

STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

ILOPROST

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
ILOPROST	VENTAVIS	26287		GPI-10	
TROMETHAMINE				(4017006000)	

GUIDELINES FOR USE

INITIAL CRITERIA (NOTE: FOR RENEWAL CRITERIA SEE BELOW)

- 1. Does the patient have a diagnosis of pulmonary arterial hypertension (PAH) (WHO Group 1) **AND** meet the following criterion?
 - Therapy is prescribed by or in consultation with a cardiologist or pulmonologist

If yes, continue to #2. If no, do not approve.

DENIAL TEXT: See initial denial text at the end of the guideline.

- 2. Does the patient have documentation (e.g., chart note, lab result, diagnostic test result, etc.) confirming PAH diagnosis based on right heart catheterization with **ALL** of the following parameters?
 - Mean pulmonary artery pressure (PAP) of greater than 20 mmHg
 - Pulmonary capillary wedge pressure (PCWP) of less than or equal to 15 mmHg
 - Pulmonary vascular resistance (PVR) of greater than 2 Wood units

If yes, continue to #3. If no, do not approve.

DENIAL TEXT: See initial denial text at the end of the guideline.

- 3. Has the patient had a trial of or contraindication to **TWO** of the following agents from different drug classes?
 - Oral endothelin receptor antagonist (e.g., Letairis [ambrisentan], Tracleer [bosentan], Opsumit [macitentan])
 - Oral phosphodiesterase-5 inhibitor (e.g., Revatio [sildenafil], Adcirca [tadalafil])
 - Oral cGMP stimulator (e.g., Adempas [riociguat])
 - IV/SQ prostacyclin (e.g., Flolan [epoprostenol], Remodulin [Treprostinil])

If yes, approve for 12 months by HICL or GPI-10.

If no, do not approve.

DENIAL TEXT: See the initial denial text at the end of the guideline.

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Revised: 5/23/2023 Page 1 of 3



STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

ILOPROST

INITIAL CRITERIA (CONTINUED)

INITIAL DENIAL TEXT: *Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **ILOPROST (Ventavis)** requires the following rule(s) be met for approval:

- A. You have pulmonary arterial hypertension (PAH: type of high blood pressure that affects arteries in the lungs and in the heart) World Health Organization (WHO Group 1: a way to classify the severity of disease)
- B. Therapy is prescribed by or in consultation with a cardiologist (heart doctor) or pulmonologist (lung doctor)
- C. There is documentation (such as chart note, lab result, diagnostic test result) showing you have pulmonary arterial hypertension based on all of the following lab values by putting a catheter (narrow flexible tube) into the right side of your heart:
 - 1. Mean pulmonary artery pressure (PAP) greater than 20 mmHg
 - 2. Pulmonary capillary wedge pressure (PCWP) less than or equal to 15 mmHg
 - 3. Pulmonary vascular resistance (PVR) greater than 2 Wood units
- D. You had a trial of or contraindication (harmful for) to TWO of the following agents from different drug classes:
 - 1. Oral endothelin receptor antagonist (such as Letairis [ambrisentan], Tracleer [bosentan], Opsumit [macitentan])
 - 2. Oral phosphodiesterase-5 inhibitor (such as Revatio [sildenafil], Adcirca [tadalafil])
 - 3. Oral cGMP stimulator (such as Adempas [riociguat])
 - 4. Intravenous or subcutaneous prostacyclin (such as Flolan [epoprostenol], Remodulin [Treprostinil])

Your doctor told us [INSERT PT SPECIFIC INFO PROVIDED]. We do not have information showing you [INSERT UNMET CRITERIA]. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

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Revised: 5/23/2023 Page 2 of 3



STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

ILOPROST

GUIDELINES FOR USE (CONTINUED)

RENEWAL CRITERIA

- 1. Does the patient have a diagnosis of pulmonary arterial hypertension (PAH) (WHO Group 1) and meet **ONE** of the following criteria?
 - The patient shown improvement from baseline in the 6-minute walk distance test
 - The patient remains stable from baseline in the 6-minute walk distance test AND the patient's WHO functional class remained stable or has improved

If yes, approve for 12 months by HICL or GPI-10. If no, do not approve.

RENEWAL DENIAL TEXT: *Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **ILOPROST** (Ventavis) requires the following rule(s) be met for renewal:

- A. You have pulmonary arterial hypertension (PAH: type of high blood pressure that affects arteries in the lungs and in the heart) World Health Organization (WHO Group 1: a way to classify the severity of disease)
- B. You meet ONE of the following:
 - 1. You have shown improvement from baseline in the 6-minute walk distance test
 - 2. You remain stable in the 6-minute walk distance test AND your World Health Organization functional class has remained stable or improved (WHO-FC: classification system for heart failure)

Your doctor told us [INSERT PT SPECIFIC INFO PROVIDED]. We do not have information showing you [INSERT UNMET CRITERIA]. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

RATIONALE

For further information, please refer to the prescribing information and/or drug monograph for Ventavis.

REFERENCES

Ventavis [Prescribing Information]. South San Francisco, CA: Actelion; January 2021.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Created: 01/08

Commercial Effective: 07/01/23 Client Approval: 05/23 P&T Approval: 04/23

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Revised: 5/23/2023 Page 3 of 3