

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: 8/23/2018

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.
2. Medi-Span
3. https://regence.myprime.com/content/dam/prime/memberportal/forms/AuthorForms/Cambia/Program_Summaries/dru444reg.pdf.
4. https://ahca.myflorida.com/medicaid/Prescribed_Drug/drug_criteria_pdf/Esbriet_Criteria.pdf.

HISTORICAL TRACKING OF CHANGES MADE TO POLICY

Date	Notes/Changes
8/23/2018	1. Changed obsolete URL in Reference #3 (http://blue.regence.com/trgmedpol/drugs/dru368.pdf) to https://regence.myprime.com/content/dam/prime/memberportal/forms/AuthorForms/Cambia/Program_Summaries/dru444reg.pdf .
11/28/2017	1. Changed "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" under Quantity/Days Supply Restrictions .