

**OJEMDA  
(tovorafenib)**

**Pre - PA Allowance**

None

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**Prior-Approval Requirements**

**Age** 6 months of age or older

**Diagnosis**

Patient must have the following:

1. Relapsed or refractory pediatric low-grade glioma (LGG)
  - a. Patient has **ONE** of the following:
    - i. BRAF fusion or rearrangement
    - ii. BRAF V600 mutation

**AND ALL** of the following:

1. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Ojemda and for 28 days after the last dose
2. Males with female partners of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Ojemda and for 2 weeks after the last dose

**Prior - Approval Limits**

**Quantity**

Strength/Dosage Form	Quantity
100 mg tablet	72 tablets per 84 days <b>OR</b>
25 mg/mL oral suspension	24 bottles per 84 days

**Duration** 12 months

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**Prior – Approval *Renewal* Requirements**

**Age** 6 months of age or older

**Diagnosis**

Patient must have the following:

1. Relapsed or refractory pediatric low-grade glioma (LGG)



**BlueCross  
BlueShield**

Federal Employee Program.

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- a. **NO** disease progression or unacceptable toxicity

**AND ALL** of the following:

1. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Ojemda and for 28 days after the last dose
2. Males with female partners of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Ojemda and for 2 weeks after the last dose

**Prior - Approval *Renewal* Limits**

Same as above