

**Kevzara® (sarilumab)
Effective 02/20/19**

Plan	<input checked="" type="checkbox"/> MassHealth <input type="checkbox"/> Commercial/Exchange	Program Type	<input checked="" type="checkbox"/> Prior Authorization <input type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
Benefit	<input checked="" type="checkbox"/> Pharmacy Benefit <input type="checkbox"/> Medical Benefit (NLX)		
Specialty Limitations	This medication has been designated specialty and must be filled at a contracted specialty pharmacy.		
Contact Information	Specialty Medications		
	All Plans	Phone: 866-814-5506	Fax: 866-249-6155
	Non-Specialty Medications		
	MassHealth	Phone: 877-433-7643	Fax: 866-255-7569
	Commercial	Phone: 800-294-5979	Fax: 888-836-0730
	Exchange	Phone: 855-582-2022	Fax: 855-245-2134
	Medical Specialty Medications (NLX)		
	All Plans	Phone: 844-345-2803	Fax: 844-851-0882
Exceptions	N/A		

Overview

Kevzara® (sarilumab) is an interleukin-6 (IL-6) receptor antagonist indicated for treatment of adult patients with moderately to severely active rheumatoid arthritis who have had an inadequate response or intolerance to one or more disease-modifying antirheumatic drugs (DMARDs).

Coverage Guidelines

Authorization may be granted when the following criteria are met:

- Member has a diagnosis of RA **AND**
 - Member is at least 18 years of age **AND**
 - Prescriber has provided documentation of ONE of the following:
 - a. Inadequate response, adverse reaction, or contraindication to at least ONE traditional DMARD (hydroxychloroquine, methotrexate, sulfasalazine)
 - b. Inadequate response or adverse reaction to ONE biologic DMARD that is FDA-approved for the requested indication
- AND**
- Dosing is appropriate (see appendix)

Continuation of Therapy

Reauthorization requires physician documentation indicating a positive response to therapy.

Limitations

1. Initial approvals will be for 6 months.
2. Reauthorizations will be for 12 months.

Appendix



Kevzara® (sarilumab)	Rheumatoid arthritis: 200 mg subcutaneously every two weeks, reduce dose to 150 mg for management of neutropenia, thrombocytopenia and elevated liver enzymes
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References

1. Kevzara (sarilumab) [prescribing information]. Bridgewater, NJ: Sanofi-Aventis; April 2018.
2. Genovese MC, Fleischmann R, Kivitz AJ, et al. Sarilumab plus methotrexate in patients with active rheumatoid arthritis and inadequate response to methotrexate: results of a phase III study. *Arthritis Rheumatol.* June 2015;67(6):1424-37.
3. Strand V, Reaney M, Chen C, et al. Sarilumab improves patient-reported outcomes in rheumatoid arthritis patients with inadequate response/intolerance to tumour necrosis factor inhibitors. *RMD Open.* 2017; 3:e000416. doi: 10.1136/rmdopen-2016-000416.

Review History

03/01/18 – Effective

02/20/19 – Reviewed in P&T Meeting

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