension is unknown. The two diastereoisomers of iloprost differ in their potency in dilating blood vessels, with the 4S isomer substantially more potent than the 4R isomer.

12.3 Pharmacokinetics

Genetics: In pharmacokinetic studies in animals, there was no evidence of interconversion of the two diastereoisomers of iloprost. In human pharmacokinetic studies, the two diastereoisomers were not individually assayed.

Iloprost administered intravenously has linear pharmacokinetics over the dose range of 1 to 3 mg/kg/min. The half-life of iloprost (5 mcg) is 20 to 30 minutes. Following inhalation of iloprost (5 mcg) patients with pulmonary hypertension have iloprost peak plasma levels of approximately 150 pg/mL. Iloprost was generally not detectable in plasma 30 minutes to 1 hour after inhalation.

Absorption and Distribution: The absolute bioavailability of inhaled iloprost has not been determined. Following intravenous infusion, the apparent steady-state volume of distribution was 0.7 to 0.8 L/kg in healthy subjects. Iloprost is approximately 60% protein-bound, mainly to albumin, and this ratio is concentration-independent in the range of 30 to 3000 pg/mL.

Metabolism and Excretion: In vitro studies reveal that cytochrome P450-dependent metabolism plays only a minor role in the biotransformation of iloprost. Iloprost is metabolized principally via β-oxidation of the carboxyl side chain. The main metabolite is tetranor-iloprost, which is found in the urine in the free and conjugated form. In animal experiments, tetranor-iloprost was pharmacologically inactive. Clearance in normal subjects was approximately 20 mL/min/kg.

A mass-balance study using intravenously and orally administered [3H]-iloprost in healthy subjects (n=8) showed recovery of 81% of total radioactivity over 14 hours post-dose, with 68% and 12% recoveries in urine and feces, respectively.

13. NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Iloprost was not mutagenic in bacterial and mammalian cells in the presence or absence of extrinsic metabolic activation. Iloprost did not cause chromosomal aberrations in vitro in human lymphocytes and was not clastogenic in vivo in NMRI/SP mice. There was no evidence of a tumorigenic effect of iloprost clathrate (13% iloprost by weight) in Sprague-Dawley rats dosed orally for up to 8 months at doses of up to 125 mg/kg/day (Cmax of 45 ng/mL serum), followed by 16 months at 100 mg/kg/day, or in Crl:CD-1®(ICR)BR albino mice dosed orally for up to 24 months at doses of up to 125 mg/kg/day (Cmax of 156 ng/mL serum). The recommended clinical dosage regimen for iloprost (5 mcg) affords a serum Cmax of 156 ng/mL. Fertility of males or females was not impaired in Han-Wistar rats at intravenous doses up to 1 mg/kg/day.

The absolute change in 6-minute walk distance (Figure 2) measured (using all available data and no imputation) 30 minutes after inhalation among patients with PAH was greater in the iloprost group compared to the placebo group at all time points. At Week 12, the placebo-corrected difference was 40 meters (p=0.01). When walk distance was measured immediately prior to inhalation, the improvement compared to placebo was approximately 60% of the effect seen at 30 minutes after inhalation.

![Figure 1: Composite Primary Endpoint for PAH Patients (WHO Group I)](image1)

![Figure 2: Change (Mean ± SEM) in 6-Minute Walk Distance 30 Minutes Post-inhalation in PAH Patients (WHO Group I)](image2)
VENTAVIS® (iloprost) Inhalation Solution

Table 2: Treatment Effects by Subgroup among PAH Patients (WHO Group 1)

