



Drug Coverage Policy

Effective Date11/01/2025
Coverage Policy Number.....PSM017
Policy Title.....Inflammatory Conditions
Preferred Specialty Management Policy
for Employer Plans: Legacy Prescription
Drug Lists

Inflammatory Conditions Preferred Specialty Management Policy for Employer Plans: Legacy Prescription Drug Lists

Tumor Necrosis Factor Inhibitors
• Adalimumab Products* ◦ adalimumab-adaz subcutaneous injection (Sandoz/Novartis) ◦ adalimumab-adbm subcutaneous injection (Boehringer Ingelheim) ◦ adalimumab-ryvk subcutaneous injection (Alvotect/Teva) ◦ Cyltezo® (adalimumab-adbm subcutaneous injection – Boehringer Ingelheim) ◦ Humira® (adalimumab subcutaneous injection – AbbVie, Cordavis) ◦ Simlandi (adalimumab-ryvk subcutaneous injection – Alvotect/Teva) • Cimzia® (certolizumab pegol subcutaneous injection – UCB) • Enbrel® (etanercept subcutaneous injection – Amgen) • Simponi® (golimumab subcutaneous injection – Janssen Biotech/Johnson & Johnson) • Zymfentra® (infliximab-dyyb subcutaneous injection – Celltrion)
Interleukin-6 Blockers
• Tocilizumab Subcutaneous Products ◦ Actemra® (tocilizumab subcutaneous injection – Genentech/Roche) ◦ Tyenne® (tocilizumab-aazg subcutaneous injection – Fresenius Kabi) • Kevzara™ (sarilumab subcutaneous injection – Regeneron)
Interleukin-17 Blockers
• Bimzelx® (bimekizumab subcutaneous injection – UCB) • Cosentyx® (secukinumab subcutaneous injection – Novartis) • Siliq™ (brodalumab subcutaneous injection – Valeant) • Taltz® (ixekizumab subcutaneous injection – Eli Lilly)
Interleukin-23 Blockers
• Ilumya™ (tildrakizumab-asmn subcutaneous injection – Sun/Merck) • Omvoh® (mirakizumab-mrkz subcutaneous injection – Eli Lilly) • Skyrizi™ (risankizumab-rzaa subcutaneous injection – AbbVie) • Tremfya™ (guselkumab subcutaneous injection – Janssen Biotech/Johnson & Johnson)
Interleukin 12/23 Blocker
• Ustekinumab Subcutaneous Products* ◦ Selarsdi™ (ustekinumab-aekn subcutaneous injection – Alvotect/Teva) ◦ Stelara® (ustekinumab subcutaneous injection – Janssen Biotech/Johnson & Johnson) ◦ ustekinumab-ttwe subcutaneous injection (Quallent) ◦ Yesintek™ (ustekinumab-kfce subcutaneous injection – Biocon)

Interleukin-1 Blocker
• Kineret® (anakinra subcutaneous injection – Swedish Orphan Biovitrim)
T-Cell Costimulation Modulator
• Orenia® (abatacept subcutaneous injection – Bristol Myers Squibb)
Integrin Receptor Antagonist
• Entyvio® (vedolizumab subcutaneous injection – Takeda)
Janus Kinases Inhibitors
• Olumiant® (baricitinib tablets – Eli Lilly)
• Rinvoq™ (upadacitinib extended-release tablets – AbbVie)
• Rinvoq® LQ (upadacitinib oral solution – AbbVie)
• Xeljanz® (tofacitinib tablets, tofacitinib oral solution – Pfizer)
• Xeljanz® XR (tofacitinib extended-release tablets – Pfizer)
Phosphodiesterase Type 4 Inhibitor
• Otezla® (apremilast tablets – Amgen)
Sphingosine 1-Phosphate Receptor Modulator
• Velsipity™ (etrasimod tablets – Pfizer)
• Zeposia® (ozanimod capsules – Celgene)
Tyrosine Kinase 2 Inhibitor
• Sotyktu™ (deucravacitinib tablets – Bristol Myers Squibb)

* For Non-Preferred products, refer to the *respective Inflammatory Conditions Preferred Specialty Management Policy for Employer Plans: Legacy Prescription Drug Lists*

INSTRUCTIONS FOR USE

The following Coverage Policy applies to health benefit plans administered by Cigna Companies. Certain Cigna Companies and/or lines of business only provide utilization review services to clients and do not make coverage determinations. References to standard benefit plan language and coverage determinations do not apply to those clients. Coverage Policies are intended to provide guidance in interpreting certain standard benefit plans administered by Cigna Companies. Please note, the terms of a customer's particular benefit plan document [Group Service Agreement, Evidence of Coverage, Certificate of Coverage, Summary Plan Description (SPD) or similar plan document] may differ significantly from the standard benefit plans upon which these Coverage Policies are based. For example, a customer's benefit plan document may contain a specific exclusion related to a topic addressed in a Coverage Policy. In the event of a conflict, a customer's benefit plan document always supersedes the information in the Coverage Policies. In the absence of a controlling federal or state coverage mandate, benefits are ultimately determined by the terms of the applicable benefit plan document. Coverage determinations in each specific instance require consideration of 1) the terms of the applicable benefit plan document in effect on the date of service; 2) any applicable laws/regulations; 3) any relevant collateral source materials including Coverage Policies and; 4) the specific facts of the particular situation. Each coverage request should be reviewed on its own merits. Medical directors are expected to exercise clinical judgment where appropriate and have discretion in making individual coverage determinations. Where coverage for care or services does not depend on specific circumstances, reimbursement will only be provided if a requested service(s) is submitted in accordance with the relevant criteria outlined in the applicable Coverage Policy, including covered diagnosis and/or procedure code(s). Reimbursement is not allowed for services when billed for conditions or diagnoses that are not covered under this Coverage Policy (see "Coding Information" below). When billing, providers must use the most appropriate codes as of the effective date of the submission. Claims submitted for services that are not accompanied by covered code(s) under the applicable Coverage Policy will be denied as not covered. Coverage Policies relate exclusively to the administration of health benefit plans. Coverage Policies are not recommendations for treatment and should never be used as treatment guidelines. In certain markets, delegated vendor guidelines may be used to support

medical necessity and other coverage determinations.

OVERVIEW

Several products are available for use in inflammatory conditions such as rheumatoid arthritis, ankylosing spondylitis, juvenile idiopathic arthritis, psoriatic arthritis, plaque psoriasis, Crohn's disease, and ulcerative colitis.¹⁻²⁵ This policy involves the use of the products listed above.

The FDA-approved indications for each product listed in this policy are documented in [Appendix A](#). For more information on criteria within a Prior Authorization program by specific condition refer to the respective standard *Prior Authorization Policy*.

Coverage Policy

Policy Statement

For all medications, this program requires the patient to meet the respective standard *Prior Authorization Policy* criteria. Additionally, this program requires trial(s) of the Preferred Product(s) according to the table below, when clinically appropriate, prior to the approval of the Non-Preferred Products. There are also situations when trials of Non-Preferred Products will be considered; see criteria below. Other details of the program are as follows:

- **Continuation of Therapy:** Approval for a patient continuing therapy with a Non-Preferred subcutaneous or oral Product must be supported with verification, noted in the criteria as either **[verification in prescription claims history required]** or, if not available, as **[verification by prescriber required]**.
 - If the patient has at least 130 days of prescription claims history on file, claims history must support that the patient has received the Non-Preferred Product for the specified period of time (90 or 120 days) within a 130-day look-back period; OR
 - When 130 days of the patient's prescription claim history file is unavailable for verification, the prescriber must verify that the patient has been receiving the Non-Preferred Product for a specified period of time (90 or 120 days), AND that the patient has been receiving the Non-Preferred Product via paid claims (e.g., patient has not been receiving samples or coupons or other types of waivers in order to obtain access to the Non-Preferred Product).
 - For a patient continuing therapy, other conditions may also apply. Refer to criteria below.
- **Approval Duration:** All approvals for continuation of therapy for Preferred and Non-Preferred Products are provided for 1 year unless noted otherwise below. In cases where the initial approval is authorized in months, 1 month is equal to 30 days.

