]	Member Name: {{MEMFIF	SST}} {{MEMLAST}} <b>DOB:</b> {{MEMBERDOB}} <b>PA Number:</b> {{PANUMBER}}
{ {F	ANUMCODE}}	
	DISPLAY_PAGNAME}} ACDESCRIPTION}}	
for:	ns to {{COMPANY_NAME	secure location as required by HIPAA regulations. Fax complete signed and dated }} at {{CLIENT_PAG_FAX}}. Please contact {{COMPANY_NAME}} at th questions regarding the prior authorization process. When conditions are met, we {DRUGNAME}}.
Pat Phy Spe Phy	cient's ID: {{MEMBERID}} ysician's Name: {{PHYFIR ecialty: ysician Office Telephone: {	T}} {{MEMLAST}} Date: {{TODAY}} Patient's Date of Birth: {{MEMBERDOB}} ST}} {{PHYLAST}} Patient Phone: << MEMPHONE>> NPI#: [{PHYSICIANPHONE}} Physician Office Fax: {{PHYSICIANFAX}} PHYADDRESS1>> << PHYADDRESS2>> << PHYCITY>>, << PHYSTATE>>
<<]	PHYZIP>>	
	ug Name: {{DRUGNAME}	
1.	What is the prescribed production of the Different of the	luct?  4mg □ generic pirfenidone (excluding 534mg) □ Esbriet
2.	What is the diagnosis? ☐ Idiopathic pulmonary fib	prosis (IPF)
3.	What is the ICD-10 code?	
Sec		Complete this section if pirfenidone 534mg is prescribed
4.	three of the formulary medialternatives for the requeste	•
5.	Has the patient tried and hat the formulary medications, medications should be presented.	d a documented inadequate response or intolerable adverse reaction to at least three of or all of the formulary alternatives if there are fewer than three? Note: Formulary cribed first unless the patient is unable to use or receive treatment with the alternative. fev and generic pirfenidone (excluding 534mg).   Yes No
	If Yes, indicate the formulary alternative(s) the patient has tried and the reason(s) for treatment failure and skip to #7.	
	Drug name:	Reason for treatment failure:
	Drug name:	Reason for treatment failure:
6.	Does the patient have a documented contraindication to all or at least three of the formulary alternative(s): Ofev and generic pirfenidone (excluding 534mg)?    Yes   No	
	If Yes, specify the formulary alternative(s) the patient is unable to take and describe the contraindication(s):	
	Drug name:	Contraindication:
	Drug name:	Contraindication:
7.	contraindication to at least fewer than three been attac indicating prior treatment	ocumentation supporting the inadequate response, intolerable adverse reaction, or three of the formulary medications, or all of the formulary alternatives if there are hed? ACTION REQUIRED: Submit chart note(s) or other documentation failure, severity of the adverse event (if any), and dosage and duration of the prior tion to formulary alternatives.   Yes  No Skip to Section C: All Requests

Member Name: {{MEMFIRST}} {{MEMLAST}} DOB: {{MEMBERDOB}} PA Number: {{PANUMBER}}		
Section B: Preferred Product - Complete this section if Esbriet is prescribed  8. The preferred products for your patient's health plan are Ofev and generic pirfenidone. Can the patient's treatment be switched to a preferred product?  If Yes, pirfenidone, please fax a new prescription to the pharmacy and skip to Section C: All Requests.  Yes, please specify:  No - Continue request for non-preferred product		
9. Does the patient have a documented intolerable adverse event to the preferred product, generic pirfenidone? <i>ACTION REQUIRED: If Yes, attach supporting chart note(s).</i> □ Yes □ No		
Was the documented intolerable adverse event an expected adverse event attributed to the active ingredient (pirfenidone) as described in the prescribing information? <i>ACTION REQUIRED: If No, attach supporting chart note(s).</i> □ Yes □ No		
11. Does the patient have a documented inadequate response to the preferred product, Ofev?  **ACTION REQUIRED: If Yes, attach supporting chart note(s) and skip to Section C: All Requests  □ Yes □ No		
12. Does the patient have a documented intolerable adverse event to the preferred product, Ofev? <i>ACTION REQUIRED: If Yes, attach supporting chart note(s).</i> □ Yes □ No		
Section C: All Requests  13. Is the patient currently receiving treatment with the requested medication?   Yes   No If No, skip to #15		
<ul> <li>14. Is the patient currently receiving the requested medication through samples or a manufacturer's patient assistance program?</li> <li>☐ Yes</li> <li>☐ No, no further questions.</li> <li>☐ Unknown</li> </ul>		
15. Have other known causes of interstitial lung disease (e.g., domestic and occupational environmental exposures, connective tissue disease, drug toxicity) been excluded? □ Yes □ No If No, no further questions.		
16. Has the patient undergone a high-resolution computed tomography (HRCT) study of the chest? <i>ACTION REQUIRED: If Yes, attach the radiology report.</i> □ Yes □ No <i>If No, skip to #20</i>		
17. Please indicate what the high-resolution computed tomography (HRCT) scan demonstrates.  ☐ Usual interstitial pneumonia (UIP) pattern, <i>no further questions</i> ☐ Other (e.g., probable UIP, indeterminate for UIP, or alternative diagnosis)		
18. Has the diagnosis of idiopathic pulmonary fibrosis been supported by a lung biopsy? <i>ACTION REQUIRED: If Yes, attach the pathology report and no further questions.</i> □ Yes □ No		
<ul> <li>19. Has the diagnosis of idiopathic pulmonary fibrosis been supported by a multidisciplinary discussion between at least a pulmonologist and a radiologist who are experienced in idiopathic pulmonary fibrosis?</li> <li>□ Yes □ No No further questions.</li> </ul>		
20. Has the patient undergone a lung biopsy? <i>ACTION REQUIRED: If Yes, attach the pathology report.</i> ☐ Yes ☐ No <i>If No, no further questions.</i>		
21. Please indicate what the biopsy report demonstrates.  ☐ Usual interstitial pneumonia (UIP) pattern ☐ Other		
I attest that the medication requested is medically necessary for this patient. I further attest that the information provided is accurate and true, and that the documentation supporting this information is available for review if requested by the claims processor, the health plan sponsor, or, if applicable a state or federal regulatory agency.		

Prescriber (Or Authorized) Signature and Date