

**PEMAZYRE  
(pemigatinib)**

**Pre - PA Allowance**

None

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**Prior-Approval Requirements**

**Age** 18 years of age or older

**Diagnoses**

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

1. Unresectable locally advanced or metastatic cholangiocarcinoma
  - a. Fibroblast growth factor receptor 2 (FGFR2) fusion or other rearrangement, as detected by an FDA-approved test
  - b. Patient has had at least one prior therapy
2. Relapsed or refractory myeloid/lymphoid neoplasms (MLNs)
  - a. Fibroblast growth factor receptor 1 (FGFR1) rearrangement

**AND ALL** of the following:

- a. Baseline ophthalmological examination has been done and patient will be monitored for retinal pigment epithelial detachment (RPED)
- b. Prescriber agrees to monitor for hyperphosphatemia and agrees to initiate a low phosphate diet or phosphate lowering therapy, as clinically indicated
- c. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Pemazyre and for 1 week after the last dose
- d. Males with female partners of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Pemazyre and for 1 week after the last dose

**Prior - Approval Limits**

**Quantity**

Diagnosis	Quantity
Cholangiocarcinoma	56 tablets per 84 days <b>OR</b>
Myeloid/Lymphoid Neoplasms	84 tablets per 84 days

**Duration** 12 months

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**BlueCross  
BlueShield**

Federal Employee Program.

**PEMAZYRE  
(pemigatinib)**

**Prior – Approval *Renewal* Requirements**

**Age** 18 years of age or older

**Diagnosis**

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

1. Unresectable locally advanced or metastatic cholangiocarcinoma
2. Relapsed or refractory myeloid/lymphoid neoplasms (MLNs)

**AND ALL** of the following:

- a. **NO** disease progression or unacceptable toxicity
- b. Patient will be monitored for retinal pigment epithelial detachment (RPED)
- c. Prescriber agrees to monitor for hyperphosphatemia and agrees to initiate a low phosphate diet or phosphate lowering therapy, as clinically indicated
- d. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Pemazyre and for 1 week after the last dose
- e. Males with female partners of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Pemazyre and for 1 week after the last dose

**Prior - Approval *Renewal* Limits**

Same as above