Documentation: When documentation is required, the prescriber must provide written documentation supporting the trials of these other Products, noted in the criteria as **[documentation required]**. Documentation may include, but is not limited to, chart notes, prescription claims records, and/or prescription receipts.

Preferred and Non-Preferred Products– Rheumatology Indications.^{YQ}

	Rheumatology				
	RA	JIA	AS	nr-axSpA	PsA
Step 1 Preferred	•Enbrel •Adalimumab Products^ – Cyltezo/ adalimumab-adbm,	•Enbrel •Adalimumab Products^ – Cyltezo/ adalimumab-adbm, Simlandi/ adalimumab-	•Enbrel •Adalimumab Products^ – Cyltezo/ adalimumab-adbm, Simlandi/	•Cimzia •Taltz	•Enbrel •Adalimumab Products^ – Cyltezo/ adalimumab-adbm, Simlandi/

	Simlandi/ adalimumab- ryvk, Humira (NDCs starting with 0074)	ryvk, Humira (NDCs starting with 0074)	adalimumab- ryvk, Humira (NDCs starting with 0074) • Taltz		adalimumab- ryvk, Humira (NDCs starting with 00074) • Otezla • Skyrizi SC [#] • Ustekinumab SC Products ^κ - Stelara SC, Selarsdi SC, ustekinumab- ttwe SC, Yesintek SC ^k • Taltz • Tremfya SC
Step 2a Non-Preferred (directed to ONE Step 1 Product)	• Tocilizumab SC Products – Actemra SC, Tyenne SC <i>Directed to adalimumab specifically.</i> • Rinvoq Xeljanz tablets/ Xeljanz XR tablets	• Tocilizumab SC Products - Actemra SC, Tyenne SC <i>Directed to adalimumab specifically. JIA Step SC is for PJIA.</i> • Rinvoq/Rinvoq LQ • Xeljanz tablets/ Xeljanz oral solution	• Rinvoq <i>Directed specifically to Enbrel or adalimumab.</i> • Xeljanz tablets/ Xeljanz XR tablets <i>Directed specifically to Enbrel or adalimumab.</i>	• Rinvoq <i>Directed specifically to Cimzia.</i>	• Rinvoq/ Rinvoq LQ <i>Directed specifically to Enbrel or adalimumab.</i> • Xeljanz tablets/ Xeljanz XR tablets <i>Directed specifically to Enbrel or adalimumab.</i>
Step 2b Non-Preferred (directed to ONE Step 1 Product)	--	--	• Bimzelx	• Bimzelx	• Bimzelx
Step 3a Non-Preferred (directed to TWO Step 1 or 2a Products) [documentation required] [*]	• Cimzia • Kevzara • Kineret • Olumiant • Orencia SC • Simponi SC	• Cimzia • Kevzara • Orencia SC	• Cimzia • Cosentyx SC • Simponi SC	• Cosentyx SC	• Cimzia • Cosentyx SC • Orencia SC • Simponi SC

^κ For Non-Preferred Products, refer to the *Inflammatory Conditions – Adalimumab Products Preferred Specialty Management Policy (PSM003)*; ^Ω For Non-Preferred Products, refer to the *Inflammatory Conditions – Ustekinumab Subcutaneous Products Preferred Specialty Management Policy for Legacy Prescription Drug List Plans (PSM022)*; Note: Ustekinumab subcutaneous injection (Janssen Biotech) 45mg vial is covered as a Preferred Product on the medical benefit only. RA – Rheumatoid arthritis; ^ A trial of more than one adalimumab product counts as ONE Preferred Product; ^{*} A trial of more than one ustekinumab product counts as ONE Preferred Product; JIA – Juvenile idiopathic arthritis; AS – Ankylosing spondylitis; nr-axSpA – Nonradiographic axial spondyloarthritis; PsA – Psoriatic arthritis; SC – Subcutaneous; [#] Pen and syringe; PJIA – Polyarticular juvenile idiopathic arthritis; ^{*} The prescriber must provide written documentation supporting the trial of Preferred Products, noted in the criteria as **[documentation required]**. Documentation may include, but is not limited to, chart notes, prescription claims records, and/or prescription receipts.

Preferred and Non-Preferred Products – Dermatology and Gastroenterology Indications.^{¥Ω}

	Dermatology		Gastroenterology	
	HS	Psoriasis	CD	UC
Step 1 Preferred	• Adalimumab Products[^] – Cyltezo/ adalimumab-adbm, Simlandi/ adalimumab-ryvk, Humira (NDCs starting with 0074) • Cosentyx SC	• Enbrel • Adalimumab Products[^] – Cyltezo/ adalimumab-adbm, Simlandi/ adalimumab-ryvk, Humira (NDCs starting with 0074) • Otezla • Skyrizi SC[#] • Sotyktu • Ustekinumab SC Products[*] – Stelara SC, Selarsdi SC, ustekinumab-ttwe SC, Yesintek SC • Taltz • Tremfya SC	• Adalimumab Products[^] – Cyltezo/ adalimumab-adbm, Simlandi/ adalimumab-ryvk, Humira (NDCs starting with 0074) • Omvoh SC • Skyrizi SC (on-body injector) • Tremfya SC • Ustekinumab SC Products[*] – Stelara SC, Selarsdi SC, ustekinumab-ttwe SC, Yesintek SC • Zymfentra	• Adalimumab Products[^] – Cyltezo/ adalimumab-adbm, Simlandi/ adalimumab-ryvk, Humira (NDCs starting with 0074) • Omvoh SC • Skyrizi SC (on-body injector) • Ustekinumab SC Products[*] – Stelara SC, Selarsdi SC, ustekinumab-ttwe SC, Yesintek SC • Tremfya SC • Velsipity • Zymfentra
Step 2a Non-Preferred (directed to ONE Step 1 Product)	--	--	• Cimzia • Rinvoq Directed to adalimumab specifically.	• Rinvoq Directed to adalimumab specifically. • Simponi SC. • Xeljanz tablets/ Xeljanz/ XR tablets Directed to adalimumab specifically.
Step 2b Non-Preferred (directed to ONE Step 1 Product)	• Bimzelx	• Bimzelx	--	--
Step 3a Non-Preferred (directed to TWO Step 1 or 2a Products) [documentation required]*	--	• Cimzia • Cosentyx SC • Ilumya • Siliq	• Entyvio SC	• Entyvio SC
Step 3b Non-Preferred (directed to TWO Step 1 Products)	--	--	--	• Zeposia Refer to MS and UC – Zeposia PSM Policy

[¥] For Non-Preferred Products, refer to the *Inflammatory Conditions – Adalimumab Products Preferred Specialty Management Policy (PSM003)*; ^Ω For Non-Preferred Products, refer to the *Inflammatory Conditions – Ustekinumab Subcutaneous Products Preferred Specialty Management Policy for Legacy Prescription Drug List Plans (PSM022)*; Note: Ustekinumab subcutaneous injection (Janssen Biotech) 45mg vial is covered as a Preferred Product on the medical benefit only. RA – Rheumatoid arthritis; [^] A trial of more than one adalimumab product counts as ONE Preferred Product; ^{*} A trial of more than one ustekinumab product counts as ONE Preferred Product; JIA – Juvenile idiopathic arthritis; AS – Ankylosing spondylitis; nr-axSpA – Nonradiographic axial spondyloarthritis; PsA – Psoriatic arthritis; SC – Subcutaneous; [#] Pen and syringe; PJIA – Polyarticular juvenile idiopathic arthritis; * The prescriber must provide written documentation supporting

the trial of Preferred Products, noted in the criteria as **[documentation required]**. Documentation may include, but is not limited to, chart notes, prescription claims records, and/or prescription receipts.

Inflammatory Conditions non-preferred products are considered medically necessary when the following non-preferred product exception criteria are met. Any other exception is considered not medically necessary.

Non-Preferred Product	Exception Criteria
Tumor Necrosis Factor Inhibitors	
Cimzia	1. <u>Rheumatoid Arthritis – Initial Therapy.</u> A) Approve for 6 months if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria; AND ii. Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [documentation required]. <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Step 1 or Step 2 Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 2. <u>Ankylosing Spondylitis – Initial Therapy.</u> A) Approve for 6 months if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria; AND ii. Patient has tried TWO of Enbrel, an adalimumab product, Rinvoq, Taltz, or Xeljanz/XR [documentation required]. <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product.

