RAdvance

Clinical Policy Title:	apremilast
Policy Number:	RxA.742
Drug(s) Applied:	Otezla®
Original Policy Date:	04/18/2022
Last Review Date:	12/1/2023
Line of Business Policy Applies to:	All lines of business (except Medicare)

Criteria

Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria. The provision of provider samples does not guarantee coverage under the terms of the pharmacy benefit administered by RxAdvance. All criteria for initial approval must be met in order to obtain coverage.

I. Initial Approval Criteria

- A. Psoriatic Arthritis (must meet all):
 - 1. Diagnosis of PsA;
 - 2. Prescribed by or in consultation with a dermatologist or a rheumatologist.

Approval Duration All Lines of Business (except Medicare): 12 months

- B. Plaque Psoriasis (must meet all):
 - 1. Diagnosis of Plaque Psoriasis (PsO);
 - 2. Trial and failure of ≥ 3 months of at least <u>one</u> (1) conventional systemic therapy, unless contraindicated or clinically significant adverse effects are experienced:
 - a. Methotrexate [MTX]
 - b. Cyclosporine
 - c. Acitretin
 - d. Phototherapy (psoralen plus ultraviolet A light [PUVA]);
 - 3. Prescribed by or in consultation with a dermatologist or a rheumatologist;

Approval Duration

All Lines of Business (except Medicare): 12 months

- C. Behçet's Disease (must meet all):
 - 1. Diagnosis of oral ulcers in members with BD;
 - 2. Trial and failure of at least one (1) one systemic therapy (e.g. colchicine, corticosteroids, azathioprine) at maximally indicated doses unless contraindicated or significantly adverse effects are experienced;
 - 3. Prescribed by or in consultation with a dermatologist or a rheumatologist;

Approval duration

All Lines of Business (except Medicare): 12 months

II. Continued Therapy Approval

This clinical policy has been developed to authorize, modify, or determine coverage for individuals with similar conditions. Specific care and treatment may vary depending on individual need and benefits covered by the plan. This policy is not intended to dictate to providers how to practice medicine, nor does it constitute a contract or guarantee regarding payment or results. This document may contain prescription brand name drugs that are trademarks of pharmaceutical manufacturers that are not affiliated with RxAdvance.

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All Indications:

1. Member is currently receiving medication, excluding manufacturer samples.

Approval Duration

All Lines of Business (except Medicare): 12 months

References

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- Menter A, Korman, NJ, Elmets CA, et al. American Academy of Dermatology. Guidelines of care for the management of psoriasis and psoriatic arthritis. Section 3. Guidelines of care for the management and treatment of psoriasis with topical therapies. J Am Acad Dermatol. 2009; 60:643-659. Available at: https://pubmed.ncbi.nlm.nih.gov/19217694/. Accessed March 27, 2023
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- Hatemi G, Mahr A, Takeno M, et al. Improvements and correlations in oral ulcers, disease activity, and QOL in behçet's syndrome patients treated with apremilast: a phase 3 randomized, double-blind, placebo-controlled study. Rheumatology, Volume 58, Issue Supplement_2) Available at: <u>https://doi.org/10.1093/rheumatology/kez062.023</u>. Accessed March 27, 2023
- Hatemi G, Christensen R, Bang D, et al. 2018 update of the EULAR recommendations for the management of Behçet's syndrome Annals of the Rheumatic Diseases 2018;77:808-818. Available at: <u>https://ard.bmj.com/content/annrheumdis/early/2018/04/06/annrheumdis-2018-213225.full.pdf</u>. Accessed March 27, 2023

Review/Revision History	Review/Revision Date	P&T Approval Date
RxA.592.Biologic_DMARDs was last reviewed and updated on 01/05/2022 and archived on 04/18/2022. For details, please refer to RxA.592.Biologic_DMARDs.	01/05/2022	4/18/2022
Drug specific policy for Otezla was created	2/14/2022	4/18/2022

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based o	on RxA.592.Biologic_DMARDs		
1.			
	Updated to include renal impairment		
	dosing information for indication PsO,		
	PsA, BD.		
2.	Initial Approval Criteria I.B.3: Updated		
	trial and failure criteria to rephrase and		
	include phototherapy (psoralen plus		
	ultraviolet A light [PUVA]).		
3.	Appendix A: Updated to include		
	abbreviations PUVA.		
4.	Appendix B, Drug Name: Updated to		
	remove discontinued brand-name		
-	therapeutic alternative Soriatane [®] .		
э.	Appendix B, Drug Name: Updated to include brand-name therapeutic		
	alternative of other biological DMARDs.		
6.	Disclaimer about contraindications		
0.	"Contraindications listed reflect		
	statements made in the manufacturer's		
	package insert" was added to		
	Appendix C.		
7.	Appendix D, General Information:		
	Updated to include new information		
	regarding Warnings and Precautions.		
8.	References were reviewed and		
	updated.		
	vas reviewed:	03/27/2023	04/13/2023
1.	References were reviewed and		
	updated.		
Policy v	vas reviewed.	12/1/2023	12/1/2023
Policy v	vas reviewed.	3/1/2024	3/1/2024