

PHARMACY / MEDICAL POLICY – 5.01.645 Pharmacologic Treatment of Psoriatic Arthritis

Effective Date:	Oct. 3, 2025*	RELATED MEDICAL POLICIES:
Last Revised:	Jun. 10, 2025	5.01.550 Pharmacotherapy of Arthropathies
Replaces:	N/A	5.01.563 Pharmacotherapy of Inflammatory Bowel Disorder
		5.01.564 Pharmacotherapy of Miscellaneous Autoimmune Diseases
*This policy has been revised. Click		5.01.607 Continuity of Coverage for Maintenance Medications
here to view the current policy.		5.01.628 Pharmacologic Treatment of Atopic Dermatitis
		5.01.629 Pharmacologic Treatment of Psoriasis
		5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies
		11.01.523 Site of Service: Infusion Drugs and Biologic Agents

The Site of Service Medical Necessity criteria within this policy DOES NOT apply to Alaska fully-insured members; refer to the infusion and injection drug Medical Necessity criteria only.

Site of Service *and* the infusion and injection drug Medical Necessity criteria apply to all other plan members.

Please contact Customer Service for more information.

Select a hyperlink below to be directed to that section.

POLICY CRITERIA | DOCUMENTATION REQUIREMENTS | CODING RELATED INFORMATION | EVIDENCE REVIEW | REFERENCES | HISTORY

Clicking this icon returns you to the hyperlinks menu above.

Introduction

Psoriatic arthritis is an inflammatory disease of the joints and areas where tendons and ligaments connect to bone. The most common symptoms are joint pain and stiffness, skin rashes, and changes in your fingernails and toenails. This policy discusses when biologics and other drugs are considered medically necessary for the treatment of psoriatic arthritis.

Note: The Introduction section is for your general knowledge and is not to be taken as policy coverage criteria. The rest of the policy uses specific words and concepts familiar to medical professionals. It is intended for providers. A provider can be a person, such as a doctor, nurse, psychologist, or dentist. A provider also can be a place where medical care is given, like a hospital, clinic, or lab. This policy informs them about when a service may be covered.

Policy Coverage Criteria

Site of Service (SOS) Medical Necessity criteria applies ONLY to medical benefit reviews. SOS Medical Necessity criteria does NOT apply to Alaska fully-insured members; refer to the infusion drug Medical Necessity criteria only. Please contact Customer Service for more information.

We will review specific intravenous (IV) and injectable drugs for medical necessity for all ages.

For those age 13 and older, we also will review the site of service for medical necessity. Site of service is defined as the location where the drug is administered, such as a hospital-based outpatient setting, an infusion center, a physician's office, or at home. Click **here** to be directed to the site of service review criteria.

Drugs subject to site of service review addressed in this policy are:

- Avsola (infliximab-axxq) IV
- Cosentyx (secukinumab) IV
- Inflectra (infliximab-dyyb) IV
- Infliximab (Janssen unbranded) IV
- Orencia (abatacept) IV
- Remicade (infliximab) IV
- Renflexis (infliximab-abda) IV
- Simponi Aria (golimumab) IV

Site of Service	Medical Necessity
Administration	
Medically necessary sites of service • Physician's office • Infusion center • Home infusion	 IV infusion and injection therapy of various medical or biologic agents will be covered in the most appropriate, safe and costeffective site: These are the preferred medically necessary sites of service for specified drugs.
Hospital-based outpatient setting Outpatient hospital IV infusion department	IV infusion and injection therapy of various medical or biologic agents will be covered in the most appropriate, safe and cost-effective site.
Hospital-based outpatient clinical level of care	 This site is considered medically necessary for the first 90 days for the following: The initial course of infusion or injection of a pharmacologic or biologic agent OR Re-initiation of an agent after 6 months or longer following discontinuation of therapy* *Note: This does not include when standard dosing between infusions or injections is 6 months or longer
	This site is considered medically necessary when there is no outpatient infusion center within 50 miles of the individual's home and there is no contracted home infusion agency that will travel to their home, or a hospital is the only place that offers infusions or injections of this drug. This site is considered medically necessary only when the individual has a clinical condition which puts him or her at increased risk of complications for infusions or injections, including any 1 of the following: • Known cardiac condition (e.g., symptomatic cardiac arrhythmia) or pulmonary condition (e.g., significant respiratory disease, serious obstructive airway disease, %FVC less than or equal to 40%) that may increase the risk of an adverse reaction



Site of Service	Medical Necessity
Administration	
	 Unstable renal function which decreases the ability to respond to fluids Difficult or unstable vascular access Acute mental status changes or cognitive conditions that impact the safety of infusion or injection therapy A known history of severe adverse drug reactions and/or anaphylaxis to prior treatment with a related or similar drug
	This site is considered medically necessary when the individual has cytokine release syndrome (CRS) and all the following are met:
	 CRS is grade 3 or 4 as evidenced by ALL the following: Temperature at least 38 °C
	 Hypotension that requires 1 or more vasopressors Hypoxia that requires oxygen through a high-flow nasal cannula, face mask, non-rebreather mask, or Venturi mask OR positive pressure (continuous positive airway pressure [CPAP], bilevel positive airway pressure [BiPAP], intubation, or mechanical ventilation)
	AND
	The individual will be admitted into an inpatient setting as soon as possible
Hospital-based outpatient	These sites are considered not medically necessary for infusion
setting	and injectable therapy services of various medical and biologic
Outpatient hospital IV	agents when the site-of-service criteria in this policy are not
infusion department	met.
 Hospital-based outpatient clinical level of care 	

This policy contains separate criteria to be used based on the member's formulary. Please check the member Plan booklet or member ID card for coverage and click the links below to navigate to the appropriate section:

Section 1: Open, Preferred, and Select Formulary Plans (Rx Plan A1, A2, B3, B4, C4, F1, and G3) and Plans with No Pharmacy Benefit Coverage



Section 3: Individual/Small Group/Student ISHIP Metallic Formulary Plans (Rx Plan M1, M2, and M4)

The following section applies to Open, Preferred, and Select formulary plans (Rx Plan A1, A2, B3, B4, C4, F1, and G3) and plans with no pharmacy benefit coverage only. Please refer to the member plan booklet or member ID card.

Step therapy tiers are listed below; please refer to the Policy section for details.

Section 1: Open, Preferred, and Select Formulary Plans (Rx Plan A1, A2, B3, B4, C4, F1, and G3) and Plans with No Pharmacy Benefit Coverage ONLY

	Arthritis –	

Psoriatic Arthritis – First Line					
TNF-α	IL-17	IL-12/23	IL-23	Janus	PDE-4
Inhibitors	Inhibitor	Inhibitors	Inhibitors	Kinase	Inhibitor
				Inhibitors	
 Inflectra (IV) Infliximab (Janssen – unbranded) (IV) Remicade (IV) Simponi Aria (IV) Adalimumabadaz (Hyrimoz unbranded) (SC) Adalimumabadbm (Cyltezo unbranded) (SC) Adalimumabaryvk (Simlandi unbranded) (SC) Cyltezo (SC) Enbrel (SC) Simlandi (SC) 	• Taltz (SC)	 Stelara (SC) Steqeyma (SC) Yesintek (SC) 	 Skyrizi (SC) Tremfya (SC) [pen, syringe, and injector] Skyrizi (SC) 	 Rinvoq / Rinvoq LQ (oral) Xeljanz / Xeljanz XR (oral) 	Otezla (oral)

Section 1: Open, Preferred, and Select Formulary Plans (Rx Plan A1, A2, B3, B4, C4, F1, and G3) and Plans with No Pharmacy Benefit Coverage ONLY

