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|-------------------------|------------------------------|
| BILLING CODE            | Must use valid NDC           |
| BENEFIT TYPE            | Pharmacy                     |
| SITE OF SERVICE ALLOWED | Home                         |
| STATUS                  | Prior Authorization Required |

Esbriet is a pyridone oral antifibrotic drug approved by the FDA in 2014. Idiopathic pulmonary fibrosis (IPF) is an interstitial lung disease characterized by chronic, progressive scarring of the lungs and the pathological hallmark of usual interstitial pneumonia (UIP).

Esbriet (pirfenidone) will be considered for coverage when the following criteria are met:

## Idiopathic Pulmonary Fibrosis (IPF)

For initial authorization:

- 1. Member is at least 18 years of age; AND
- 2. Medication must be prescribed by or in consultation with a pulmonologist; AND
- 3. Member has a diagnosis of IPF confirmed by high resolution computed tomography (HRCT) or lung biopsy (results must be submitted for review); AND
- 4. Documentation of member's baseline forced vital capacity (FVC) must be equal to or greater than 50% predicted; AND
- 5. Member does not have severe hepatic impairment (Child Pugh Class C); AND
- 6. Member is not a current smoker and provider attests the member will not smoke during treatment.
- Dosage allowed/Quantity limit: Titrate as follows, to 801 mg three times per day (2403 mg/day total) (90 tablets per 30 days).

| Treatment days  | Dosage                              |  |  |  |  |  |  |
|---|-------------------------------------|--|--|--|--|--|--|
| Benezi Arminitary and 26 months of mondaile (201 unighted and |                                     |  |  |  |  |  |  |
| s daily (1602 mg/day)   | Days 8 through 14 534 mg three time |  |  |  |  |  |  |
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If all the above requirements are met , the medication will be approved for 6 months.

## For reauthorization:

- 1. Member continues to abstain from smoking; AND
- 2. Chart notes must show improvement or stabilized signs and symptoms of disease demonstrated by reduced rate of FVC decline.

If all the above requirements are met , the medication will be approved for an additional 12 months.