

ANTIFIBROTIC RESPIRATORY AGENTS PRIOR AUTHORIZATION FORM

Prior authorization guidelines for **Antifibrotic Respiratory Agents** and **Quantity Limits/Daily Dose Limits** are available on the DHS Pharmacy Services website at https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/default.aspx.

☐New request ☐Renewal request	# of pages:	Prescriber name:				
Name of office contact:	Specialty:					
Contact's phone number:		NPI:		State license #:		
LTC facility contact/phone:		Street address:				
Beneficiary name:		City/State/Zip:				
Beneficiary ID#:	DOB:	Phone:		Fax:		
CLINICAL INFORMATION						
Drug requested:	Strength:	Formulation (powder, tablet, etc.):				
Dose/directions:			Quantit	Quantity: Refills:		
Diagnosis (submit documentation):			Dx code (<u>required</u>):			
Is the medication being prescribed by or in consultation with a pulmonologist, rheumatologist, or other specialist?			☐Yes Submit documentation of ☐No consultation, if applicable.			
Is the beneficiary currently being treated with the requested medication?			□Yes □No	TI VAS SUNMIT ANCUMENTATION I		
If applicable, has the dose of the requested medication been adjusted for the beneficiary's degree of liver impairment, concomitant medications, adverse effects, etc.?				☐Yes Submit documentation.		
INITIAL requests						
For a non-preferred Antifibrotic Respiratory Agent, does the beneficiary have a history of trial and failure of or a contraindication or an intolerance to the preferred Antifibrotic Respiratory Agents appropriate for the beneficiary's diagnosis or indication? Refer to https://papdl.com/preferred-drug-list for a list of preferred and non-preferred drugs in this class.				Yes Submit documentation. No		
Is the beneficiary a current smoker? If yes, did the prescriber advise the beneficiary to stop smoking?			□Yes □Yes		Submit cumentation.	
RENEWAL requests						
Has the beneficiary experienced a positive clinical response to the requested medication?				☐Yes ☐No Submit documentation.		
Did the beneficiary experience any adverse reactions that require dose adjustment as described in the FDA-approved product labeling (e.g., liver enzyme elevations, GI reaction, photosensitivity reaction, rash)?				☐Yes ☐No Submit documentation.		
PLEASE <u>FAX</u> COMPLETED FORM WITH <u>REQUIRED CLINICAL DOCUMENTATION</u> TO DHS – PHARMACY DIVISION						
Prescriber Signature:			Date:			

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