

Kaiser Foundation Health Plan of Washington Kaiser Foundation Health Plan of Washington Options, Inc. CONTRACT MANAGER NAME Provider Communications, RCR-A3W-04 PO Box 34262, Seattle WA 98124-1262

FEBRUARY 28, 2024

GOLIMUMAB (SIMPONI ARIA) UPDATED PRIOR AUTHORIZATION CRITERIA

Dear Provider,

Golimumab (Simponi Aria) is on the **non-Medicare** list of office-administered drugs requiring prior authorization. **Effective June 1, 2024**, the criteria for golimumab (Simponi Aria) will expand to include a quantity limit for psoriatic arthritis (PsA) and ankylosing spondylitis (AS) indications. **This letter is a notification of the change in prior authorization criteria required before administering this medication under the medical benefit.**

Kaiser Foundation Health Plan of Washington and Kaiser Foundation Health Plan of Washington Options, Inc. (Kaiser Permanente) requires prior authorization for a select group of injectable drugs that may be administered under the medical benefit in a physician's office or by home infusion. These reviews are intended to ensure consistent benefit adjudication as well as appropriate utilization in accordance with the Kaiser Permanente Pharmacy & Therapeutics Committee's evidence-based criteria for coverage.

Prior Authorization Criteria for Golimumab (Simponi Aria) (changes are in bold):

DRUG NAME	COVERAGE CRITERIA
GOLIMUMAB	Covered for:
	 Patients with rheumatoid arthritis (RA) who have failure, contraindication, or intolerance to methotrexate, two formulary anti-TNFs (e.g., adalimumab [e.g., Amjevita], infliximab [e.g., Inflectra]), abatacept, and one other biologic DMARD
	2. Psoriatic arthritis (PsA) in patients with failure, contraindication, or intolerance to:
	 At least one conventional synthetic disease modifying anti-rheumatic drug (csDMARD) (methotrexate preferred), and Two of the following biologics (one of which must be adalimumab or infliximab) and adalimumab (e.g., Amjevita) infliximab (e.g., Inflectra) secukinumab etanercept Guselkumab, and at least one of the following biologic DMARDs (ustekinumab, risankizumab, abatacept)
	Note: csDMARD not required for patients with axial disease or severe (rapidly progressive, erosive) disease
	3. Patients with active ankylosing spondylitis (AS) who have failure, contraindication, or intolerance to two formulary anti-TNFs (e.g., adalimumab [Amjevita] or infliximab [Inflectra]), and secukinumab

DRUG NAME

COVERAGE CRITERIA

Not covered for use in combination with other biologic therapies including (but not limited to):

• Infliximab, adalimumab, etanercept, vedolizumab, rituximab, abatacept, tocilizumab, certolizumab, ustekinumab, canakinumab

Quantity Limit for RA/PsA/AS:

Induction: 2 mg/kg at weeks 0 and 4Maintenance: 2 mg/kg every 8 weeks

Note: Must be administered in a non-hospital setting. See <u>Site of Care: Infusion Therapy and Clinic Administered Medicines*</u> for criteria, reauthorization, and exceptions for new starts.

Members will have in-network benefit coverage for select home infused medications and supplies only when they get these medicines and supplies through Kaiser Permanente Specialty Home Infusion. There is no out-of-network benefit coverage for home infusion. See Infused Drugs Restricted to Kaiser Permanente Washington's Specialty Pharmacy Network** for medications impacted by this change.

https://wa-provider.kaiserpermanente.org/static/pdf/provider/clinical-review/infusion-site-care-policy.pdf
**Infused Drugs Restricted URL

https://healthy.kaiserpermanente.org/content/dam/kporg/final/documents/formularies/wa/infused-drugs-wa-en.pdf

Additional Information

A complete list of office-administered injectable drugs requiring prior authorization can be found on the Kaiser Permanente provider website at https://wa-provider.kaiserpermanente.org/provider-manual/clinical-review/officeinject.

To request prior authorization review, please use the Referral Request online form (login required) located on the Kaiser Permanente provider website at https://wa-provider.kaiserpermanente.org/provider-manual/clinical-review/preservice. You can also fax your request to the Review Services department toll-free at 1-888-282-2685.

Thank you for the care you provide to our members, your patients. If you have any questions about this process, please call Review Services at 1-800-289-1363, Monday through Friday from 8 a.m. to 5 p.m.

Sincerely,

Ravi Ubriani, MD, Chair

Pharmacy & Therapeutics Committee

^{*}Site of Care: Infusion Therapy and Clinic Administered Medicines URL