

**KOSELUGO
(selumetinib)**

Pre - PA Allowance

None

Prior-Approval Requirements

Age 2 years of age or older

Diagnosis

Patient must have the following:

Neurofibromatosis Type 1 (NF1)

AND ALL of the following:

- a. Patient is symptomatic
- b. Patient has plexiform neurofibromas (PN) that are inoperable
- c. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Koselugo 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 Koselugo and for 1 week after the last dose
- e. Baseline ophthalmic assessment has been done and prescriber agrees to monitor for ocular toxicities
- f. Baseline left ventricular ejection fraction (LVEF) has been assessed and prescriber agrees to monitor LVEF

Prior - Approval Limits

Quantity 720 capsules per 90 days

Duration 12 months

Prior – Approval *Renewal* Requirements

Age 2 years of age or older

Diagnosis

Patient must have the following:

Neurofibromatosis Type 1 (NF1)



**BlueCross
BlueShield**

Federal Employee Program.

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AND ALL of the following:

- a. **NO** disease progression or unacceptable toxicity
- b. Females of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Koselugo and for 1 week after the last dose
- c. Males with female partners of reproductive potential **only**: patient will be advised to use effective contraception during treatment with Koselugo and for 1 week after the last dose
- d. Prescriber agrees to monitor for ocular toxicities
- e. Prescriber agrees to monitor left ventricular ejection fraction (LVEF)

Prior - Approval *Renewal* Limits

Same as above