Psoriatic Arthritis – Second Line				
TNF-α Inhibitors	IL-17 IL-12/23		T-Cell Costimulation	
	Inhibitors	Inhibitors	Modulator	
Avsola (IV)	Bimzelx (SC)	Imuldosa (SC)	Orencia (IV/SC)	
Renflexis (IV)		Otulfi (SC)		
Cimzia (SC)	Cosentyx(IV/SC)	Pyzchiva (SC)		
Simponi(SC)		Selarsdi (SC)		
Abrilada (SC)	-	Brand		
Adalimumab-aacf (Idacio		ustekinumab (Stelara		
unbranded) (SC)		unbranded)		
Adalimumab-aaty (Yuflyma unbranded) (SC)		(SC)		
Adalimumab-fkjp (Hulio unbranded) (SC)		 Brand ustekinumab- aekn (Selarsdi 		
Amjevita (SC)		unbranded)		
Hadlima (SC)		(SC)		
Hulio (SC)		 Brand ustekinumab- 		
Humira (SC)		ttwe (Pyzchiva		
Hyrimoz (SC)		unbranded)		
Idacio (SC)		(SC)		
Yuflyma (SC)		Wezlana (SC)		
Yusimry (SC)				

	TINF-G Alitagonists - First Line			
Agent		Medical Necessity, Psoriatic Arthritis		
•	Cyltezo (adalimumab- adbm) SC Simlandi (adalimumab-	Simponi Aria (golimumab) IV is subject to review for site of service administration.		
	ryvk) SC	Cyltezo (adalimumab-adbm), Simlandi (adalimumab-ryvk), adalimumab-adaz (Hyrimoz unbranded), adalimumab-adbm		



- Adalimumab-adaz (Hyrimoz unbranded) SC
- Adalimumab-adbm (Cyltezo unbranded) SC
- Adalimumab-ryvk (Simlandi unbranded) SC
- Enbrel (etanercept) SC
- Simponi Aria (golimumab) IV

(Cyltezo unbranded), adalimumab-ryvk (Simlandi unbranded), Enbrel (etanercept) or Simponi Aria (golimumab) may be considered medically necessary for the treatment of active psoriatic arthritis when:

The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies. Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

- Inflectra (infliximabdyyb) IV
- Infliximab (Janssen unbranded) IV
- Remicade (infliximab) IV

Inflectra (infliximab-dyyb), Infliximab (Janssen – unbranded), and Remicade (infliximab) are subject to review for site of service administration.

Inflectra (infliximab-dyyb), Infliximab (Janssen – unbranded), and Remicade (infliximab) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

IL-17 Inhibitor - First Line

Taltz (ixekizumab) SC

Taltz (ixekizumab) SC may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

IL-12/23 Inhibitor - First Line

- Stelara (ustekinumab) SC
- Steqeyma (ustekinumabstba) SC
- Yesintek (ustekinumabkfce) SC

Steqeyma (ustekinumab-stba) SC and Yesintek (ustekinumab-kfce) SC may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 6 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

Stelara (ustekinumab) SC may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 6 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

AND

- Has had an inadequate response or intolerance to ALL the following agents:¹
 - Stegeyma (ustekinumab-stba) SC
 - Yesintek (ustekinumab-kfce) SC

¹**Note:** Only applies to individuals not previously treated with requested therapy.

IL-23 Inhibitors - First Line

Tremfya (guselkumab) SC (pen, syringe, and injector)

Tremfya (guselkumab) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

Skyrizi (risankizumab-rzaa) SC

Skyrizi (risankizumab-rzaa) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older



 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

Janus Kinase Inhibitors - First Line

- Rinvoq (upadacitinib) oral
- Rinvoq LQ (upadacitinib) oral

Rinvoq (upadacitinib) and Rinvoq LQ (upadacitinib) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 2 years or older

AND

 The individual has had an inadequate response or intolerance to 1 or more TNF blockers

AND

- Medication is being prescribed by or in consultation with a rheumatologist or dermatologist
- Xeljanz (tofacitinib) oral
- Xeljanz XR (tofacitinib extended-release) oral

Xeljanz (tofacitinib) and Xeljanz XR (tofacitinib extendedrelease) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Has had an inadequate response or intolerance to 1 or more TNF blockers

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

PDE4 Inhibitor - First Line

Otezla (apremilast) oral

Otezla (apremilast) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

TNF-α Antagonists – Second Line

- Abrilada (adalimumabafzb) SC
- Adalimumab-aacf (Idacio unbranded) SC

Abrilada (adalimumab-afzb), adalimumab-aacf (Idacio unbranded), adalimumab-aaty (Yuflyma unbranded), adalimumab-fkjp (Hulio unbranded), Amjevita (adalimumab-atto), Hadlima (adalimumab-bwwd), Hulio (adalimumab-fkjp),



- Adalimumab-aaty (Yuflyma unbranded) SC
- Adalimumab-fkjp (Hulio unbranded) SC
- Amjevita (adalimumabatto) SC
- Hadlima (adalimumabbwwd) SC
- Hulio (adalimumab-fkjp)
 SC
- Humira (adalimumab) SC
- Hyrimoz (adalimumabadaz) SC
- Idacio (adalimumab-aacf)
 SC
- Yuflyma (adalimumabaaty) SC
- Yusimry (adalimumabaqvh) SC

Humira (adalimumab), Hyrimoz (adalimumab-adaz), Idacio (adalimumab-aacf), Yuflyma (adalimumab-aaty), and Yusimry (adalimumab-aqvh) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

AND

- Has had an inadequate response or intolerance to ALL the following agents:
 - Cyltezo (adalimumab-adbm) OR adalimumab-adbm (Cyltezo unbranded)
 - Adalimumab-adaz (Hyrimoz unbranded)
 - Simlandi (adalimumab-ryvk) OR adalimumab-ryvk (Simlandi unbranded)

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies.

Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

- Cimzia (certolizumab pegol) SC
- Simponi (golimumab) SC

Cimzia (certolizumab pegol) and Simponi (golimumab) SC may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

- Has had an inadequate response or intolerance to 2 of the following agents:
 - Enbrel (etanercept)



- Cyltezo (adalimumab-adbm) OR Simlandi (adalimumabryvk) OR adalimumab-adaz (Hyrimoz unbranded) OR adalimumab-adbm (Cyltezo unbranded) OR adalimumabryvk (Simlandi unbranded)
- Otezla (apremilast)
- Rinvoq (upadacitinib) or Rinvoq LQ (upadacitinib)
- Skyrizi (risankizumab-rzaa) SC
- Stelara (ustekinumab) SC OR Steqeyma (ustekinumab-stba)
 SC OR Yesintek (ustekinumab-kfce) SC
- Taltz (ixekizumab)
- Tremfya (guselkumab)
- Xeljanz (tofacitinib) OR Xeljanz XR (tofacitinib extendedrelease)

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies.

Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

- Avsola (infliximab-axxq)
- Renflexis (infliximababda) IV

Avsola (infliximab-axxq) and Renflexis (infliximab-abda) are subject to review for site of service administration.