<table>
<thead>
<tr>
<th>Composite Clinical Endpoint</th>
<th>6-Minute Walk (m)*</th>
</tr>
</thead>
<tbody>
<tr>
<td>n VENTAVIS</td>
<td>n (%)</td>
</tr>
<tr>
<td>All Subjects with PAH**</td>
<td>68</td>
</tr>
<tr>
<td>NYHA III</td>
<td>40</td>
</tr>
<tr>
<td>NYHA IV</td>
<td>28</td>
</tr>
<tr>
<td>Male</td>
<td>23</td>
</tr>
<tr>
<td>Female</td>
<td>45</td>
</tr>
<tr>
<td>Age ≤ 55</td>
<td>41</td>
</tr>
<tr>
<td>Age &gt; 55</td>
<td>27</td>
</tr>
</tbody>
</table>

* Change from baseline to 12 Weeks with measurement 30 minutes after dosing, based on all available data.
** Etiologies of PAH, WHO Group 1 included idiopathic in 62% (n=90), associated with connective tissue disease, including DREST and scleroderma, in 17%, (n=25), associated with anorexigen use in 6% (n=9), heritable PAH in 3% (n=5), other PPH in 3% (n=5), SLE in 1% (n=2), post-partum in 1% (n=2), and overlap/other in 5% (n=8).

Hemodynamic assessments obtained at week 12 before inhalation in both groups (at least 2 hours after a previous dose, trough) and after inhalation in the iloprost group (approximately 15 minutes after a dose, peak), are shown in Table 3. The relationship between hemodynamic changes and clinical effects is unknown.

Table 3: Hemodynamic Parameters Before and After Iloprost Inhalation: Change from Baseline at Week 12*

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Baseline Mean (± SD) change from baseline at Week 12</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Before Inhalation</td>
</tr>
<tr>
<td></td>
<td>Iloprost</td>
</tr>
<tr>
<td>PVR (dyn•s•cm⁻⁵)</td>
<td>1029 ± 390</td>
</tr>
<tr>
<td>mPAP (mmHg)</td>
<td>53 ± 12</td>
</tr>
<tr>
<td>CO (L/min)</td>
<td>3.8 ± 1.1</td>
</tr>
<tr>
<td>SVO₂ (%)</td>
<td>60 ± 8</td>
</tr>
</tbody>
</table>

*Patients of all etiologies of PAH, including CTEPH.

In a small, randomized, double-blind, placebo-controlled study (the STEP trial), 34 patients treated with bosentan 125 mg bid for at least 16 weeks tolerated the addition of inhaled iloprost (up to 5 mcg 6 to 9 times per day during waking hours). The mean daily inhaled dose was 27 mcg and the mean number of inhalations per day was 5.6.

16. HOW SUPPLIED/STORAGE AND HANDLING
VENTAVIS (iloprost) Inhalation Solution is supplied in cartons of 30 x 1 mL clear glass single-use ampules as follows:
1 mL ampule containing iloprost 10 mcg per mL, carton of 30 (NDC 68215-302-30)
1 mL ampule containing iloprost 20 mcg per mL, carton of 30 (NDC 68215-303-30)

STORAGE
Store at 20 – 25°C (68 – 77°F)
Excursions permitted to 15 – 30°C (59 – 86°F)
[See USP Controlled Room Temperature]

17. PATIENT COUNSELING INFORMATION
Patients receiving VENTAVIS should be advised to use the drug only as prescribed with the I-neb® AAD® System, following the manufacturer’s instructions. Patients should be trained in proper administration techniques including dosing frequency, ampule dispensing, I-neb® AAD® System operation, and equipment cleaning.
Advise patients that they may have a fall in blood pressure with VENTAVIS, so they may become dizzy or even faint. They should stand up slowly when they get out of a chair or bed. If fainting gets worse, patients should consult their physicians about dose adjustment.
Advise patients that VENTAVIS should be inhaled at intervals of not less than 2 hours and that the acute benefits of VENTAVIS may not last 2 hours. Thus patients may want to adjust times of administration to cover planned activities.
See FDA-approved patient labeling.

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ACT20131125

Manufactured for:
Actelion Pharmaceuticals US, Inc.
5000 Shoreline Court, Ste. 200
South San Francisco, CA 94080
VENTAVIS® Inhalation Solution

VENTAVIS® (ven TAY vis)

• You should not take VENTAVIS more than every 2 hours. The benefits of VENTAVIS may not last if you use it more frequently.