B) If the patient has met criterion 2Ai (the standard *Inflammatory Conditions – Cimzia Prior Authorization Policy* criteria), but criterion 2Aii is not met: a request for a Step 1 or Step 2 Product (Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Taltz, Xeljanz tablets, or Xeljanz XR) may be reviewed using the respective standard *Inflammatory Conditions Prior Authorization Policy* criteria.

3. Juvenile Idiopathic Arthritis – Initial Therapy.

A) Approve for 6 months if the patient meets BOTH of the following (i and ii):

i. Patient meets the standard *Inflammatory Conditions – Cimzia Prior Authorization Policy* criteria; AND

ii. Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq/Rinvoq LQ, and Xeljanz **[documentation required]**; OR

Note: Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of both tocilizumab products counts as **ONE** product. Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as **ONE** product. A trial of either or both Xeljanz products (Xeljanz tablets and Xeljanz oral solution) collectively counts as **ONE** product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as **ONE** product. A trial of a tocilizumab intravenous product (Actemra intravenous, biosimilar), Kevzara, Orencia intravenous or subcutaneous, an infliximab product (e.g., Remicade, biosimilars), or Simponi Aria also counts **[documentation required]**.

B) If the patient has met criterion 3Ai (the standard *Inflammatory Conditions – Cimzia Prior Authorization Policy* criteria), but criterion 3Aii is not met: a request for a Step 1 or Step 2a Product (Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-ryvk, Humira [NDCs starting with 00074], Simlandi, Rinvoq, Rinvoq LQ, Xeljanz tablets, or Xeljanz oral solution) may be reviewed using the respective standard *Inflammatory Conditions – Prior Authorization Policy* criteria.

4. Psoriatic Arthritis – Initial Therapy.

A) Approve for 6 months if the patient meets BOTH of the following (i and ii):

i. Patient meets the standard *Inflammatory Conditions – Cimzia Prior Authorization Policy* criteria; AND

ii. Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, an ustekinumab subcutaneous product, Taltz, Tremfya subcutaneous, and Xeljanz/XR **[documentation required]**.

Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, adalimumab-adbm, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn,

	Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. B) If the patient has met criterion 4Ai (the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria), but criterion 4Aii is not met: a request for a Step 1 or Step 2 Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz tablets, or Xeljanz XR</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 5. <u>Plaque Psoriasis – Initial Therapy.</u> A) Approve for 3 months if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria; AND ii. Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, an ustekinumab subcutaneous product, Taltz, or Tremfya subcutaneous [documentation required]. <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, adalimumab-adbm, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. B) If the patient has met criterion 5Ai (the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria), but criterion 5Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 6. <u>Crohn’s Disease – Initial Therapy.</u> A) Approve for 6 months if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of an adalimumab product, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), an ustekinumab subcutaneous product, Tremfya subcutaneous, or Zymfentra.
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	<u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, adalimumab-adbm, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, Yesintek. A trial of an infliximab intravenous product (e.g., Remicade, biosimilars), Omvoh intravenous, Skyrizi intravenous, Tremfya intravenous, or ustekinumab intravenous also counts. B) If the patient has met criterion 6Ai (the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria), but criterion 6Aii is not met: a request for a Preferred Product (<u>Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous [on-body injector], Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Tremfya subcutaneous, or Zymfentra</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 7. <u>Rheumatoid Arthritis, Ankylosing Spondylitis, Juvenile Idiopathic Arthritis, Psoriatic Arthritis, Plaque Psoriasis, or Crohn’s Disease – Patient is Currently Receiving Cimzia.</u> A) Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a, b, c, d, e, <u>or</u> f): a) Patient has <u>Rheumatoid Arthritis</u> and has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. b) Patient has <u>Ankylosing Spondylitis</u> and has tried TWO of Enbrel, an adalimumab product, Rinvoq, Taltz, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. c) Patient has <u>Juvenile Idiopathic Arthritis</u> and has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq/Rinvoq LQ, and Xeljanz [documentation required]; OR
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	<u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of both tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz tablets and Xeljanz oral solution) collectively counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. A trial of a tocilizumab intravenous product (Actemra intravenous, biosimilar), Kevzara, Orencia intravenous or subcutaneous, an infliximab product (e.g., Remicade, biosimilars), or Simponi Aria also counts [documentation required]. d) Patient has <u>Psoriatic Arthritis</u> and has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, an ustekinumab subcutaneous product, Taltz, Tremfya subcutaneous, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. e) Patient has <u>Plaque Psoriasis</u> and has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, ustekinumab subcutaneous product, Taltz, or Tremfya subcutaneous [documentation required]; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, adalimumab-adbm, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. f) Patient has <u>Crohn's Disease</u> and has tried one of an adalimumab product, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), an ustekinumab subcutaneous product, Tremfya subcutaneous, or Zymfentra; OR
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	<u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, Yesintek. A trial of an infliximab intravenous product (e.g., Remicade, biosimilars), Omvoh intravenous, Skyrizi intravenous, Tremfya intravenous, or ustekinumab intravenous also counts. g) Patient has been established on Cimzia for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Cimzia was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]. <u>Note:</u> In cases when 130 days of the patient's prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Cimzia for at least 90 days AND the patient has been receiving Cimzia via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Cimzia). B) If the patient has met criterion 7Ai (the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria), but criterion 7Aii is not met: a request for one of the following Products may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria: i. Rheumatoid Arthritis: <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR.</u> ii. Juvenile Idiopathic Arthritis: <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Rinvoq LQ, Xeljanz tablets, or Xeljanz oral solution.</u> iii. Ankylosing Spondylitis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Taltz, Xeljanz tablets, or Xeljanz XR.</u> iv. Psoriatic Arthritis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz tablets, or Xeljanz XR.</u> v. Plaque Psoriasis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous (pen or syringe), Sotyktu, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous.</u> vi. Crohn's Disease: <u>Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Omvoh</u>
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	subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Tremfya subcutaneous, or Zymfentra. 8. Other Conditions. Approve <u>Cimzia</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria.
Enbrel	All Conditions. Approve <u>Enbrel</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Enbrel Prior Authorization Policy</i> criteria.
Humira (NDCs starting with 00074) Adalimumab-adbm Cyltezo Simlandi adalimumab-ryvk	All Conditions. Approve (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Adalimumab Products Prior Authorization Policy</i> criteria.
Simponi Subcutaneous	1. Rheumatoid Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Step 1 or Step 2 Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 2. Ankylosing Spondylitis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient has tried TWO of Enbrel, an adalimumab product, Rinvoq, Taltz, or Xeljanz/XR [documentation required].