Avsola (infliximab-axxq) and Renflexis (infliximab-abda) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

AND

 Has had a documented trial and treatment failure with Inflectra (infliximab-dyyb), Infliximab (Janssen – unbranded), or Remicade (infliximab)

IL-12/23 Inhibitors - Second Line

- Imuldosa (ustekinumabsrlf) SC
- Otulfi (ustekinumab-aauz) SC
- Pyzchiva (ustekinumabttwe) SC
- Selarsdi (ustekinumabackn) SC
- Ustekinumab (Stelara unbranded) SC
- Ustekinumab-aekn (Selarsdi unbranded) SC
- Ustekinumab-ttwe (Pyzchiva unbranded) SC
- Wezlana (ustekinumabauub) SC

Imuldosa (ustekinumab-srlf) SC, Otulfi (ustekinumab-aauz) SC, Pyzchiva (ustekinumab-ttwe) SC, Selarsdi (ustekinumab-ackn) SC, ustekinumab (Stelara unbranded) SC, ustekinumab-aekn (Selarsdi unbranded) SC, ustekinumab-ttwe (Pyzchiva unbranded) SC, and Wezlana (ustekinumab-auub) SC may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 6 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

AND

- Has had an inadequate response or intolerance to ALL the following agents:
 - Stelara (ustekinumab) SC
 - Stegeyma (ustekinumab-stba) SC
 - Yesintek (ustekinumab-kfce) SC

IL-17 Inhibitors - Second Line

Bimzelx (bimekizumabbkzx) SC

Bimzelx (bimekizumab-bkzx) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

- Has had an inadequate response or intolerance to 1 of the following agents:
 - Enbrel (etanercept)
 - Cyltezo (adalimumab-adbm) OR Simlandi (adalimumabryvk) OR adalimumab-adaz (Hyrimoz unbranded) OR adalimumab-adbm (Cyltezo unbranded) OR adalimumabryvk (Simlandi unbranded)
 - Otezla (apremilast)
 - Skyrizi (risankizumab-rzaa)
 - Stelara (ustekinumab) OR Steqeyma (ustekinumab-stba) SC
 OR Yesintek (ustekinumab-kfce) SC
 - Taltz (ixekizumab)
 - Tremfya (guselkumab)



AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies. Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

Cosentyx (secukinumab) IV/SC

Cosentyx (secukinumab) IV is subject to review for site of service administration.

Cosentyx (secukinumab) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 2 years or older

AND

- Has had an inadequate response or intolerance to 2 of the following agents:
 - Enbrel (etanercept)
 - Cyltezo (adalimumab-adbm) OR Simlandi (adalimumabryvk) OR adalimumab-adaz (Hyrimoz unbranded) OR adalimumab-adbm (Cyltezo unbranded) OR adalimumabryvk (Simlandi unbranded)
 - Otezla (apremilast)
 - Rinvoq (upadacitinib) OR Rinvoq LQ (upadacitinib)
 - Skyrizi (risankizumab-rzaa)
 - Stelara (ustekinumab) OR Steqeyma (ustekinumab-stba) SC
 OR Yesintek (ustekinumab-kfce) SC
 - Taltz (ixekizumab)
 - Tremfya (guselkumab)
 - Xeljanz (tofacitinib) OR Xeljanz XR (tofacitinib extendedrelease)



 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies.

Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

T-Cell Costimulation Modulators – Second Line

Orencia (abatacept) IV/SC

Orencia (abatacept) IV is subject to review for site of service administration.

Orencia (abatacept) IV/SC may be considered medically necessary for the treatment of adults with active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

- Has had an inadequate response or intolerance to 2 of the following agents:
 - Enbrel (etanercept)
 - Cyltezo (adalimumab-adbm) OR Simlandi (adalimumabryvk) OR adalimumab-adaz (Hyrimoz unbranded) OR adalimumab-adbm (Cyltezo unbranded) OR adalimumabryvk (Simlandi unbranded)
 - Otezla (apremilast)
 - Rinvog (upadacitinib) **OR** Rinvog LQ (upadacitinib)
 - Skyrizi (risankizumab-rzaa)
 - Stelara (ustekinumab) SC OR Steqeyma (ustekinumab-stba)
 SC OR Yesintek (ustekinumab-kfce) SC
 - Taltz (ixekizumab)



- Tremfya (guselkumab)
- Xeljanz (tofacitinib) OR Xeljanz XR (tofacitinib extendedrelease)

OR

 Has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, or a demyelinating disorder

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies.

Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

Orencia (abatacept) SC may be considered medically necessary for the treatment of pediatric individuals with active psoriatic arthritis when:

• The individual is aged 2 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

AND

- Has had an inadequate response or intolerance to ONE of the following agents:
 - Enbrel (etanercept)
 - Rinvoq (upadacitinib) **OR** Rinvoq LQ (upadacitinib)
 - Stelara (ustekinumab) SC OR Steqeyma (ustekinumab-stba)
 SC OR Yesintek (ustekinumab-kfce) SC

OR

 Has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, or a demyelinating disorder



Step therapy tiers are listed below; please refer to the Policy section for details.

Section 2: Essentials Formulary Plans (Rx Plan E1, E3, and E4) ONLY					
Psoriatic Arthritis –	First Line				
TNF-α Inhibitors	IL-17	IL-12/23	IL-23	Janus	PDE-4
	Inhibitor	Inhibitors	Inhibitors	Kinase	Inhibitor
				Inhibitors	
 Inflectra (IV) Infliximab (Janssen – unbranded) (IV) Remicade (IV) Simponi Aria (IV) Adalimumab-adaz (Hyrimoz unbranded) (SC) Adalimumab-adbm (Cyltezo unbranded) (SC) Adalimumab-ryvk (Simlandi unbranded) (SC) Cyltezo (SC) 	• Taltz (SC)	 Steqeyma (SC) Yesintek (SC) 	 Skyrizi (SC) Tremfya (SC) [pen, syringe, and injector] Skyrizi (SC) 	 Rinvoq / Rinvoq LQ (oral) Xeljanz / Xeljanz XR (oral) 	• Otezla (oral)
Enbrel (SC)					
• Simlandi (SC)					
Psoriatic Arthritis – Second Line					
TNF-α Inhibitors		IL-17	IL-12/23	T-Cell Costin	nulation
		Inhibitors	Inhibitors	Modulator	
Avsola (IV)		Bimzelx	Imuldosa (SC)	Orencia (IV/S)	iC)
Renflexis (IV)		(SC)	Otulfi (SC)		
Cimzia (SC)		Cosentyx (IV/SC)	Pyzchiva (SC)		
Simponi(SC)		, , ,	 Selarsdi (SC) 		

- Abrilada (SC)
- Adalimumab-aacf (Idacio unbranded) (SC)
- Adalimumab-aaty (Yuflyma unbranded) (SC)
- Adalimumab-fkjp (Hulio unbranded) (SC)
- Amjevita (SC)
- Hadlima (SC)
- Hulio (SC)
- Humira (SC)
- Hyrimoz (SC)
- Idacio (SC)
- Yuflyma (SC)
- Yusimry (SC)

- Stelara (SC)
- Brand
 ustekinumab
 (Stelara
 unbranded)
 (SC)
- Brand ustekinumabaekn (Selarsdi unbranded) (SC)
- Brand ustekinumabttwe (Pyzchiva unbranded) (SC)
- Wezlana (SC)

Section 2: Essentials Formulary Plans (Rx Plan E1, E3, and E4) ONLY TNF- α Antagonists – First Line

Agent

- Cyltezo (adalimumabadbm) SC
- Simlandi (adalimumabryvk) SC
- Adalimumab-adaz (Hyrimoz unbranded) SC
- Adalimumab-adbm (Cyltezo unbranded) SC
- Adalimumab-ryvk (Simlandi unbranded) SC
- Enbrel (etanercept) SC
- Simponi Aria
 (golimumab) IV

Medical Necessity, Psoriatic Arthritis

Simponi Aria (golimumab) IV is subject to review for site of service administration.