• Take VENTAVIS exactly as your doctor tells you to take it. Your doctor may change your dose if needed.

See the end of this leaflet for detailed instructions for using VENTAVIS.

How should I take VENTAVIS?

- medicines used to treat high blood pressure or heart problems

Especially tell your doctor if you take:

VENTAVIS and other medicines may affect each other causing side effects. VENTAVIS may affect the way other medicines work, and other medicines may affect how VENTAVIS works.

Tell your doctor about all the medicines you take, including prescription and nonprescription medicines, vitamins, and herbal supplements.

VENTAVIS and other medicines may affect each other causing side effects; VENTAVIS may affect the way other medicines work, and other medicines may affect how VENTAVIS works. Especially tell your doctor if you take:

- medicines used to treat high blood pressure or heart problems

- medicines that lessen blood clotting (warfarin, Coumadin, Jantoven)

Know the medicines you take. Keep a list of your medicines and show it to your doctor and pharmacist when you get a new medicine.

How should I take VENTAVIS?

See the end of this leaflet for detailed instructions for using VENTAVIS.

- Take VENTAVIS exactly as your doctor tells you to take it. Your doctor may change your dose if needed.

- You should not take VENTAVIS more than every 2 hours. The benefits of VENTAVIS may not last 2 hours, so you may adjust the times that you use it to cover planned activities.

- Do not drink VENTAVIS.

- Do not let VENTAVIS come into contact with your skin or eyes. If it does, rinse your skin or eyes with water right away.

- Do not allow other people to be exposed to VENTAVIS while you are breathing it, especially babies and pregnant women.

- If you take too much VENTAVIS, you may have a headache, red face, dizziness, nausea, vomiting and diarrhea. If this happens stop taking VENTAVIS. If your symptoms do not go away, call your doctor or get emergency help right away.

What are the possible side effects of VENTAVIS?

VENTAVIS may cause side effects, including feeling dizzy, lightheaded and faint. If you have any of these side effects, you should stand up slowly when you get out of chairs or bed. Tell your doctor if your fainting gets worse during treatment with VENTAVIS. Your doctor may need to change your dose or your treatment.

Do not drive a car or operate any tools or machines if dizziness or fainting from low blood pressure is a problem for you.

You may have trouble breathing after taking VENTAVIS because it may cause the muscles around your airway to tighten (bronchospasm). Get emergency help right away if you have trouble breathing.

Other important side effects of VENTAVIS include:

- bleeding

- red face (flushing)

- increased cough

- low blood pressure

- headaches

- nausea

- spasm of your jaw muscles that makes it hard to open your mouth

Talk to your doctor if you have any side effect that bothers you or that does not go away. These are not all the possible side effects of VENTAVIS. For more information, ask your doctor or pharmacist.

Call your doctor for medical advice about side effects. You may report side effects to the FDA at 1-800-FDA-1088.

How should I store VENTAVIS?

- Store VENTAVIS between 68°F to 77°F (20°C to 25°C).

- Safely throw away VENTAVIS that is out of date or no longer needed.

Keep VENTAVIS and all medicines out of the reach of children.

General Information about VENTAVIS

Medicines are sometimes prescribed for conditions that are not listed in the patient leaflet. Do not use VENTAVIS for a condition for which it was not prescribed. Do not give VENTAVIS to other people, even if they have the same symptoms you have. It may harm them.

This patient information leaflet summarizes the most important information about VENTAVIS. If you would like more information about VENTAVIS talk with your doctor. You can ask your doctor or pharmacist for information about VENTAVIS that is written for health professionals. For more information go to www.VENTAVIS.com or call 1-888-228-3546.

What are the ingredients in VENTAVIS?

Active ingredient: iloprost

Inactive ingredients: tromethamine, ethanol, sodium chloride, hydrochloric acid for pH adjustment, and water for injection.

