

**Generic Name:** Pirfenidone

**Preferred:** N/A

**Therapeutic Class or Brand Name:** Esbriet®

**Non-preferred:** N/A

**Applicable Drugs (if Therapeutic Class):** N/A

**Date of Origin:** 8/30/2016

**GPI Code:** 4555006000

**Date Last Reviewed / Revised:** 8/28/2019

## PRIOR AUTHORIZATION CRITERIA

(May be considered medically necessary when criteria I through VI are met)

- I. Documented diagnosis of idiopathic pulmonary fibrosis (IPF).
- II. Documentation that patient has a baseline percent predicted forced vital capacity (%FVC) greater than or equal to 50%.
- III. Documentation that the patient has a baseline carbon monoxide (%DLCO) greater than or equal to 30%.
- IV. Documentation that patient does not smoke and has not smoked for a minimum of six weeks.
- V. Minimum age requirement: 18 years old.
- VI. The prescriber is a Pulmonologist.

## EXCLUSION CRITERIA

- Coadministration of Esbriet® with Ofev® (nintedanib).
- Coadministration of Esbriet® with any of the following:
  - Fluvoxamine or other strong CYP1A2 inhibitors (i.e. enoxacin).
  - Moderate or strong inhibitors of both CYP1A2 and one or more other CYP isoenzymes involved in the metabolism of Esbriet® (i.e. CYP2C9, 2C19, 2D6, and 2E1).
  - Strong CYP1A2 inducers.

## OTHER CRITERIA

- N/A.

## QUANTITY / DAYS SUPPLY RESTRICTIONS

- 267 mg strength: Quantities of up to 270 capsules/tablets per 30 days.
- 801 mg strength: Quantities of up to 90 tablets per 30 days.

**APPROVAL LENGTH**

- **Authorization:** 6 months
- **Re-Authorization:** An updated letter of medical necessity or progress notes showing current medical necessity criteria are met and that the medication is effective. Documentation of both a and b are also required:
  - a. There is less than a 200 ml decrease in FVC or less than a 10% decline in %FVC.
  - b. The patient does not smoke.

**APPENDIX**

N/A.

**REFERENCES**

1. [http://www.gene.com/download/pdf/esbriet\\_prescribing.pdf](http://www.gene.com/download/pdf/esbriet_prescribing.pdf).
2. Medi-Span®.

**HISTORICAL TRACKING OF CHANGES MADE TO POLICY**

Date	Notes/Changes
8/28/2019	1. <b>Removed under References:</b> <a href="https://regence.myprime.com/content/dam/prime/memberportal/forms/AuthorForms/Cambia/Program_Summaries/dru444reg.pdf">https://regence.myprime.com/content/dam/prime/memberportal/forms/AuthorForms/Cambia/Program_Summaries/dru444reg.pdf</a> .  <a href="https://ahca.myflorida.com/medicaid/Prescribed_Drug/drug_criteria_pdf/Esbriet_Criteria.pdf">https://ahca.myflorida.com/medicaid/Prescribed_Drug/drug_criteria_pdf/Esbriet_Criteria.pdf</a> .
8/23/2018	2. <b>Changed</b> obsolete URL in Reference #3 ( <a href="http://blue.regence.com/trgmedpol/drugs/dru368.pdf">http://blue.regence.com/trgmedpol/drugs/dru368.pdf</a> ) to <a href="https://regence.myprime.com/content/dam/prime/memberportal/forms/AuthorForms/Cambia/Program_Summaries/dru444reg.pdf">https://regence.myprime.com/content/dam/prime/memberportal/forms/AuthorForms/Cambia/Program_Summaries/dru444reg.pdf</a> .
11/28/2017	1. <b>Changed</b> "Quantities of up to 270 capsules per 30 days" to "267 mg strength: Quantities of up to 270 capsules/tablets per 30 days; 801 mg strength: Quantities of up to 90 tablets per 30 days" <b>under Quantity/Days Supply Restrictions.</b>