	<u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: a request for a Step 1 or Step 2 Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Taltz, Xeljanz tablets, or Xeljanz XR</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 3. Psoriatic Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, an ustekinumab subcutaneous product, Taltz, Tremfya subcutaneous, or Xeljanz/XR [documentation required]. <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. B) If the patient has met criterion 3Ai (the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria), but criterion 3Aii is not met: a request for a Step 1 or Step 2 Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz tablets, or Xeljanz XR</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 4. Ulcerative Colitis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of an adalimumab product, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), an
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	ustekinumab subcutaneous product, Tremfya subcutaneous, Velsipity, or Zymfentra. <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, Yesintek. A trial of an infliximab intravenous product (e.g., Remicade, biosimilars), Omvoh intravenous, Skyrizi intravenous, Tremfya intravenous, or ustekinumab intravenous also counts. B) If the patient has met criterion 4Ai (the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria), but criterion 4Aii is not met: a request for a Preferred Product (<u>Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Tremfya subcutaneous, Velsipity, or Zymfentra</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 5. <u>Rheumatoid Arthritis, Ankylosing Spondylitis, Psoriatic Arthritis, or Ulcerative Colitis – Patient is Currently Receiving Simponi Subcutaneous or Aria.</u> A) Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a, b, c, d, e, <u>or</u> f): a) Patient has <u>Rheumatoid Arthritis</u> and has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. b) Patient has <u>Ankylosing Spondylitis</u> and has tried TWO of Enbrel, an adalimumab product, Rinvoq, Taltz, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. c) Patient has <u>Psoriatic Arthritis</u> and has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi
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	subcutaneous, an ustekinumab subcutaneous product, Taltz, Tremfya subcutaneous, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. d) Patient has <u>Ulcerative Colitis</u> and has tried one of an adalimumab product, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), an ustekinumab subcutaneous product, Tremfya subcutaneous, Velsipity, or Zymfentra; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, Yesintek. A trial of an infliximab intravenous product (e.g., Remicade, biosimilars), Omvoh intravenous, Skyrizi intravenous, Tremfya intravenous, or ustekinumab intravenous also counts. e) According to the prescriber, the patient has been established on Simponi Aria for at least 90 days; OR f) Patient has been established on Simponi subcutaneous for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Simponi subcutaneous was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]. <u>Note:</u> In cases when 130 days of the patient's prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Simponi subcutaneous for at least 90 days AND the patient has been receiving Simponi subcutaneous via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Simponi subcutaneous). B) If the patient has met criterion 5Ai (the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria), but criterion 5Aii is not met: a request for one of the following Products may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria: i. Rheumatoid Arthritis: <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira (NDCs starting with 00074).</u>
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	<u>adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR.</u> ii. Ankylosing Spondylitis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Taltz, Xeljanz tablets, or Xeljanz XR.</u> iii. Psoriatic Arthritis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz tablets, or Xeljanz XR.</u> iv. Ulcerative Colitis: <u>Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Tremfya subcutaneous, Velsipity, or Zymfentra.</u> 6. Other Conditions. Approve <u>Simponi subcutaneous</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria.
Zymfentra	All Conditions. Approve <u>Zymfentra</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Zymfentra Prior Authorization Policy</i> criteria.
Interleukin-6 Blockers	
Actemra Subcutaneous Tyenne Subcutaneous	1. Polyarticular Juvenile Idiopathic Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Tocilizumab Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a <u>or</u> b): a) Patient has tried one adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Enbrel, Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi Aria also counts. b) According to the prescriber, the patient has heart failure or a previously treated lymphoproliferative disorder. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Tocilizumab Subcutaneous Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 2. Rheumatoid Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Tocilizumab Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a <u>or</u> b):

	a) Patient has tried one adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, Enbrel, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. b) According to the prescriber, the patient has heart failure or a previously treated lymphoproliferative disorder. B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Tocilizumab Subcutaneous Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 3. <u>Polyarticular Juvenile Idiopathic Arthritis or Rheumatoid Arthritis – Patient is Currently Receiving Tocilizumab Subcutaneous or Intravenous.</u> A) Approve for 1 year if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Tocilizumab Subcutaneous Policy</i> criteria; AND ii. Patient meets ONE of the following (a, b, c, d, or e): a) Patient has <u>Polyarticular Juvenile Idiopathic Arthritis</u> and has tried one adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Enbrel, Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi Aria also counts. b) Patient has <u>Rheumatoid Arthritis</u> and has tried one adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, Enbrel, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. c) According to the prescriber, the patient has heart failure or a previously treated lymphoproliferative disorder; OR d) According to the prescriber, the patient has been established on tocilizumab intravenous for at least 90 days; OR e) Patient has been established on tocilizumab subcutaneous for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of tocilizumab subcutaneous was dispensed within the past 130 days [verification in prescription claims history required]</u>, or if claims history is not available, according to the prescriber <u>[verification by prescriber required]</u>. <u>Note:</u> In cases when 130 days of the patient’s prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving tocilizumab subcutaneous for at least
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	90 days AND the patient has been receiving tocilizumab subcutaneous via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to tocilizumab subcutaneous). B) If the patient has met criterion 3Ai (the standard <i>Inflammatory Conditions – Tocilizumab Subcutaneous Prior Authorization Policy</i> criteria), but criterion 3Aii is not met: a request for a Preferred Product may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria: i. Polyarticular Juvenile Idiopathic Arthritis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi.</u> ii. Rheumatoid Arthritis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi.</u> 4. All Other Conditions (including systemic juvenile idiopathic arthritis). Approve <u>tocilizumab subcutaneous</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Tocilizumab Subcutaneous Prior Authorization Policy</i> criteria.
Kevzara	1. Rheumatoid Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Kevzara Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a <u>or</u> b): a) Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of tocilizumab intravenous (Actemra intravenous, biosimilar), Cimzia, an infliximab product (e.g., Remicade, biosimilars), Orencia (intravenous or subcutaneous), or Simponi (Aria or subcutaneous) also counts [documentation required]. b) According to the prescriber, the patient has heart failure or a previously treated lymphoproliferative disorder. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Kevzara Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Step 1 or Step 2 Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.

	2. <u>Juvenile Idiopathic Arthritis/Juvenile Rheumatoid Arthritis – Initial Therapy.</u> A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Kevzara Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following conditions (a <u>or</u> b): a) Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq/Rinvoq LQ, or Xeljanz [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. A trial of Cimzia, a tocilizumab intravenous product (Actemra intravenous, biosimilar), Orencia intravenous or subcutaneous, an infliximab product (e.g., Remicade, biosimilars), or Simponi Aria also counts [documentation required]. b) According to the prescriber, the patient has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, OR a demyelinating disorder. B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions –Kevzara Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: a request for a Step 1 or Step 2 Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Rinvoq LQ, or Xeljanz tablets</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 3. <u>Juvenile Idiopathic Arthritis or Rheumatoid Arthritis – Patient is Currently Receiving Kevzara.</u> A) Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Kevzara Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a, b, c, <u>or</u> d): a) Patient has <u>Rheumatoid Arthritis</u> and has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively
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	counts as ONE product. A trial of tocilizumab intravenous (Actemra intravenous, biosimilar), Cimzia, an infliximab product (e.g., Remicade, biosimilars), Orenzia (intravenous or subcutaneous), or Simponi (Aria or subcutaneous) also counts [documentation required]. b) Patient has <u>Juvenile Idiopathic Arthritis</u> and has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, Rinvoq LQ, or Xeljanz [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. A trial of Cimzia, a tocilizumab intravenous product (Actemra intravenous, biosimilar), Orenzia intravenous or subcutaneous, an infliximab product (e.g., Remicade, biosimilars), or Simponi Aria also counts [documentation required]. c) According to the prescriber, the patient has heart failure or a previously treated lymphoproliferative disorder; OR d) Patient has been established on Kevzara for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Kevzara was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]. <u>Note:</u> In cases when 130 days of the patient's prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Kevzara for at least 90 days AND the patient has been receiving Kevzara via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Kevzara). A) If the patient has met criterion 3Ai (the standard <i>Inflammatory Conditions – Kevzara Prior Authorization Policy</i> criteria), but criterion 3Aii is not met: a request for one of the following Products may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. i. Rheumatoid Arthritis: <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR.</u> ii. Juvenile Idiopathic Arthritis: <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Rinvoq LQ, or Xeljanz tablets.</u> 3. Other Conditions. Approve <u>Kevzara</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the
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	standard <i>Inflammatory Conditions – Kevzara Prior Authorization Policy</i> criteria.
Interleukin-17 Blockers	
Bimzelx	1. <u>Ankylosing Spondylitis – Initial Therapy.</u> A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of Enbrel, an adalimumab product, or Taltz; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Preferred Product (<u>Enbrel, Cyltezo, adalimumab-adbm, Humira [NDCs starting with 00074], Simlandi, adalimumab-ryvk, or Taltz</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 2. <u>Hidradenitis Suppurativa – Initial Therapy.</u> A) Approve for 3 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria for hidradenitis suppurativa; AND ii. Patient has tried ONE of an adalimumab product or Cosentyx subcutaneous. <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: offer to review for a Preferred Product (<u>Cyltezo, adalimumab-adbm, Humira [NDCs starting with 00074], Simlandi, adalimumab-ryvk or Cosentyx subcutaneous</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria 3. <u>Non-Radiographic Spondyloarthritis (nr-axSpA) – Initial Therapy.</u> A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Bimzelx Authorization Policy</i> criteria; AND ii. Patient has tried one of Cimzia or Taltz. <u>Note:</u> A trial of Enbrel, an adalimumab product, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry.