VENTAVIS is a clear, colorless solution.

Patient Instructions for Using VENTAVIS

To take VENTAVIS, you will need to use the I-neb Adaptive Aerosol Delivery (AAD) System. This system is used to give you the right dose of VENTAVIS. You should not use other systems to take VENTAVIS as other systems may not give you the amount of VENTAVIS you need.

Do not use VENTAVIS until your doctor has showed you how to use this system the right way. Make sure you understand all the instructions or ask questions until you do.

I-neb AAD System

With the I-neb System, you will receive two medicine chambers (one with a red latch and one with a purple latch) and two color-matching dosing discs to use with the 10 micrograms per 1 mL of VENTAVIS.

You should use the red dosing disc with the red latched medicine chamber (gives you a 2.5 microgram dose). You should use the purple dosing disc with the purple-latched medicine chamber (gives you a 5 microgram dose). Always use all of the contents of only 1 ampule when using the I-neb System.

If you are using the I-neb System and usually have long treatment times, your doctor may ask you to switch to a third medicine chamber (one with a gold latch). The medicine chamber with the gold latch and matching dosing disc are only for use with the 20 micrograms per 1 mL ampule of VENTAVIS. You should use the gold dosing disc with the gold latched medicine chamber (gives you a 5 microgram dose).

Do not change the medicine chamber and dosing disc in your I-neb System without talking to your doctor.

Do not put any medicines other than VENTAVIS in your I-neb System.

VENTAVIS® (iloprost) Inhalation Solution

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Do not change the medicine chamber and dosing disc in your I-neb System without talking to your doctor.

Do not put any medicines other than VENTAVIS in your I-neb System.
To Use VENTAVIS:
Open the small glass bottle (ampule) of VENTAVIS by using either an ampule breaker or a rubber
pad. Use either the ampule breaker or the rubber pad. You do not need to use both methods to open
a VENTAVIS ampule.

When using an ampule breaker:
Step 1. Line up the blue dot on the VENTAVIS ampule with the dot on the ampule breaker, if available,
and then insert the top of the ampule into the ampule breaker.

Step 2. Gently break open the neck of the ampule by pushing away from the dot on the VENTAVIS
ampule to snap off the ampule lid.

Step 3. Carefully throw away the top of the ampule into a safe container.

When using a rubber pad:
Step 1. Hold the ampule with the blue dot facing away from your body.

Step 2. Wrap the rubber pad around the ampule to protect yourself from getting cut.

Step 3. Use your thumbs to break open the neck of the ampule by snapping the top toward you.

Step 4. Using the small tube (pipette) that comes with VENTAVIS, draw-up the entire amount of
one ampule of VENTAVIS and empty it into the center of the I-neb System medicine chamber. The
amount of VENTAVIS you receive will be controlled by either the dosing disc or the medicine chamber.

I-neb System

Step 5. Throw away in a safe container the:
• top of the ampule (the ampule lid)
• open ampule
Keep both the ampule and the pipette out of the reach of children.

Step 6. To breathe in your dose of VENTAVIS, follow the instructions that come with your I-neb
System. Each treatment session with VENTAVIS lasts about 4 to 10 minutes. Call your doctor if you
usually have longer treatment times as your dose may need to be changed.
The I-neb System allows you to stop your treatment for up to ten minutes with no effect on the
final dose you get. If your treatment is stopped for more than ten minutes, the I-neb System will
reset itself. If that happens, throw away the solution in the chamber and wait at least two
hours before taking your next dose. If you take a second dose right away you could get
too much medicine.

Step 7. Throw away any VENTAVIS that is left in the medicine chamber after each treatment. Do not
use the rest of the VENTAVIS because it will not give you the right dose.

Step 8. Clean your system after each treatment. Follow the instructions that come with your system.
Step 9. Make sure you have access to a back-up I-neb System to use for VENTAVIS treatments. This
is important if your original system does not work for some reason.