

VENTAVIS® (iloprost) Inhalation Solution is 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 predominantly patients with NYHA Functional Class III-IV symptoms and etiologies of idiopathic or heritable PAH (65%) or PAH associated with connective tissue disease (23%).

VENTAVIS Patient Enrollment Form for VA Medical Facilities

Patient name:

IMPORTANT SAFETY INFORMATION

WARNINGS AND PRECAUTIONS

Risk of Syncope

- Vital signs should be monitored while initiating VENTAVIS. Hypotension leading to syncope has been observed; VENTAVIS should therefore not be initiated in patients with systolic blood pressure less than 85 mmHg.

Pulmonary Venous Hypertension

- Stop VENTAVIS immediately if signs of pulmonary edema occur; this may be a sign of pulmonary venous hypertension.

Bronchospasm

- VENTAVIS inhalation may cause bronchospasm and patients with a history of hyperreactive airway disease may be more sensitive.

ADVERSE REACTIONS

Serious Adverse Events

- Serious adverse events reported include congestive heart failure, chest pain, supraventricular tachycardia, dyspnea, peripheral edema, and kidney failure.

Adverse Events

- Adverse events reported in a Phase 3 clinical trial occurring with a $\geq 3\%$ difference between VENTAVIS patients and placebo patients were vasodilation (flushing) (27% vs 9%), increased cough (39% vs 26%), headache (30% vs 20%), trismus (12% vs 3%), insomnia (8% vs 2%), nausea (13% vs 8%), hypotension (11% vs 6%), vomiting (7% vs 2%), alkaline phosphatase increased (6% vs 1%), flu syndrome (14% vs 10%), back pain (7% vs 3%), tongue pain (4% vs 0%), palpitations (7% vs 4%), syncope (8% vs 5%), GGT increased (6% vs 3%), muscle cramps (6% vs 3%), hemoptysis (5% vs 2%), and pneumonia (4% vs 1%).

DRUG INTERACTIONS

Antihypertensives and Vasodilators

- VENTAVIS has the potential to increase the hypotensive effect of vasodilators and antihypertensive agents.

Anticoagulants and Platelet Inhibitors

- VENTAVIS also has the potential to increase risk of bleeding, particularly in patients maintained on anticoagulants or platelet inhibitors.

Please see accompanying full Prescribing Information.



VENTAVIS® (iloprost) Inhalation Solution—Veterans Administration

SPECIALTY PHARMACY SERVICES ENROLLMENT FORM



All fields must be completed to expedite prescription fulfillment						
Physician information	Name:		DEA # (optional):		NPI #:	
	Name of facility:		MD specialty:		UPIN #:	
	Contact name and phone #:		State license #:		Phone #:	
	Address:		Suite:	City:	State:	ZIP:
	PCP (if applicable/different from prescribing MD): _____				Phone #: _____	Fax #: _____
Patient information	Name:			DOB:		
	Address:		City:	State:	ZIP:	
	Preferred language (if not English):			Phone #:	Sex: Male Female	
	Caregiver name (if applicable):				Alternate phone #:	
VA pharmacy information	Name of facility:					
	Address:		Suite:	City:	State: ZIP:	
	Contact name:			Contact phone #:	Contact fax #:	
	Ship-to address:		Suite:	City:	State: ZIP:	
	Purchase order #:					
	Ship to: Patient VA location					
Prescription	Statement of medical necessity					
	DIAGNOSIS:					
	Primary Arterial Pulmonary Hypertension – ICD-9 416.0 Date of Onset ___/___/___		Secondary Arterial Pulmonary Hypertension – ICD-9 416.8 Date of Onset ___/___/___		Other _____ ICD-9 _____ Date of Onset ___/___/___	
	New York Heart Association (NYHA) Functional Classification I II III IV					
	FAX COPIES OF MOST RECENT:					
	History and Physical	ANA Results	VQ/Perfusion Scan	Cardiac Catheter Report	Echocardiogram	
	Pain Management Protocol	Physical Statement	6-Minute Walk Test			
	(Treatment with oral calcium blocking agents has been tried and failed or has been considered and ruled out)					
	NURSING NEEDS (check all that apply):					
	Start of Care Date ___/___/___	Number of Visits _____	Pre-Hospital/Pre-Home Teaching	In-Hospital Teaching	Nursing Follow-Up	
ORDERS:						
Start of Care Date ___/___/___	Patient Status	Urgent/Patient in Hospital	Projected Start Date ___/___/___	Hospital Contact _____		
Rx						
VENTAVIS Equipment I-neb® AAD® Device(s) 2.5 mcg Initial Dose, Then 5.0 mcg Ongoing Frequency _____ Times Per Day (Waking Hours) Dispense _____ Month Supply Ancillary Supplies Provided as Needed for Administration.		Prescriber's Notes Refill: PRN _____ Times In _____ Months Dispense As Written Substitution Allowed				
Prescriber's Signature				Date		

Please provide completed form to the VA pharmacy for review and forwarding to Caremark Specialty Pharmacy.

Caremark Telephone: 1-877-242-2738 Fax: 1-877-943-1000

Please see accompanying full Prescribing Information.

Caremark is committed to protecting the privacy of your health information. We will hold your health information in confidence and will only use and disclose it in accordance with applicable law.

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