B) If the patient has met criterion 3Ai (the standard *Inflammatory Conditions – Bimzelx Prior Authorization Policy* criteria), but criterion 2Aii is not met: a request for a Preferred Product (Cimzia or Taltz) may be reviewed using the respective standard *Inflammatory Conditions – Prior Authorization Policy* criteria.

4. Plaque Psoriasis – Initial Therapy.

A) Approve for 3 months if the patient meets BOTH of the following (i and ii):

- i.** Patient meets the standard *Inflammatory Conditions – Bimzelx Prior Authorization Policy* criteria for plaque psoriasis; AND
- ii.** Patient has tried ONE of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, an ustekinumab subcutaneous product, Taltz, or Tremfya subcutaneous.

Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek.

B) If the patient has met criterion 4Ai (the standard *Inflammatory Conditions – Bimzelx Prior Authorization Policy* criteria), but criterion 1Aii is not met: a request for a Preferred Product (Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous) may be reviewed using the respective standard *Inflammatory Conditions – Prior Authorization Policy* criteria.

5. Psoriatic Arthritis – Initial Therapy.

A) Approve for 6 months if the patient meets BOTH of the following (i and ii):

- i.** Patient meets the standard *Inflammatory Conditions – Bimzelx Prior Authorization Policy* criteria; AND
- ii.** Patient has tried one of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, an ustekinumab subcutaneous product, Taltz, or Tremfya subcutaneous; OR

Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts.

B) If the patient has met criterion 5Ai (the standard *Inflammatory Conditions – Bimzelx Prior Authorization Policy* criteria), but criterion 4Aii is not met: a request for a Preferred Product (Enbrel, Cyltezo, adalimumab-adbm, Humira [NDCs starting with 00074], Simlandi, adalimumab-ryvk, Otezla, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous)

	may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 6. <u>Ankylosing Spondylitis, Hidradenitis Suppurativa, nr-axSpA, Plaque Psoriasis or Psoriatic Arthritis – Patient is Currently Receiving Bimzelx.</u> A) Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a, b, c, d, <u>or</u> e): a) Patient has <u>Ankylosing Spondylitis</u> and has tried one of Enbrel, an adalimumab product, or Taltz; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. b) Patient has <u>Hidradenitis Suppurativa</u> and has tried one of an adalimumab product or Cosentyx subcutaneous; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. c) Patient has <u>nr-axSpA</u> and has tried one of Cimzia or Taltz; OR <u>Note:</u> A trial of Enbrel, an adalimumab product, an infliximab product (Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry d) Patient has <u>Plaque Psoriasis</u> and has tried ONE of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, an ustekinumab subcutaneous product, Taltz, or Tremfya subcutaneous; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. e) Patient has <u>Psoriatic Arthritis</u> and has tried one of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, an ustekinumab subcutaneous product, Taltz, or Tremfya subcutaneous; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio,
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	Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. f) Patient has been established on Bimzelx for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Bimzelx was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]. <u>Note:</u> In cases when 130 days of the patient's prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Bimzelx for at least 90 days AND the patient has been receiving Bimzelx via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Bimzelx). B) If the patient has met criterion 6Ai (the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria), but criterion 6Aii is not met: a request for one of the following Preferred Products may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. i. Ankylosing Spondylitis: <u>Enbrel, Cyltezo, adalimumab-adbm, adalimumab-ryvk, Humira [NDCs starting with 00074], Simlandi, or Taltz.</u> ii. Hidradenitis Suppurativa: <u>Cyltezo, adalimumab-adbm, adalimumab-ryvk, Humira [NDCs starting with 00074], Simlandi, or Cosentyx subcutaneous.</u> iii. nr-axSpA: <u>Cimzia or Taltz.</u> iv. Plaque Psoriasis: <u>Enbrel, Cyltezo, adalimumab-adbm, adalimumab-ryvk, Humira [NDCs starting with 00074], Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous.</u> v. Psoriatic Arthritis: <u>Enbrel, Cyltezo, adalimumab-adbm, adalimumab-ryvk, Humira [NDCs starting with 00074], Simlandi, Otezla, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous.</u> 7. Other Conditions. Approve Bimzelx (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria.
Cosentyx SC	1. Ankylosing Spondylitis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria; AND

	ii. Patient has tried TWO of Enbrel, an adalimumab product, Rinvoq, Taltz, or Xeljanz/XR [documentation required]. <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of Cimzia, an infliximab product (e.g. Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts [documentation required]. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Step 1 or Step 2 Product (<u>Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Enbrel, Rinvoq, Taltz, Xeljanz tablets, or Xeljanz XR</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 2. <u>Non-Radiographic Spondyloarthritis (nr-axSpA) – Initial Therapy.</u> A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient has tried TWO of Cimzia, Taltz, or Rinvoq [documentation required]. <u>Note:</u> A trial of Enbrel, an adalimumab product (e.g., Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry), an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts [documentation required]. A trial of multiple adalimumab products counts as ONE product. B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: a request for a Step 1 or Step 2 Product (<u>Cimzia, Taltz, or Rinvoq</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 3. <u>Plaque Psoriasis – Initial Therapy.</u> A) Approve for 3 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, an ustekinumab subcutaneous product, Taltz, or Tremfya <u>subcutaneous</u> [documentation required]. <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple
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	adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. B) If the patient has met criterion 3Ai (the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria), but criterion 3Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 4. Psoriatic Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a or b): a) Patient is ≥ 18 years of age AND has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, an ustekinumab subcutaneous product, Taltz, Tremfya <u>subcutaneous</u>, or Xeljanz/XR [documentation required]; OR b) Patient is < 18 years of age AND has tried ONE of Enbrel, Otezla, Rinvoq/Rinvoq LQ, or an ustekinumab subcutaneous product [documentation required]. <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (subcutaneous or Aria) also counts toward a trial of a TNFi [documentation required]. For a patient < 18 years of age, a trial of another TNFi counts towards a trial of Enbrel [documentation required]. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. B) If the patient has met criterion 4Ai (the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria), but criterion 4Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz, or Xeljanz XR</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.
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5. Ankylosing Spondylitis; nr-axSpA; Plaque Psoriasis; or Psoriatic Arthritis – Patient is Currently Receiving Cosentyx (Subcutaneous or Intravenous).

A) Approve for 1 year if the patient meets BOTH of the following (i and ii):

i. Patient meets the standard *Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy* criteria; AND

ii. Patient meets ONE of the following (a, b, c, d, e, f, or g):

a) Patient has Ankylosing Spondylitis and has tried TWO of Enbrel, an adalimumab product, Rinvoq, Taltz, or Xeljanz/XR
[documentation required]; OR

Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as **ONE** product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as **ONE** product. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts **[documentation required]**.

b) Patient has nr-axSpA and has tried TWO of Cimzia, Taltz, or Rinvoq **[documentation required]; OR**

Note: A trial of Enbrel, an adalimumab product (e.g., Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry), an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts **[documentation required]**. A trial of multiple adalimumab products counts as **ONE** product.

c) Patient has Plaque Psoriasis and has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, an ustekinumab subcutaneous product, Taltz, or Tremfya subcutaneous **[documentation required]; OR**

Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as **ONE** product. A trial of multiple ustekinumab products counts as ONE product.

d) Patient is ≥ 18 years of age with Psoriatic Arthritis and has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, an ustekinumab subcutaneous product, Taltz, Tremfya subcutaneous, or Xeljanz/XR
[documentation required]; OR

Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi,

	ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts [documentation required]. e) Patient is < 18 years of age with <u>Psoriatic Arthritis</u> and has tried ONE of Enbrel, Otezla, Rinvoq/Rinvoq LQ, or an ustekinumab subcutaneous product [documentation required]; OR <u>Note:</u> A trial of another TNFi counts towards a trial of Enbrel [documentation required]. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. f) According to the prescriber, the patient with Ankylosing Spondylitis, Non-Radiographic Spondyloarthritis, or Psoriatic Arthritis has been established on Cosentyx intravenous for at least 90 days; OR g) Patient has been established on Cosentyx subcutaneous for at least 90 days and prescription claims history indicates <u>at least a 90-day supply of Cosentyx SC was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]. <u>Note:</u> In cases when 130 days of the patient's prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Cosentyx SC for at least 90 days AND the patient has been receiving Cosentyx SC via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Cosentyx SC). B) If the patient has met criterion 5Ai (the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria), but criterion 5Aii is not met: a request for one of the following Products may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria: i. Ankylosing Spondylitis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Taltz, Xeljanz tablets, or Xeljanz XR.</u> ii. nr-axSpA: <u>Cimzia, Taltz, or Rinvoq.</u> iii. Plaque Psoriasis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous (pen or syringe), Sotyktu, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous.</u>
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	iv. Psoriatic Arthritis in a Patient $\geq$ 18 years of age: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz, or Xeljanz XR.</u> v. Psoriatic Arthritis in a Patient $<$ 18 years of age: <u>Enbrel, Otezla, Rinvoq, Rinvoq LQ, Stelara SC, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, or Yesintek subcutaneous.</u> 6. Other Conditions. Approve <u>Cosentyx SC</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria.
Siliq	1. Plaque Psoriasis – Initial Therapy. A) Approve for 3 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Siliq Prior Authorization Policy</i> criteria for plaque psoriasis; AND ii. Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, an ustekinumab subcutaneous product, Taltz, or Tremfya subcutaneous [documentation required]. <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Siliq Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 2. Plaque Psoriasis – Patient is Currently Receiving Siliq. A) Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Siliq Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a <u>or</u> b): a) Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, an ustekinumab subcutaneous product, Taltz, or Tremfya subcutaneous [documentation required]; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry.

	Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. b) Patient has been established on Siliq for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Siliq was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]. <u>Note:</u> In cases when 130 days of the patient's prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Siliq for at least 90 days AND the patient has been receiving Siliq via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Siliq). B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Siliq Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 3. Other Conditions. Approve <u>Siliq</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Siliq Prior Authorization Policy</i> criteria.
Taltz	All Conditions. Approve <u>Taltz</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Taltz Prior Authorization Policy</i> criteria.
Interleukin-23 Blockers	
Ilumya	1. Plaque Psoriasis – Initial Therapy. A) Approve for 3 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Ilumya Prior Authorization Policy</i> criteria; AND ii. Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, an ustekinumab subcutaneous product, Taltz, or Tremfya subcutaneous [documentation required]. <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product.

B) If the patient has met criterion 1Ai (the standard *Inflammatory Conditions – Ilumya Prior Authorization Policy* criteria), but criterion 1Aii is not met: a request for a Preferred Product (Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous) may be reviewed using the respective standard *Inflammatory Conditions – Prior Authorization Policy* criteria.

2. Plaque Psoriasis – Patient is Currently Receiving Ilumya.

A) Approve for 1 year if the patient meets BOTH of the following (i and ii):

i. Patient meets the standard *Inflammatory Conditions – Ilumya Prior Authorization Policy* criteria; AND

ii. Patient meets ONE of the following (a or b):

a) Patient has plaque psoriasis and has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, ustekinumab subcutaneous product, Taltz, or Tremfya subcutaneous **[documentation required]**; OR
Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as **ONE** product. A trial of multiple ustekinumab products counts as ONE product.

b) Patient has been established on Ilumya for at least 90 days and prescription claims history indicates at least a 90-day supply of Ilumya was dispensed within the past 130 days **[verification in prescription claims history required]**, or if claims history is not available, according to the prescriber **[verification by prescriber required]**.

Note: In cases when 130 days of the patient's prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Ilumya for at least 90 days AND the patient has been receiving Ilumya via paid claims (e.g., patient has not been receiving samples or coupons or other types of waivers in order to obtain access to Ilumya).

B) If the patient has met criterion 2Ai (the standard *Inflammatory Conditions – Ilumya Prior Authorization Policy* criteria), but criterion 2Aii is not met: a request for a Preferred Product (Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous) may be reviewed using the respective standard *Inflammatory Conditions – Prior Authorization Policy* criteria.

3. Other Conditions. Approve Ilumya (initial therapy for a duration as directed or 1 year for a patient continuing therapy) if the patient meets the

	standard <i>Inflammatory Conditions – Ilumya Prior Authorization Policy</i> criteria.
Omvoh SC	All Conditions. Approve <u>Omvoh subcutaneous</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Omvoh Subcutaneous Prior Authorization Policy</i> criteria.
Skyrizi Subcutaneous	All Conditions. Approve <u>Skyrizi subcutaneous</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Skyrizi Subcutaneous Prior Authorization Policy</i> criteria.
Tremfya	All Conditions. Approve <u>Tremfya subcutaneous</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Tremfya Subcutaneous Prior Authorization Policy</i> criteria.
IL-12/23 Blocker	
Stelara Subcutaneous Selarsdi Subcutaneous Ustekinumab-ttwe Subcutaneous Yesintek Subcutaneous	All Conditions. Approve <u>Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, or Yesintek subcutaneous</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Ustekinumab Subcutaneous Prior Authorization Policy</i> criteria.
Integrin Receptor Antagonist	
Entyvio SC	<u>Applies only when Entyvio SC is requested for coverage under the Prescription Drug Benefit</u> 1. <u>Crohn’s Disease – Initial Therapy.</u> A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Entyvio Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a <u>or</u> b): a) Patient has tried TWO of an adalimumab product, Skyrizi subcutaneous, Tremfya subcutaneous, an ustekinumab subcutaneous product, Zymfentra, Omvoh subcutaneous, Cimzia, or Rinvoq [documentation required]; OR Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. A trial of an infliximab intravenous product (e.g., Remicade, biosimilars), Skyrizi intravenous, Omvoh intravenous, Tremfya intravenous, or

	ustekinumab intravenous also counts [documentation required]. b) According to the prescriber, the patient has already started on or is currently undergoing induction therapy with Entvyio IV. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Entvyio Subcutaneous Prior Authorization Policy</i> criteria), but criterion 1Aii is not met, a request for a Step 1 or Step 2 Product (<u>Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Skyrizi subcutaneous, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Rinvoq, Cimzia, Omvoh subcutaneous, Tremfya subcutaneous, or Zymfentra</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 2. <u>Ulcerative Colitis – Initial Therapy.</u> A) Approve for 6 months if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Entvyio Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a or b): a) Patient has tried TWO of an adalimumab product, Skyrizi subcutaneous, an ustekinumab subcutaneous product, Zymfentra, Omvoh subcutaneous, Rinvoq, Simponi subcutaneous, Tremfya subcutaneous, Velsipity, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of an infliximab intravenous product (e.g., Remicade, biosimilars), Omvoh intravenous, Skyrizi intravenous, ustekinumab intravenous, or Tremfya intravenous also counts [documentation required]. b) According to the prescriber, the patient has already started on or is currently undergoing induction therapy with Entvyio IV. B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Entvyio Subcutaneous Prior Authorization Policy</i> criteria), but criterion 2Aii is not met, a request for a Step 1 or Step 2 Product (<u>Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Omvoh subcutaneous, Rinvoq, Simponi SC, Skyrizi subcutaneous (on-body injector), Xeljanz/XR, Tremfya subcutaneous, Velsipity, or Zymfentra</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 3. <u>Crohn’s Disease and Ulcerative Colitis – Patient is Currently Receiving Entvyio Subcutaneous or Intravenous.</u>
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	A) Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Entyvio Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following conditions (a, b, c, <u>or</u> d): a) Patient has <u>Crohn’s Disease</u> and has tried TWO of an adalimumab product, Skyrizi subcutaneous, Tremfya subcutaneous, an ustekinumab subcutaneous product, Zymfentra, Omvoh subcutaneous, Cimzia, or Rinvoq [documentation required]; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. A trial of an infliximab intravenous product (e.g., Remicade, biosimilars), Skyrizi intravenous, Omvoh intravenous, Tremfya intravenous, or ustekinumab intravenous also counts [documentation required]. b) Patient has <u>Ulcerative Colitis</u> and has tried TWO of an adalimumab product, Skyrizi subcutaneous, an ustekinumab subcutaneous product, Tremfya subcutaneous, Zymfentra, Omvoh subcutaneous, Rinvoq, Simponi subcutaneous, Velsipity, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-aacf, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of an infliximab intravenous product (e.g., Remicade, biosimilars), Omvoh intravenous, Skyrizi intravenous, ustekinumabintravenous, or Tremfya intravenous also counts [documentation required]. c) According to the prescriber, the patient has been established on Entyvio intravenous for at least 90 days; OR d) Patient has been established on Entyvio subcutaneous for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Entyvio subcutaneous was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]. <u>Note:</u> In cases where 130 days of the patient’s prescription claim history file is unavailable to be verified, an exception to
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	this requirement is allowed if the prescriber has verified that the patient has been receiving Entyvio subcutaneous for at least 90 days AND the patient has been receiving Entyvio subcutaneous via paid claims (e.g., patient has not been receiving samples or coupons or other types of waivers in order to obtain access to Entyvio subcutaneous). B) If the patient has met criterion 3Ai (the standard <i>Inflammatory Conditions – Entyvio Subcutaneous Prior Authorization Policy</i> criteria), but criterion 3Aii is not met, a request for one of the following Products may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. i. Crohn’s Disease: <u>Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Skyrizi subcutaneous, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Rinvoq, Cimzia, Omvoh subcutaneous, Tremfya subcutaneous, or Zymfentra.</u> ii. Ulcerative Colitis: <u>Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Tremfya subcutaneous, Omvoh subcutaneous, Rinvoq, Simponi SC, Xeljanz/XR, Velsipity, or Zymfentra.</u> 4. Other Conditions. Approve <u>Entyvio subcutaneous</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Entyvio Subcutaneous Prior Authorization Policy</i> criteria.
Interleukin-1 Blocker	
Kineret	1. Rheumatoid Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Kineret Prior Authorization Policy</i> criteria; AND ii. Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [documentation required]. <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of tocilizumab intravenous (Actemra intravenous, biosimilar), Cimzia, Orencia (subcutaneous or intravenous), an infliximab product (e.g., Remicade, biosimilar), Kevzara, or Simponi (Aria or subcutaneous) also counts [documentation required]. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Kineret Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Step 1 or Step 2 Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira [NDCs starting</u>

