

Orencia (abatacept)
Effective 02/20/19

Plan	<input checked="" type="checkbox"/> MassHealth <input type="checkbox"/> Commercial/Exchange	Program Type	<input checked="" type="checkbox"/> Prior Authorization
Benefit	<input checked="" type="checkbox"/> Pharmacy Benefit <input type="checkbox"/> Medical Benefit (NLX)		<input type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
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

Orencia (abatacept) is a selective T cell costimulation modulator indicated for:

1. Moderately to severely active Rheumatoid Arthritis (RA) in adults
2. Moderately to severely active polyarticular Juvenile Idiopathic Arthritis (pJIA) in patients 2 years of age or older
3. Active Psoriatic Arthritis (PsA) in adults

Coverage Guidelines

Authorization may be granted when the following criteria are met:

1. Member has a diagnosis of RA, pJIA or PsA **AND**
 2. 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 or a contraindication to ALL biologic DMARDs
- AND**
3. Dosing is appropriate (see appendix for dosing)

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 up to 12 months.

Appendix

	Pediatric Dosing	Other Dosing												
Orencia® (abatacept)	<p>Juvenile Idiopathic Arthritis (≥ 6 years old, weight < 75 kg) 10 mg/kg IV at weeks 0, 2, and 4, then every 4 weeks thereafter</p> <p>Juvenile Idiopathic Arthritis (≥ 6 years old, weight > 75 kg) Refer to adult IV dosing regimen. Maximum dose: 1,000 mg.</p> <p>Juvenile Idiopathic Arthritis (≥ 2 years old, weight 10 kg to 24 kg) 50 mg SQ every week</p> <p>Juvenile Idiopathic Arthritis (≥ 2 years old, weight 25 to 49 kg) 87.5 mg SQ every week</p> <p>Juvenile Idiopathic Arthritis (≥ 2 years old, weight ≥50 kg) 125 mg SQ every week</p>	<p>Rheumatoid arthritis or Psoriatic Arthritis (IV regimen) Dose based on weight infused over 30 minutes at week 0, 2, and 4, then every 4 weeks thereafter.</p> <table border="1"> <thead> <tr> <th>Body weight of patient</th> <th>Dose</th> <th>Number of vials</th> </tr> </thead> <tbody> <tr> <td><60 kg</td> <td>500 mg</td> <td>2</td> </tr> <tr> <td>60 to 100 kg</td> <td>750 mg</td> <td>3</td> </tr> <tr> <td>>100 kg</td> <td>1 gm</td> <td>4</td> </tr> </tbody> </table> <p>Rheumatoid arthritis or Psoriatic Arthritis (SQ regimen) Weight-based IV loading dose (as per above), then 125 mg SQ within 1 day, followed by 125 mg SQ once weekly.</p> <p><u>Unable to receive an IV infusion:</u> Initiate weekly SQ injections without an IV loading dose.</p> <p><u>Transitioning from IV to SQ:</u> administer the first SQ dose instead of the next scheduled IV dose.</p>	Body weight of patient	Dose	Number of vials	<60 kg	500 mg	2	60 to 100 kg	750 mg	3	>100 kg	1 gm	4
Body weight of patient	Dose	Number of vials												
<60 kg	500 mg	2												
60 to 100 kg	750 mg	3												
>100 kg	1 gm	4												

References

- Orencia [package insert]. Princeton, NJ: Bristol-Myers Squibb; June 2017.
- Smolen JS, Landewé R, Billsma J, et al. EULAR recommendations for the management of rheumatoid arthritis with synthetic and biological disease-modifying antirheumatic drugs: 2016 update. *Ann Rheum Dis.* 2017;0:1-18.
- Mease P, Genovese MC, Gladstein G, et al. Abatacept in the treatment of patients with psoriatic arthritis: results of a six-month, multicenter, randomized, double-blind, placebo-controlled, phase II trial. *Arthritis Rheum* 2011; 63:939.
- Beukelman T, Patkar NM, Saag KG, et al. 2011 American College of Rheumatology recommendations for the treatment of juvenile idiopathic arthritis: initiation and safety monitoring of therapeutic agents for the treatment of arthritis and systemic features. *Arthritis Care Res.* 2011;63(4):465-482.
- Ruperto N, Lovell DJ, Quartier P, et al. Abatacept in children with juvenile idiopathic arthritis: a randomised, double-blind, placebo-controlled withdrawal trial. *Lancet* 2008; 372:383.

Review History

06/26/06 – Reviewed and revised
 08/15/06 – Effective
 02/25/08 – Reviewed
 02/23/09 – Reviewed
 02/22/10 – Reviewed
 02/28/11 – Reviewed
 10/24/11 – Reviewed and revised (09/12/11 drug file Orencia SQ)
 02/27/12 – Reviewed



02/25/13 – Reviewed

02/24/14 – Reviewed

02/23/15 – Reviewed

02/22/16 – Reviewed

02/27/17 – Reviewed and revised (adopted SGM & ST) in P&T Meeting

03/01/18 – Reviewed and revised (adopted MH RS)

02/20/19 – Reviewed in P&T Meeting

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