

REVATIO, LIQREV (sildenafil)**Pre - PA Allowance**None

Prior-Approval Requirements**Diagnoses**Patient must have **ONE** of the following

1. Pulmonary Arterial Hypertension - WHO Group I
 - a. NYHA functional classification of physical activity - Class II or III
 - b. Prescribed by or recommended by a cardiologist or pulmonologist
2. Raynaud's syndrome
 - a. Inadequate treatment response, intolerance, or contraindication to **TWO** of the following:
 - i. Calcium channel blockers
 - ii. Alpha adrenergic receptor blockers
 - iii. Angiotensin II receptor antagonist

AND NONE of the following:

1. Concurrent therapy with any nitrates (in any form)
2. Concurrent therapy with another phosphodiesterase-5 (PDE5) inhibitor
3. Concurrent therapy with Guanylate Cyclase (GC) Stimulators
4. Concurrent therapy with alpha blockers

AND ALL of the following:

1. Prescriber agrees to counsel and evaluate the patient for sudden loss of vision or hearing associated with this medication
2. **Brand Revatio only:** Patient **MUST** have tried the preferred product (generic Revatio: sildenafil) unless the patient has a valid medical exception (e.g. inadequate treatment response, intolerance, contraindication)

Prior - Approval Limits**Duration** 2 years

Prior – Approval *Renewal* Requirements



Federal Employee Program.

REVATIO, LIQREV (sildenafil)

Diagnoses

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2. Raynaud's syndrome

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2. Concurrent therapy with another phosphodiesterase-5 (PDE5) inhibitor
3. Concurrent therapy with Guanylate Cyclase (GC) Stimulators
4. Concurrent therapy with alpha blockers

AND ALL of the following:

1. Symptoms have improved or stabilized
2. Prescriber agrees to counsel and evaluate the patient for sudden loss of vision or hearing associated with this medication
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Prior – Approval *Renewal* Limits

Same as above