



Pre – PA Allowance

None

Prior - Approval Requirements

Age 18 years of age or older

Diagnoses

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

1. Advanced or metastatic gastric cancer or gastro-esophageal junction adenocarcinoma
 - a. Used as a single agent (monotherapy) or combination therapy with paclitaxel
 - b. Patient has received prior chemotherapy containing fluoropyrimidine or platinum and experienced disease progression on or after therapy
2. Metastatic non-small cell lung cancer (NSCLC) and **ONE** of the following:
 - a. Used in combination with docetaxel
 - i. Patient has received prior chemotherapy containing platinum and experienced disease progression on or after therapy
 - ii. Positive EGFR or ALK tumor expression
only: patient has had disease progression on FDA-approved therapy
 - b. Used in combination with erlotinib
 - i. First-line treatment
 - ii. Tumors have epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 (L858R) mutations
3. Metastatic colorectal cancer
 - a. Combination therapy with FOLFIRI
 - b. Patient has received prior chemotherapy containing bevacizumab, oxaliplatin, and a fluoropyrimidine and experienced disease progression on or after therapy
4. Hepatocellular carcinoma (HCC)



CYRAMZA
(ramucirumab)

- a. Used as a single agent (monotherapy)
- b. Alpha fetoprotein \geq (AFP) 400 ng/mL
- c. Patient has previously been treated with (sorafenib)

AND the following for **ALL** diagnoses:

1. Confirmation that patient does not have the following and if condition develops, therapy will be discontinued:
 - a. Hemorrhage or any severe bleeding event
 - b. Arterial thromboembolic events (ATEs)

Prior - Approval Limits

Duration 12 months

Prior – Approval *Renewal* Requirements

Age 18 years of age or older

Diagnoses

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

1. Advanced or metastatic gastric cancer or gastro-esophageal junction adenocarcinoma
2. Metastatic non-small cell lung cancer (NSCLC)
3. Metastatic colorectal cancer
4. Hepatocellular carcinoma (HCC)

AND ALL of the following:

1. Patient has not experienced disease progression or unacceptable toxicity
2. Confirmation that patient does not have the following and if condition develops, therapy will be discontinued:
 - a. Hemorrhage or any severe bleeding event
 - b. Arterial thromboembolic events (ATEs)

Prior – Approval *Renewal* Limits

Same as above