	with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 2. Rheumatoid Arthritis – Patient is Currently Receiving Kineret. A) Approve for 1 year if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Kineret Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a or b): a) Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of tocilizumab intravenous (Actemra intravenous, biosimilar), Cimzia, Orencia (subcutaneous or intravenous), an infliximab product (e.g., Remicade, biosimilar), Kevzara, or Simponi (Aria or subcutaneous) also counts [documentation required]. b) Patient has been established on Kineret at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Kineret was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]. <u>Note:</u> In cases when 130 days of the patient’s prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Kineret for at least 90 days AND the patient has been receiving Kineret via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Kineret). B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Kineret Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: a request for a Step 1 or Step 2 Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 3. Other Conditions. Approve Kineret (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Kineret Prior Authorization Policy</i> criteria. <u>Note:</u> This includes Cryopyrin-Associated Periodic Syndromes (CAPS), Systemic Juvenile Idiopathic Arthritis.
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T-Cell Costimulation Modulator	
Orencia Subcutaneous	1. Rheumatoid Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Orencia Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a <u>or</u> b): a) Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of tocilizumab intravenous (Actemra intravenous, biosimilar), Cimzia, an infliximab product (e.g., Remicade, biosimilars), Kevzara, or Simponi (Aria or subcutaneous) also counts [documentation required]. b) According to the prescriber, the patient has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, OR a demyelinating disorder. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Orencia Subcutaneous Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Step 1 or Step 2 Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 2. Juvenile Idiopathic Arthritis/Juvenile Rheumatoid Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Orencia Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a <u>or</u> b): a) Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq/Rinvoq LQ, or Xeljanz [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz tablets and Xeljanz oral solution)

	collectively counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. A trial of Cimzia, tocilizumab intravenous (Actemra intravenous, biosimilar), Kevzara, Orencia intravenous, an infliximab product (e.g., Remicade, biosimilar), or Simponi Aria also counts [documentation required]. b) According to the prescriber, the patient has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, OR a demyelinating disorder. B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Orencia Subcutaneous Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: a request for a Step 1 or Step 2 Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Rinvoq LQ, Xeljanz tablets, or Xeljanz oral solution</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 3. Psoriatic Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Orencia Subcutaneous Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a, b, or c): a) Patient is ≥ 18 years of age AND has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, an ustekinumab subcutaneous product, Taltz, Tremfya subcutaneous, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), Simponi (Aria or subcutaneous), Cosentyx, or Bimzelx also counts [documentation required]. b) Patient is < 18 years of age AND has tried ONE of Enbrel, Otezla, Rinvoq/Rinvoq LQ, or an ustekinumab subcutaneous product [documentation required]; OR <u>Note:</u> A trial of another TNFi counts towards a trial of Enbrel [documentation required]. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product.
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	c) According to the prescriber, the patient has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, OR a demyelinating disorder. B) If the patient has met criterion 3Ai (the standard <i>Inflammatory Conditions – Orencia Subcutaneous Prior Authorization Policy</i> criteria), but criterion 3Aii is not met: a request for a Step 1 or Step 2a Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz tablets, or Xeljanz XR</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 4. <u>Rheumatoid Arthritis, Juvenile Idiopathic Arthritis, or Psoriatic Arthritis – Patient is Currently Receiving Orencia (Subcutaneous or Intravenous).</u> A) Approve for 1 year if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Orencia Subcutaneous Policy</i> criteria; AND ii. Patient meets ONE of the following (a, b, c, d, e, f, or g): a) Patient has <u>Rheumatoid Arthritis</u> and has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of tocilizumab intravenous (Actemra intravenous, biosimilar), Cimzia, an infliximab product (e.g., Remicade, biosimilars), Kevzara, or Simponi (Aria or subcutaneous) also counts [documentation required]. b) Patient has <u>Juvenile Idiopathic Arthritis</u> and has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq/Rinvoq LQ, or Xeljanz tablets or oral solution [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz tablets and Xeljanz oral solution) collectively counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. A trial of Cimzia, tocilizumab intravenous (Actemra intravenous, biosimilar), Kevzara, Orencia intravenous,
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	an infliximab product (e.g., Remicade, biosimilars), or Simponi Aria also counts [documentation required]. c) Patient is ≥ 18 years of age with <u>Psoriatic Arthritis</u> AND has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, an ustekinumab subcutaneous product, Taltz, Tremfya subcutaneous, or Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. A trial of multiple adalimumab products counts as ONE product. A trial of multiple ustekinumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), Simponi (Aria or subcutaneous), Cosentyx, or Bimzelx also counts [documentation required]. d) Patient is < 18 years of age with <u>Psoriatic Arthritis</u> AND has tried ONE of Enbrel, Otezla, Rinvoq/Rinvoq LQ, or an ustekinumab subcutaneous product [documentation required]; OR <u>Note:</u> A trial of another TNFi counts towards a trial of Enbrel [documentation required]. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as ONE product. Examples of ustekinumab products include Stelara/ustekinumab, Imuldosa, Otulfi, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek. e) According to the prescriber, the patient has been established on Orencia intravenous for at least 90 days; OR f) According to the prescriber, the patient has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, OR a demyelinating disorder; OR g) Patient has been established on Orencia subcutaneous for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Orencia subcutaneous was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]. <u>Note:</u> In cases when 130 days of the patient's prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Orencia subcutaneous for at least 90 days AND the patient has been receiving Orencia subcutaneous via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Orencia subcutaneous). B) If the patient has met criterion 4Ai (the standard <i>Inflammatory Conditions – Orencia Subcutaneous Prior Authorization Policy</i> criteria),
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	but criterion 4Aii is not met, a request for one of the following Products may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. i. Rheumatoid Arthritis: <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR.</u> ii. Juvenile Idiopathic Arthritis: <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Rinvoq LQ, Xeljanz tablets, or Xeljanz oral solution.</u> iii. Psoriatic Arthritis in a Patient ≥ 18 Years of Age: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz tablets, or Xeljanz XR.</u> iv. Psoriatic Arthritis in a Patient < 18 Years of Age: <u>Enbrel, Otezla, Rinvoq, Rinvoq LQ, Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, or Yesintek subcutaneous.</u> 5. Other Conditions. Approve <u>Orencia subcutaneous</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Orencia Subcutaneous Prior Authorization Policy</i> criteria.
Janus Kinases Inhibitors	
Olumiant	1. Rheumatoid Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Olumiant Prior Authorization Policy</i> criteria; AND ii. Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [documentation required]. <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of tocilizumab intravenous (Actemra intravenous, biosimilar), Cimzia, an infliximab product (e.g., Remicade, biosimilars), Kevzara, Orencia (intravenous or subcutaneous), or Simponi (Aria or subcutaneous) also counts [documentation required]. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Olumiant Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Step 1 or Step 2 Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira [NDCs starting</u>

