

Immunomodulators: Olumiant

Member Information

1. Member Last Name: _____ 2. First Name: _____
3. Member ID #: _____ 4. Member Date of Birth: _____ 5. Member Gender: _____

Prescriber Information

6. Prescribing Provider NPI #: _____
7. Requester Contact Information - Name: _____ Phone #: _____ Ext. _____

Drug Information

8. Drug Name: _____ 9. Strength: _____ 10. Quantity Per 30 Days: _____
11. Length of Therapy (in days): ☐ up to 30 Days ☐ 60 Days ☐ 90 Days ☐ 120 Days ☐ 180 Days ☐ 365 Days ☐
Other _____

Clinical Information

Request for Rheumatoid Arthritis

1. Does the member have a diagnosis of Rheumatoid Arthritis? ☐ Yes ☐ No
2. Is the member not on another injectable biologic immunomodulator? ☐ Yes ☐ No
3. Has the member individual risks and benefits been considered prior to initiating or continuing therapy in those at higher risk for malignancy and/or major adverse cardiovascular events (MACE)? ☐ Yes ☐ No
4. Is the member NOT considered to be at high risk for thrombosis? ☐ Yes ☐ No
5. Has the member been considered and screened for the presence of latent tuberculosis? ☐ Yes ☐ No
6. Has the member been tested with Hep B SAG and Core Ab? ☐ Yes ☐ No
7. Will the member NOT receive live vaccines during therapy? ☐ Yes ☐ No
8. Has the member experienced a therapeutic failure/inadequate response, with at least one Tumor Necrosis Factor Blocker)? ☐ Yes ☐ No
9. Is the member unable to receive Tumor Necrosis Factor Blockers due to contraindications or intolerabilities? ☐ Yes ☐ No
10. Has the member had a trial and failure of Enbrel or Humira or a clinical reason member cannot try Enbrel or Humira? ☐ Yes ☐ No

Signature of Prescriber: _____ Date: _____

(Prescriber Signature Mandatory)

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.