Cyltezo (adalimumab-adbm), Simlandi (adalimumab-ryvk), adalimumab-adaz (Hyrimoz unbranded), adalimumab-adbm (Cyltezo unbranded), adalimumab-ryvk (Simlandi unbranded), Enbrel (etanercept) or Simponi Aria (golimumab) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist



Section 2: Essentials Formulary Plans (Rx Plan E1, E3, and E4) ONLY Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical **Necessity Criteria for Custom Open and Preferred Formularies.** Please check the member Plan booklet or member ID card to determine whether this policy criteria applies. Inflectra (infliximab-Inflectra (infliximab-dyyb), Infliximab (Janssen – unbranded), dyyb) IV and Remicade (infliximab) are subject to review for site of Infliximab (Janssen service administration. unbranded) IV Remicade (infliximab) IV Inflectra (infliximab-dyyb), Infliximab (Janssen – unbranded), and Remicade (infliximab) may be considered medically necessary for the treatment of active psoriatic arthritis when: The individual is aged 18 years or older **AND** Medication is being prescribed by or in consultation with a rheumatologist or dermatologist IL-17 Inhibitor - First Line Taltz (ixekizumab) SC Taltz (ixekizumab) SC may be considered medically necessary for the treatment of active psoriatic arthritis when: The individual is aged 18 years or older **AND** Medication is being prescribed by or in consultation with a rheumatologist or dermatologist IL-12/23 Inhibitor - First Line Steqeyma (ustekinumab-Stegeyma (ustekinumab-stba) SC and Yesintek (ustekinumabstba) SC kfce) SC may be considered medically necessary for the Yesintek (ustekinumabtreatment of active psoriatic arthritis when: kfce) SC The individual is aged 6 years or older **AND** Medication is being prescribed by or in consultation with a rheumatologist or dermatologist **IL-23 Inhibitors - First Line** Tremfya (guselkumab) SC Tremfya (guselkumab) may be considered medically necessary for the treatment of active psoriatic arthritis when: (pen, syringe, and injector) The individual is aged 18 years or older

Section 2: Essentials Form	ulary Plans (Rx Plan E1, E3, and E4) ONLY		
	AND		
	Medication is being prescribed by or in consultation with a		
	dermatologist or a rheumatologist		
Skyrizi (risankizumab-rzaa)	Skyrizi (risankizumab-rzaa) may be considered medically		
SC	necessary for the treatment of active psoriatic arthritis when:		
	The individual is aged 18 years or older		
	AND		
	Medication is being prescribed by or in consultation with a		
	dermatologist or a rheumatologist		
Janus Kinase Inhibitors - I	irst Line		
• Rinvoq (upadacitinib) oral	Rinvoq (upadacitinib) and Rinvoq LQ (upadacitinib) may be		
 Rinvoq LQ (upadacitinib) 	considered medically necessary for the treatment of active		
oral	psoriatic arthritis when:		
	The individual is aged 2 years or older		
	AND		
	The individual has had an inadequate response or intolerance		
	to 1 or more TNF blockers		
	AND		
	Medication is being prescribed by or in consultation with a		
	rheumatologist or dermatologist		
Xeljanz (tofacitinib) oral Xelianz VD (tofacitinib)	Xeljanz (tofacitinib) and Xeljanz XR (tofacitinib extended-		
 Xeljanz XR (tofacitinib extended-release) oral 	release) may be considered medically necessary for the		
extended-release) oral	treatment of active psoriatic arthritis when:		
	The individual is aged 18 years or older		
	AND		
	Has had an inadequate response or intolerance to 1 or more TNF blockers		
	AND		
	Medication is being prescribed by or in consultation with a should be a second by the second b		
DDE4 Inhibitor First Line	rheumatologist or dermatologist		
PDE4 Inhibitor - First Line	Otople (sprewilest) may be sensidered readically reserves for		
Otezla (apremilast) oral	Otezla (apremilast) may be considered medically necessary for		
	the treatment of active psoriatic arthritis when:		
	The individual is aged 18 years or older		
	AND		

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

TNF-α Antagonists – Second Line

- Abrilada (adalimumabafzb) SC
- Adalimumab-aacf (Idacio unbranded) SC
- Adalimumab-aaty (Yuflyma unbranded) SC
- Adalimumab-fkjp (Hulio unbranded) SC
- Amjevita (adalimumabatto) SC
- Hadlima (adalimumabbwwd) SC
- Hulio (adalimumab-fkjp)
 SC
- Humira (adalimumab) SC
- Hyrimoz (adalimumabadaz) SC
- Idacio (adalimumab-aacf)
 SC
- Yuflyma (adalimumabaaty) SC
- Yusimry (adalimumabagvh) SC

Abrilada (adalimumab-afzb), adalimumab-aacf (Idacio unbranded), adalimumab-aaty (Yuflyma unbranded), adalimumab-fkjp (Hulio unbranded), Amjevita (adalimumab-atto), Hadlima (adalimumab-bwwd), Hulio (adalimumab-fkjp), Humira (adalimumab), Hyrimoz (adalimumab-adaz), Idacio (adalimumab-aacf), Yuflyma (adalimumab-aaty), and Yusimry (adalimumab-aqvh) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

AND

- Has had an inadequate response or intolerance to ALL the following agents:
 - Cyltezo (adalimumab-adbm) OR adalimumab-adbm (Cyltezo unbranded)
 - Adalimumab-adaz (Hyrimoz unbranded)
 - Simlandi (adalimumab-ryvk) OR adalimumab-ryvk (Simlandi unbranded)

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies. Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

- Cimzia (certolizumab pegol) SC
- Simponi (golimumab) SC

Cimzia (certolizumab pegol) and Simponi (golimumab) SC may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

AND

- Has had an inadequate response or intolerance to 2 of the following agents:
 - Enbrel (etanercept)
 - Cyltezo (adalimumab-adbm) OR Simlandi (adalimumabryvk) OR adalimumab-adaz (Hyrimoz unbranded) OR adalimumab-adbm (Cyltezo unbranded) OR adalimumabryvk (Simlandi unbranded)
 - Otezla (apremilast)
 - Rinvoq (upadacitinib) or Rinvoq LQ (upadacitinib)
 - Skyrizi (risankizumab-rzaa) SC
 - Steqeyma (ustekinumab-stba) SC OR Yesintek (ustekinumab-kfce) SC
 - Taltz (ixekizumab)
 - Tremfya (guselkumab)
 - Xeljanz (tofacitinib) OR Xeljanz XR (tofacitinib extendedrelease)

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies.

Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

- Avsola (infliximab-axxq)
 IV
- Renflexis (infliximababda) IV

Avsola (infliximab-axxq) and Renflexis (infliximab-abda) are subject to review for site of service administration.

