



2024 PRIOR AUTHORIZATION REQUEST FORM
Individual and Family Plans

Uptravi

Fax back to: (833) 605-4407

Phone: (215) 991-4300

Jefferson Health Plans manages the pharmacy drug benefit for your patient. Certain requests for coverage require review with the prescribing physician. Please answer the following questions and fax this form to the number listed above.

PLEASE NOTE: Any information (patient, prescriber, drug, labs) left blank, illegible, or not attached WILL delay the review process.

Form with fields: Patient Name, Prescriber Name, Member Number, Fax, Phone, Date of Birth, Office Contact, Line of Business, NPI, State Lic ID, Address, City, State ZIP, Primary Phone, Specialty/facility name (if applicable)

REQUEST FOR EXPEDITED REVIEW: By checking this box and signing below, I certify that the standard review timeframe may seriously jeopardize the life or health of the enrollee or the enrollee's ability to regain maximum function.

Form with fields: Drug Name, Strength, Directions / SIG

Please attach any pertinent medical history including labs and information for this member that may support approval. Please answer the following questions and sign.

Q1. Is Uptravi being prescribed by or in consultation with a cardiologist or pulmonologist?

Yes No

Q2. Is the patient 18 years of age or older?

Yes No

Q3. Does the patient have a diagnosis of World Health Organization (WHO) group 1 pulmonary arterial hypertension (PAH)?

Yes No

Q4. Has the diagnosis of PAH been confirmed by a complete right catheterization (RHC) (please attach RHC report)? PAH is defined as: I. A mean pulmonary arterial pressure (mPAP) greater than 20 mmHg; II. A pulmonary capillary wedge pressure (PCWP) less than or equal to 15 mmHg; III. A pulmonary vascular resistance (PVR) greater than 3 Wood units

Yes No



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<b>Patient Name:</b>	<b>Prescriber Name:</b>
Q5. Does the patient have a World Health Organization (WHO) functional class of: II (Slight limitation of physical activity but comfortable at rest. Ordinary physical activity causes undue dyspnea or fatigue, chest pain, or near syncope), or III (Marked limitation of physical activity and comfortable at rest. Less than ordinary activity causes undue dyspnea or fatigue, chest pain, or near syncope)?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Q6. Are pharmacy records or chart notes provided documenting trial of or inadequate response to two of the following alternatives (used alone or in combination): I. Endothelin Receptor Antagonists (bosentan, ambrisentan, macitentan); II. Phosphodiesterase-5 inhibitors (sildenafil, tadalafil); III. Guanylate Cyclase stimulators (riociguat)?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Q7. Is there a treatment plan?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Q8. Will Uptravi be used along with a strong CYP2C8 inhibitor (eg gemfibrozil)?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Q9. Does the patient have hepatic impairment (Child Pugh class B or greater) with lab monitoring and dose adjustments as needed?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Q10. Additional Information:	
Q11. Duration:	
<input type="checkbox"/> 12 months <input type="checkbox"/> Other:	

\_\_\_\_\_  
Prescriber Signature

\_\_\_\_\_  
Date



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<b>Patient Name:</b>	<b>Prescriber Name:</b>
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2024 Prior Authorization Request