#### **ERLOTINIB**

### **Pre - PA Allowance**

None

## **Prior-Approval Requirements**

Age 18 years of age or older

#### **Diagnoses**

The patient must have **ONE** of the following:

1. Non-small cell lung cancer (NSCLC)

#### AND the following:

- a. Metastatic disease with a positive EGFR mutation (exon 19 deletions OR exon 21 L858R substitution mutations) detected by an FDA-approved test (e.g. cobas® EGFR Mutation Test)
- 2. Pancreatic cancer
  - a. Tumor is locally advanced, unresectable or metastatic
  - b. First line treatment
  - c. Used in combination with gemcitabine
- 3. Renal cell carcinoma
  - a. Relapsed or unresectable Stage IV disease with non-clear cell histology
- 4. Recurrent Chordoma
- Leptomeningeal metastases from NSCLC
  - a. Positive EGFR mutation (exon 19 deletions or exon 21 L858R substitution mutations) detected by an FDA-approved test (e.g. cobas® EGFR Mutation Test)

## **Prior - Approval Limits**

#### Quantity

Strength	Quantity
25 mg	180 tablets per 90 days <b>OR</b>
100 mg	90 tablets per 90 days OR

Federal Employee Program.

#### **ERLOTINIB**

150 mg 90 tablets per 90 days

Maximum daily limit of any combination: 150 mg

**Duration** 12 months

# Prior – Approval Renewal Requirements

Age 18 years of age or older

#### **Diagnoses**

Patient must have **ONE** of the following:

- 1. Metastatic non-small cell lung cancer (NSCLC)
- 2. Pancreatic cancer
- 3. Renal cell carcinoma
- 4. Recurrent Chordoma
- 5. Leptomeningeal metastases from NSCLC

#### **AND ALL** of the following:

a. NO disease progression or unacceptable toxicity

# Prior - Approval Renewal Limits

Same as above