Avsola (infliximab-axxq) and Renflexis (infliximab-abda) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist



AND

 Has had a documented trial and treatment failure with Inflectra (infliximab-dyyb), Infliximab (Janssen – unbranded), or Remicade (infliximab)

IL-12/23 Inhibitors - Second Line

- Imuldosa (ustekinumabsrlf) SC
- Otulfi (ustekinumab-aauz)
 SC
- Pyzchiva (ustekinumabttwe) SC
- Selarsdi (ustekinumabackn) SC
- Stelara (ustekinumab) SC
- Ustekinumab (Stelara unbranded) SC
- Ustekinumab-aekn (Selarsdi unbranded) SC
- Ustekinumab-ttwe (Pyzchiva unbranded) SC
- Wezlana (ustekinumabauub) SC

Imuldosa (ustekinumab-srlf) SC, Otulfi (ustekinumab-aauz) SC, Pyzchiva (ustekinumab-ttwe) SC, Selarsdi (ustekinumab-ackn) SC, Stelara (ustekinumab), ustekinumab (Stelara unbranded) SC, ustekinumab-aekn (Selarsdi unbranded) SC, ustekinumab-ttwe (Pyzchiva unbranded) SC, and Wezlana (ustekinumab-auub) SC may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 6 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

AND

- Has had an inadequate response or intolerance to ALL the following agents:
 - Steqeyma (ustekinumab-stba) SC
 - Yesintek (ustekinumab-kfce) SC

IL-17 Inhibitors - Second Line

Bimzelx (bimekizumabbkzx) SC

Bimzelx (bimekizumab-bkzx) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

- Has had an inadequate response or intolerance to 1 of the following agents:
 - Enbrel (etanercept)
 - Cyltezo (adalimumab-adbm) OR Simlandi (adalimumabryvk) OR adalimumab-adaz (Hyrimoz unbranded) OR adalimumab-adbm (Cyltezo unbranded) OR adalimumabryvk (Simlandi unbranded)
 - Otezla (apremilast)
 - Skyrizi (risankizumab-rzaa)



- Steqeyma (ustekinumab-stba) SC OR Yesintek (ustekinumab-kfce) SC
- Taltz (ixekizumab)
- Tremfya (guselkumab)

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies. Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

Cosentyx (secukinumab) IV/SC

Cosentyx (secukinumab) IV is subject to review for site of service administration.

Cosentyx (secukinumab) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 2 years or older

- Has had an inadequate response or intolerance to 2 of the following agents:
 - Enbrel (etanercept)
 - Cyltezo (adalimumab-adbm) OR Simlandi (adalimumabryvk) OR adalimumab-adaz (Hyrimoz unbranded) OR adalimumab-adbm (Cyltezo unbranded) OR adalimumabryvk (Simlandi unbranded)
 - Otezla (apremilast)
 - Rinvoq (upadacitinib) **OR** Rinvoq LQ (upadacitinib)
 - Skyrizi (risankizumab-rzaa)
 - Steqeyma (ustekinumab-stba) SC OR Yesintek (ustekinumab-kfce) SC
 - Taltz (ixekizumab)
 - Tremfya (guselkumab)



 Xeljanz (tofacitinib) OR Xeljanz XR (tofacitinib extendedrelease)

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies.

Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

T-Cell Costimulation Modulators – Second Line

Orencia (abatacept) IV/SC

Orencia (abatacept) IV is subject to review for site of service administration.

Orencia (abatacept) IV/SC may be considered medically necessary for the treatment of adults with active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

- Has had an inadequate response or intolerance to 2 of the following agents:
 - Enbrel (etanercept)
 - Cyltezo (adalimumab-adbm) OR Simlandi (adalimumabryvk) OR adalimumab-adaz (Hyrimoz unbranded) OR adalimumab-adbm (Cyltezo unbranded) OR adalimumabryvk (Simlandi unbranded)
 - Otezla (apremilast)
 - Rinvog (upadacitinib) **OR** Rinvog LQ (upadacitinib)
 - Skyrizi (risankizumab-rzaa)
 - Steqeyma (ustekinumab-stba) SC OR Yesintek (ustekinumab-kfce) SC



- Taltz (ixekizumab)
- Tremfya (guselkumab)
- Xeljanz (tofacitinib) OR Xeljanz XR (tofacitinib extendedrelease)

OR

 Has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, or a demyelinating disorder

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies.

Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

Orencia (abatacept) SC may be considered medically necessary for the treatment of pediatric individuals with active psoriatic arthritis when:

• The individual is aged 2 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

AND

- Has had an inadequate response or intolerance to ONE of the following agents:
 - Enbrel (etanercept)
 - Rinvoq (upadacitinib) OR Rinvoq LQ (upadacitinib)
 - Steqeyma (ustekinumab-stba) SC **OR** Yesintek (ustekinumab-kfce) SC

OR

 Has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, or a demyelinating disorder



Step therapy tiers are listed below; please refer to the Policy section for details.

Section 3: Individual/Small Group/Student ISHIP METALLIC Formulary Plans (Rx Plan M1, M2, and M4) ONLY **Psoriatic Arthritis – First Line TNF-α Inhibitors IL-17** IL-12/23 **IL-23** Janus PDE-4 **Inhibitors Inhibitors** Inhibitor Kinase Inhibitor **Inhibitors** Inflectra (IV) Taltz Steqeyma Skyrizi (SC) Rinvoq / Otezla (SC) Rinvoq LQ (oral) (SC) Infliximab (Janssen Tremfya (SC) (oral) - unbranded) (IV) Yesintek (SC) [pen, syringe, and injector] Xeljanz / Remicade (IV) Xeljanz XR Simponi Aria (IV) (oral) Adalimumab-adaz Skyrizi (SC) (Hyrimoz unbranded) (SC) Adalimumabadbm (Cyltezo unbranded) (SC) Adalimumab-ryvk (Simlandi unbranded) (SC) Cyltezo (SC) Enbrel (SC) Simlandi (SC) **Psoriatic Arthritis – Second Line** IL-12/23 **T-Cell Costimulation TNF-α Inhibitors IL-17 Inhibitors Inhibitors Modulator** Avsola (IV) Bimzelx Orencia (IV/SC) Imuldosa (SC) (SC) Renflexis (IV) Otulfi (SC) Cosentyx Cimzia (SC) Pyzchiva (SC) (IV/SC) Simponi(SC) Selarsdi (SC)

- Abrilada (SC)
- Adalimumab-aacf (Idacio unbranded) (SC)
- Adalimumab-aaty (Yuflyma unbranded) (SC)
- Adalimumab-fkjp (Hulio unbranded) (SC)
- Amjevita (SC)
- Hadlima (SC)
- Hulio (SC)
- Humira (SC)
- Hyrimoz (SC)
- Idacio (SC)
- Yuflyma (SC)
- Yusimry (SC)

- Stelara (SC)
- Brand
 ustekinumab
 (Stelara
 unbranded)
 (SC)
- Brand ustekinumabaekn (Selarsdi unbranded) (SC)
- Brand ustekinumabttwe (Pyzchiva unbranded) (SC)
- Wezlana (SC)

Section 3: Individual/Small Group/Student ISHIP METALLIC Formulary Plans (Rx Plan M1, M2, and M4) ONLY

TNF-α Antagonists – First Line

Agent

- Cyltezo (adalimumabadbm) SC
- Simlandi (adalimumabryvk) SC
- Adalimumab-adaz (Hyrimoz unbranded) SC
- Adalimumab-adbm (Cyltezo unbranded) SC
- Adalimumab-ryvk
 (Simlandi unbranded) SC
- Enbrel (etanercept) SC
- Simponi Aria (golimumab) IV

Medical Necessity, Psoriatic Arthritis

Simponi Aria (golimumab) IV is subject to review for site of service administration.

Cyltezo (adalimumab-adbm), Simlandi (adalimumab-ryvk), adalimumab-adaz (Hyrimoz unbranded), adalimumab-adbm (Cyltezo unbranded), adalimumab-ryvk (Simlandi unbranded), Enbrel (etanercept) or Simponi Aria (golimumab) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist



Note:

This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy **5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies**. Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

- Inflectra (infliximabdyyb) IV
- Infliximab (Janssen unbranded) IV
- Remicade (infliximab) IV

Inflectra (infliximab-dyyb), Infliximab (Janssen – unbranded), and Remicade (infliximab) are subject to review for site of service administration.

Inflectra (infliximab-dyyb), Infliximab (Janssen – unbranded), and Remicade (infliximab) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

IL-17 Inhibitor - First Line

Taltz (ixekizumab) SC

Taltz (ixekizumab) SC may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

IL-12/23 Inhibitor - First Line

- Steqeyma (ustekinumabstba) SC
- Yesintek (ustekinumabkfce) SC

Steqeyma (ustekinumab-stba) SC and Yesintek (ustekinumab-kfce) SC may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 6 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

IL-23 Inhibitors - First Line



Section 3: Individual/Smal	I Group/Student ISHIP METALLIC Formulary Plans (Rx Plan
M1, M2, and M4) ONLY	
Tremfya (guselkumab) SC (pen, syringe, and injector)	 Tremfya (guselkumab) may be considered medically necessary for the treatment of active psoriatic arthritis when: The individual is aged 18 years or older AND Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist
Skyrizi (risankizumab-rzaa) SC	 Skyrizi (risankizumab-rzaa) may be considered medically necessary for the treatment of active psoriatic arthritis when: The individual is aged 18 years or older AND Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist
Janus Kinase Inhibitors - F	irst Line
Rinvoq (upadacitinib) oral Rinvoq LQ (upadacitinib) oral	 Rinvoq (upadacitinib) and Rinvoq LQ (upadacitinib) may be considered medically necessary for the treatment of active psoriatic arthritis when: The individual is aged 2 years or older AND The individual has had an inadequate response or intolerance to 1 or more TNF blockers AND Medication is being prescribed by or in consultation with a rheumatologist or dermatologist
 Xeljanz (tofacitinib) oral Xeljanz XR (tofacitinib extended-release) oral 	 Xeljanz (tofacitinib) and Xeljanz XR (tofacitinib extended-release) may be considered medically necessary for the treatment of active psoriatic arthritis when: The individual is aged 18 years or older AND Has had an inadequate response or intolerance to 1 or more TNF blockers AND Medication is being prescribed by or in consultation with a rheumatologist or dermatologist
PDE4 Inhibitor - First Line	

Otezla (apremilast) oral

Otezla (apremilast) may be considered medically necessary for the treatment of active psoriatic arthritis when:

The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

TNF-α Antagonists – Second Line

- Abrilada (adalimumabafzb) SC
- Adalimumab-aacf (Idacio unbranded) SC
- Adalimumab-aaty (Yuflyma unbranded) SC
- Adalimumab-fkjp (Hulio unbranded) SC
- Amjevita (adalimumabatto) SC
- Hadlima (adalimumabbwwd) SC
- Hulio (adalimumab-fkjp)
 SC
- Humira (adalimumab) SC
- Hyrimoz (adalimumabadaz) SC
- Idacio (adalimumab-aacf)
 SC
- Yuflyma (adalimumabaaty) SC
- Yusimry (adalimumabaqvh) SC

Abrilada (adalimumab-afzb), adalimumab-aacf (Idacio unbranded), adalimumab-aaty (Yuflyma unbranded), adalimumab-fkjp (Hulio unbranded), Amjevita (adalimumab-atto), Hadlima (adalimumab-bwwd), Hulio (adalimumab-fkjp), Humira (adalimumab), Hyrimoz (adalimumab-adaz), Idacio (adalimumab-aacf), Yuflyma (adalimumab-aaty), and Yusimry (adalimumab-aqvh) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

AND

- Has had an inadequate response or intolerance to ALL the following agents:
 - Cyltezo (adalimumab-adbm) OR adalimumab-adbm (Cyltezo unbranded)
 - Adalimumab-adaz (Hyrimoz unbranded)
 - Simlandi (adalimumab-ryvk) OR adalimumab-ryvk (Simlandi unbranded)

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies.

Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.



- Cimzia (certolizumab pegol) SC
- Simponi (golimumab) SC

Cimzia (certolizumab pegol) and Simponi (golimumab) SC may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

AND

- Has had an inadequate response or intolerance to 2 of the following agents:
 - Enbrel (etanercept)
 - Cyltezo (adalimumab-adbm) OR Simlandi (adalimumabryvk) OR adalimumab-adaz (Hyrimoz unbranded) OR adalimumab-adbm (Cyltezo unbranded) OR adalimumabryvk (Simlandi unbranded)
 - Otezla (apremilast)
 - Rinvoq (upadacitinib) or Rinvoq LQ (upadacitinib)
 - Skyrizi (risankizumab-rzaa) SC
 - Steqeyma (ustekinumab-stba) SC OR Yesintek (ustekinumab-kfce) SC
 - Taltz (ixekizumab)
 - Tremfya (guselkumab)
 - Xeljanz (tofacitinib) OR Xeljanz XR (tofacitinib extendedrelease)

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies.

Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

- Avsola (infliximab-axxq)
- Renflexis (infliximababda) IV

Avsola (infliximab-axxq) and Renflexis (infliximab-abda) are subject to review for site of service administration.



Avsola (infliximab-axxq) and Renflexis (infliximab-abda) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a rheumatologist or dermatologist

AND

 Has had a documented trial and treatment failure with Inflectra (infliximab-dyyb), Infliximab (Janssen – unbranded), or Remicade (infliximab)

IL-12/23 Inhibitors - Second Line

- Imuldosa (ustekinumabsrlf) SC
- Otulfi (ustekinumab-aauz)
 SC
- Pyzchiva (ustekinumabttwe) SC
- Selarsdi (ustekinumabackn) SC
- Stelara (ustekinumab) SC
- Ustekinumab (Stelara unbranded) SC
- Ustekinumab-aekn (Selarsdi unbranded) SC
- Ustekinumab-ttwe (Pyzchiva unbranded) SC
- Wezlana (ustekinumabauub) SC

Imuldosa (ustekinumab-srlf) SC, Otulfi (ustekinumab-aauz) SC, Pyzchiva (ustekinumab-ttwe) SC, Selarsdi (ustekinumab-ackn) SC, Stelara (ustekinumab), ustekinumab (Stelara unbranded) SC, ustekinumab-aekn (Selarsdi unbranded) SC, ustekinumab-ttwe (Pyzchiva unbranded) SC, and Wezlana (ustekinumab-auub) SC may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 6 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

AND

- Has had an inadequate response or intolerance to ALL the following agents:
 - Steqeyma (ustekinumab-stba) SC
 - Yesintek (ustekinumab-kfce) SC

IL-17 Inhibitors - Second Line

Bimzelx (bimekizumabbkzx) SC Bimzelx (bimekizumab-bkzx) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Has had an inadequate response or intolerance to 1 of the following agents:



- Enbrel (etanercept)
- Cyltezo (adalimumab-adbm) OR Simlandi (adalimumabryvk) OR adalimumab-adaz (Hyrimoz unbranded) OR adalimumab-adbm (Cyltezo unbranded) OR adalimumabryvk (Simlandi unbranded)
- Otezla (apremilast)
- Skyrizi (risankizumab-rzaa)
- Steqeyma (ustekinumab-stba) SC OR Yesintek (ustekinumab-kfce) SC
- Taltz (ixekizumab)
- Tremfya (guselkumab)

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies.

Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

Cosentyx (secukinumab) IV/SC

Cosentyx (secukinumab) IV is subject to review for site of service administration.

Cosentyx (secukinumab) may be considered medically necessary for the treatment of active psoriatic arthritis when:

• The individual is aged 2 years or older

- Has had an inadequate response or intolerance to 2 of the following agents:
 - Enbrel (etanercept)
 - Cyltezo (adalimumab-adbm) OR Simlandi (adalimumabryvk) OR adalimumab-adaz (Hyrimoz unbranded) OR adalimumab-adbm (Cyltezo unbranded) OR adalimumabryvk (Simlandi unbranded)



- Otezla (apremilast)
- Rinvoq (upadacitinib) **OR** Rinvoq LQ (upadacitinib)
- Skyrizi (risankizumab-rzaa)
- Steqeyma (ustekinumab-stba) SC OR Yesintek (ustekinumab-kfce) SC
- Taltz (ixekizumab)
- Tremfya (guselkumab)
- Xeljanz (tofacitinib) OR Xeljanz XR (tofacitinib extendedrelease)

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies.

Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

T-Cell Costimulation Modulators – Second Line

Orencia (abatacept) IV/SC

Orencia (abatacept) IV is subject to review for site of service administration.

Orencia (abatacept) IV/SC may be considered medically necessary for the treatment of adults with active psoriatic arthritis when:

• The individual is aged 18 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist

- Has had an inadequate response or intolerance to 2 of the following agents:
 - Enbrel (etanercept)



- Cyltezo (adalimumab-adbm) OR Simlandi (adalimumabryvk) OR adalimumab-adaz (Hyrimoz unbranded) OR adalimumab-adbm (Cyltezo unbranded) OR adalimumabryvk (Simlandi unbranded)
- Otezla (apremilast)
- Rinvoq (upadacitinib) **OR** Rinvoq LQ (upadacitinib)
- Skyrizi (risankizumab-rzaa)
- Steqeyma (ustekinumab-stba) SC OR Yesintek (ustekinumab-kfce) SC
- Taltz (ixekizumab)
- Tremfya (guselkumab)
- Xeljanz (tofacitinib) OR Xeljanz XR (tofacitinib extendedrelease)

OR

 Has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, or a demyelinating disorder

Note: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies.

Please check the member Plan booklet or member ID card to determine whether this policy criteria applies.

Orencia (abatacept) SC may be considered medically necessary for the treatment of pediatric individuals with active psoriatic arthritis when:

• The individual is aged 2 years or older

AND

 Medication is being prescribed by or in consultation with a dermatologist or a rheumatologist



Section 3: Individual/Small Group/Student ISHIP METALLIC Formulary Plans (Rx Plan M1, M2, and M4) ONLY				
	 Has had an inadequate response or intolerance to ONE of the following agents: Enbrel (etanercept) 			
	 Rinvoq (upadacitinib) OR Rinvoq LQ (upadacitinib) Steqeyma (ustekinumab-stba) SC OR Yesintek (ustekinumab-kfce) SC 			
	OR			
	 Has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, or a demyelinating disorder 			

Drug	Investigational
As listed	The medications listed in this policy are subject to the product's US Food and Drug Administration (FDA) dosage and administration prescribing information.
	All other uses of the above-named agents when used in combination with each other or for conditions not outlined in this policy or Related Medical Policies are considered investigational.

Drug	Not Medically Necessary
As listed	All other uses of the drugs for approved conditions listed in
	this policy are considered not medically necessary.

Length of Approval		
Approval	Criteria	
Initial authorization	Non-formulary exception reviews and all other reviews for all drugs listed in policy may be approved up to 12 months.	
Re-authorization criteria	Non-formulary exception reviews and all other reviews for all drugs listed in policy may be approved up to 12 months as long as the drug-specific coverage criteria are met, and chart	

Length of Approv	al
Approval	Criteria
	notes demonstrate that the individual continues to show a
	positive clinical response to therapy.

Documentation Requirements

The individual's medical records submitted for review for all conditions should document that medical necessity criteria are met. The record should include the following:

 Office visit notes that contain the diagnosis, relevant history, physical evaluation and medication history

Coding

Code	Description
HCPCS	
J0129	Injection, abatacept (Orencia), 10 mg (code may be used for Medicare when drug administered under the direct supervision of a physician, not for use when drug is self-administered)
J0139	Injection, adalimumab (Humira), 1 mg (new code effective 01/01/25)
J0717	Injection, certolizumab pegol (Cimzia), 1 mg (code may be used for Medicare when drug administered under the direct supervision of a physician, not for use when drug is self-administered)
J1438	Injection, etanercept (Enbrel), 25mg (code may be used for Medicare when drug administered under the direct supervision of a physician, not for use when drug is self-administered)
J1602	Injection, golimumab (Simponi Aria), 1 mg, for intravenous use
J1628	Injection, guselkumab (Tremfya), 1 mg
J1745	Injection, infliximab, excludes biosimilar (Remicade or Janssen unbranded), 10mg
J3247	Injection, secukinumab, intravenous, (Cosentyx) 1 mg
J3357	Injection, ustekinumab (Stelera), 1mg
J3590	Unclassified biologics (use to report: Abrilada, Adalimumab-adaz HCF, Amjevita, Bimzelx, Cyltezo, Hadlima, Hulio, Hyrimoz LCF, Hyrimoz HCF, Imuldosa, Kevzara, Kineret, Sandoz (unbranded), Simponi, Skyrizi, Taltz, Yuflyma, Yusimry)



Code	Description
Q5098	Injection, ustekinumab-srlf (Imuldosa), biosimilar, 1 mg (new code effective 07/01/25)
Q5099	Injection, ustekinumab-stba (Steqeyma), biosimilar, 1 mg (new code effective 07/01/25)
Q5100	Injection, ustekinumab-kfce (Yesintek), biosimilar, 1 mg (new code effective 07/01/25)
Q5103	Injection, infliximab-dyyb, biosimilar (Inflectra), 10 mg
Q5104	Injection, infliximab-abda, biosimilar (Renflexis), 10 mg
Q5115	Injection, rituximab-abbs, biosimilar (Truxima), 10 mg
Q5121	Injection, infliximab-axxq, biosimilar (Avsola), 10 mg
Q5137	Injection, ustekinumab-auub (Wezlana), biosimilar, SC, 1 mg
Q5138	Injection, ustekinumab-auub (Wezlana), biosimilar, IV, 1 mg
Q5140	Injection, adalimumab-fkjp (Hulio), biosimilar, 1 mg (new code effective 01/01/25)
Q5141	Injection, adalimumab-aaty (Yuflyma), biosimilar, 1 mg (new code effective 01/01/25)
Q5142	Injection, adalimumab-ryvk biosimilar (Simlandi), 1 mg (new code effective 01/01/25)
Q5143	Injection, adalimumab-adbm, biosimilar (Cyltezo), 1 mg (new code effective 01/01/25)
Q5144	Injection, adalimumab-aacf (Idacio), biosimilar, 1 mg (new code effective 01/01/25)
Q5145	Injection, adalimumab-afzb (Abrilada), biosimilar, 1 mg (new code effective 01/01/25)
Q9996	Injection, ustekinumab-ttwe (Pyzchiva), subcutaneous, 1 mg
Q9998	Injection, ustekinumab-aekn (Selarsdi), 1 mg
Q9999	Injection, ustekinumab-aauz (Otulfi), biosimilar, 1 mg (new code effective 04/01/25)

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Related Information

Consideration of Age

Age limits specified in this policy are determined according to FDA-approved indications, where applicable.



For site of service for medical necessity the age described in this policy is 13 years of age or older. Site of service is defined as the location where the drug is administered, such as a hospital-based outpatient setting, an infusion center, a physician's office, or at home. The age criterion for site of service for medical necessity is based on the following: Pediatric individuals are not small adults. Pediatric individuals differ physiologically, developmentally, cognitively, and emotionally from adult individuals, and vary by age groups from infancy to teen. Children often require smaller doses than adults, lower infusion rates, appropriately sized equipment, the right venipuncture site determined by therapy and age, and behavioral management during administration of care. Specialty infusion training is therefore necessary for pediatric IV insertions and therapy. Due to pediatrics unique physiology and psychology, site of service review is limited to individuals above the age of 13.

Evidence Review

Psoriatic Arthritis

Psoriatic Arthritis (PsA) is characterized as a spondyloarthropathy associated with psoriasis. The true incidence is unknown and is variably reported to occur in 6-42% (25% is considered a reasonable estimate) of individuals with psoriasis, an immunologic skin disease which occurs in 2-3% of the general population. There is similarity in the histopathogenesis of PsA and RA, including the role of cytokines such as tumor necrosis factor alpha (TNF- α), although there are important differences as well. Several subsets of PsA have also been described. PsA is characterized by stiffness - both peripheral and spine inflammation and pain - joint deformities related to joint destruction, dactylitis, enthesitis (inflammation at insertion sites of tendons, ligaments, and joint capsule fibers), and psoriasis skin plaques. The course of PsA is variable, but the majority of individuals develop a chronic progressive form of the disease resulting in joint destruction, unless treated effectively. Although less well characterized than in RA, similar levels of disability, decreased quality of life, increased co-morbidities, and premature mortality are now being noted in long term registry studies.

Pharmacologic therapy combined with a physical rehabilitation program is the most effective available treatment for psoriatic arthritis (PsA). As with RA, early initiation of pharmacologic therapy is needed to avoid joint damage and disability.

NSAIDs have customarily been used in milder disease along with corticosteroids or traditional DMARDs. Moderate to severe disease requires the use of traditional DMARDs such as MTX,



sulfasalazine, or the anti-TNF agents. Azathioprine and cyclosporine are rarely used. Retinoids, phototherapy, and topical and systemic corticosteroids have also been used to treat the skin manifestations of PsA. In January 2002, etanercept, a TNF- α inhibitor became the first therapy to be approved for the indication. Adalimumab has also recently received FDA-approval for this indication. Additionally, infliximab has been demonstrated effective for this condition in at least one randomized, double-blind, controlled clinical trial. FDA has since approved the newer TNF- α inhibitors certolizumab pegol and golimumab for this indication. More recently, the IL12/IL23 inhibitor ustekinumab and the phosphodiesterase four inhibitor apremilast are now approved.

2025 Update

Moved psoriatic arthritis criteria for the infliximab products, adalimumab products, Simponi Aria, Enbrel, Taltz, Stelara, Skyrizi, Tremfya, Rinvoq, Rinvoq LQ, Xeljanz, Xeljanz XR, Otezla, Cimzia, Simponi, Bimzelx, Cosentyx, and Orencia from Policy 5.01.550 to 5.01.645. Added coverage criteria for Imuldosa (ustekinumab-srlf), Otulfi (ustekinumab-aauz), Pyzchiva (ustekinumab-ttwe), Selarsdi (ustekinumab-ackn), Stegeyma (ustekinumab-stba), ustekinumab (Stelara unbranded), ustekinumab-aekn (Selarsdi unbranded), ustekinumab-ttwe (Pyzchiva unbranded), Yesintek (ustekinumab-kfce), and Wezlana (ustekinumab-auub). Added the following to note to select criteria for Bimzelx (bimekizumab-bkzx), Cimzia (certolizumab pegol), Cosentyx (secukinumab) IV/SC, Orencia (abatacept) IV/SC, and Simponi (golimumab) SC: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formulary plans can be found in policy 5.01.647 Medical Necessity Criteria for Custom Open and Preferred Formularies. Clarified that the Site of Service Medical Necessity criteria can apply to injection drugs. Updated Stelara (ustekinumab) coverage criteria to require trial with Stegeyma (ustekinumab-aauz) and Yesintek (ustekinumab-kfce) for individuals not previously treated. New policy sections for Metallic (individual/small group/student ISHIP) formulary plans, Essentials formulary plans, and Open/Preferred/Select formulary plans and plans with no pharmacy benefit coverage. Added different coverage criteria for Metallic (individual/small group/student ISHIP) formulary and Essentials formulary plans for the following drugs: Enbrel (etanercept), adalimumab products, infliximab products, Taltz (ixekizumab), ustekinumab products, Skyrizi (risankizumab-rzaa), Tremfya (guselkumab), Otezla (apremilast), Bimzelx (bimekizumab-bkzx), Cosentyx (secukinumab), Cimzia (certolizumab pegol), Rinvoq/Rinvoq LQ (upadacitinib), Xeljanz/Xeljanz XR (tofacitinib extended-release), Simponi (golimumab), Simponi Aria (golimumab), Rinvoq LQ (upadacitinib), and Orencia (abatacept).



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- 18. Stelara (ustekinumab). Prescribing Information. Janssen Biotech, Inc. Horsham, PA. Revised November 2024.
- 19. Orencia (abatacept). Prescribing Information. Bristol-Myers Squibb. Princeton, NJ. Revised May 2024.
- 20. Simlandi (adalimumab-ryvk). Prescribing Information. Teva Pharmaceuticals. Parsippany, NJ. Revised June 2024.
- 21. Cimzia (certolizumab pegol). Prescribing Information. UCB, Inc. Smyrna, GA. Revised September 2024.
- 22. Bimzelx (bimekizumab-bkzx). Prescribing Information. UCB, Inc. Smyrna, GA. Revised November 2024.

History

Date	Comments
07/01/25	New policy, approved June 10, 2025. Moved psoriatic arthritis criteria for the infliximab products, adalimumab products, Simponi Aria, Enbrel, Taltz, Stelara, Skyrizi, Tremfya, Rinvoq, Rinvoq LQ, Xeljanz, Xeljanz XR, Otezla, Cimzia, Simponi, Bimzelx, Cosentyx, and Orencia from Policy 5.01.550 to 5.01.645. Added coverage criteria for Imuldosa (ustekinumab-srlf), Otulfi (ustekinumab-aauz), Pyzchiva (ustekinumab-ttwe), Selarsdi (ustekinumab-srlf), Steqeyma (ustekinumab-stba), ustekinumab (Stelara unbranded), ustekinumab-aekn (Selarsdi unbranded), ustekinumab-atwe (Pyzchiva unbranded), Vesintek (ustekinumab-kfce), and Wezlana (ustekinumab-auub). Added the following to note to select criteria for Bimzelx (bimekizumab-bkzx), Cimzia (certolizumab pegol), Cosentyx (secukinumab) IV/SC, Orencia (abatacept) IV/SC, and Simponi (golimumab) SC: This medical necessity criteria does not apply to one Open formulary (Formulary ID: 6062; Rx Plan F1) and one Preferred formulary (Formulary ID: 6064; Rx Plan G3). The criteria for members with these custom Open and Preferred formularies. Added HCPCS code Q9999. Also added HCPCS codes Q5137 and Q5138 for Wezlana, Q9996 and Q9998 for Steqeyma. Also added new HCPCS codes Q5098 for Imuldosa and Q5099 for Otulfi and Q5100 for Yesintek. The following policy changes are effective October 3, 2025, following 90-day provider notification. Clarified that the Site of Service Medical Necessity criteria can apply to injection drugs. Updated Stelara (ustekinumab) coverage criteria to require trial with Steqeyma (ustekinumab-aauz) and Yesintek (ustekinumab-kfce) for individuals not previously treated. New policy sections for Metallic (individual/small group/student ISHIP) formulary plans, Essentials formulary plans, and Open/Preferred/Select formulary plans and plans with no pharmacy benefit coverage. Added different coverage criteria for Metallic (individual/small group/student ISHIP) formulary plans for the following drugs: Enbrel (etanercept), adalimumab products, infliximab products, Taltz (ixek

Disclaimer: This medical policy is a guide in evaluating the medical necessity of a particular service or treatment. The Company adopts policies after careful review of published peer-reviewed scientific literature, national guidelines and local standards of practice. Since medical technology is constantly changing, the Company reserves the right to review and update policies as appropriate. Member contracts differ in their benefits. Always consult the member benefit



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