	with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 2. Rheumatoid Arthritis – Patient is Currently Receiving Olumiant. A) Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Olumiant Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a <u>or</u> b): a) Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, and Xeljanz/XR [documentation required]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as ONE product. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as ONE product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as ONE product. A trial of tocilizumab intravenous (Actemra intravenous, biosimilar), Cimzia, an infliximab product (e.g., Remicade, biosimilars), Kevzara, Orencia (intravenous or subcutaneous), or Simponi (Aria or subcutaneous) also counts [documentation required]. b) Patient has been established on Olumiant for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Olumiant was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]. <u>Note:</u> In cases when 130 days of the patient’s prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Olumiant for at least 90 days AND the patient has been receiving Olumiant via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Olumiant). B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Olumiant Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: a request for a Step 1 or Step 2 Product (Actemra subcutaneous, Tyenne subcutaneous, Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 3. Other Conditions. Approve <u>Olumiant</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Olumiant Prior Authorization Policy</i> criteria.
Rinvoq	1. Ankylosing Spondylitis – Initial Therapy.

	A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, or Taltz</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 2. Crohn's Disease – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND ii. Patient has tried one adalimumab product. <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of an infliximab product (e.g., Remicade, biosimilars; Zymfentra) or Cimzia also counts. B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: a request for a Preferred Product (<u>Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous [on-body injector], Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Tremfya subcutaneous, or Zymfentra</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 3. Juvenile Idiopathic Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars) or Simponi Aria also counts. B) If the patient has met criterion 3Ai (the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria), but criterion 3Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira</u>
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	[NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 4. <u>Non-Radiographic Spondyloarthritis (nr-axSpA) – Initial Therapy.</u> A) Approve for 6 months if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND ii. Patient has tried Cimzia. <u>Note:</u> A trial of Enbrel, an adalimumab product, an infliximab product (Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. B) If the patient has met criterion 4Ai (the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria), but criterion 4Aii is not met: a request for a Preferred Product (<u>Cimzia or Taltz</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 5. <u>Rheumatoid Arthritis – Initial Therapy.</u> A) Approve for 6 months if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. B) If the patient has met criterion 5Ai (the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria), but criterion 5Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 6. <u>Psoriatic Arthritis – Initial Therapy.</u> A) Approve for 6 months if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. B) If the patient has met criterion 6Ai (the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria), but criterion 6Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira</u>
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[NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous) may be reviewed using the respective standard *Inflammatory Conditions Prior Authorization Policy* criteria.

7. Ulcerative Colitis – Initial Therapy.

- A)** Approve for 6 months if the patient meets BOTH of the following (i and ii):
- i.** Patient meets the standard *Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy* criteria; AND
 - ii.** Patient has tried one adalimumab product.
Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of an infliximab product (e.g., Remicade, biosimilars; Zymfentra) or Simponi subcutaneous also counts.
- B)** If the patient has met criterion 7Ai (the standard *Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy* criteria), but criterion 7Aii is not met: a request for a Preferred Product (Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Tremfya subcutaneous, Velsipity, or Zymfentra) may be reviewed using the respective standard *Inflammatory Conditions Prior Authorization Policy* criteria.

8. Ankylosing Spondylitis, Crohn's Disease, nr-axSpA, Rheumatoid Arthritis, Psoriatic Arthritis, or Ulcerative Colitis – Patient is Currently Receiving Rinvoq.

- A)** Approve for 1 year if the patient meets BOTH of the following (i and ii):
- i.** Patient meets the standard *Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy* criteria; AND
 - ii.** Patient meets ONE of the following (a, b, c, d, e, f, g, or h):
 - a)** Patient has Ankylosing Spondylitis and has tried one of Enbrel or an adalimumab product; OR
Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts.
 - b)** Patient has Crohn's Disease and has tried one adalimumab product; OR
Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of an infliximab product (e.g., Remicade, biosimilars; Zymfentra) or Cimzia also counts.
 - c)** Patient has Juvenile Idiopathic Arthritis and has tried ONE of Enbrel or an adalimumab product; OR

	<u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars) or Simponi Aria also counts. d) Patient has <u>nr-axSpA</u> and has tried Cimzia; OR <u>Note:</u> A trial of Enbrel, an adalimumab product, an infliximab product (Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. e) Patient has <u>Rheumatoid Arthritis</u> and has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. f) Patient has <u>Psoriatic Arthritis</u> and has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. g) Patient has <u>Ulcerative Colitis</u> and has tried one adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of an infliximab product (e.g., Remicade, biosimilars; Zymfentra) or Simponi subcutaneous also counts. h) Patient has been established on Rinvoq for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Rinvoq was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]. <u>Note:</u> In cases when 130 days of the patient's prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Rinvoq for at least 90 days AND the patient has been receiving Rinvoq via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Rinvoq).
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	B) If the patient has met criterion 8Ai (the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria), but criterion 8Aii is not met: a request for one of the following Products may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria: i. Ankylosing Spondylitis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, or Taltz.</u> ii. Crohn’s Disease: <u>Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Tremfya subcutaneous, or Zymfentra.</u> iii. Juvenile Idiopathic Arthritis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi.</u> iv. nr-axSpA: <u>Cimzia or Taltz.</u> v. Rheumatoid Arthritis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi.</u> vi. Psoriatic Arthritis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous.</u> vii. Ulcerative Colitis: <u>Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Tremfya subcutaneous, Velsipity, or Zymfentra.</u> 9. All Other Conditions. Approve <u>Rinvoq</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria.
Rinvoq LQ	1. Juvenile Idiopathic Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars) or Simponi Aria also counts. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 2. Psoriatic Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):

	i. Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 3. <u>Juvenile Idiopathic Arthritis or Psoriatic Arthritis – Patient is Currently Receiving Rinvoq/LQ.</u> A) Approve for 1 year if the patient meets BOTH of the following (i and ii): i. Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following conditions (a, b, or c): a) Patient has <u>Juvenile Idiopathic Arthritis</u> and has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars) or Simponi Aria also counts. b) Patient has <u>Psoriatic Arthritis</u> and has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. c) Patient has been established on Rinvoq/LQ for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Rinvoq/LQ was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]. <u>Note:</u> In cases when 130 days of the patient’s prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Rinvoq/LQ for at least 90 days AND the patient has been receiving Rinvoq/LQ via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Rinvoq/LQ).
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	B) If the patient has met criterion 3Ai (the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria but criterion 3Aii is not met: a request for one of the following Products may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria: i. Juvenile Idiopathic Arthritis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi.</u> ii. Psoriatic Arthritis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous.</u> 4. Other Conditions. Approve <u>Rinvoq LQ</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria.
Xeljanz tablets, Xeljanz XR tablets	1. Ankylosing Spondylitis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, or Taltz</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 2. Rheumatoid Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 3. Juvenile Idiopathic Arthritis – Initial Therapy.

	A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars) or Simponi Aria also counts. B) If the patient has met criterion 3Ai (the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria), but criterion 3Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 4. Psoriatic Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts. B) If the patient has met criterion 4Ai (the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria), but criterion 4Aii is not met: a request for a Step 1 Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous</u>) may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria. 5. Ulcerative Colitis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND ii. Patient has tried one adalimumab product. <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of an infliximab product (e.g., Remicade, biosimilars; Zymfentra) or Simponi subcutaneous also counts. B) If the patient has met criterion 5Ai (the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria), but criterion 5Aii is not met: a request for a Preferred Product (<u>Humira [NDCs</u>
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starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Tremfya subcutaneous, Velsipity, or Zymfentra) may be reviewed using the respective standard *Inflammatory Conditions Prior Authorization Policy* criteria.

6. Ankylosing Spondylitis, Rheumatoid Arthritis, Juvenile Idiopathic Arthritis, Psoriatic Arthritis, or Ulcerative Colitis – Patient is Currently Receiving Xeljanz/XR.

- A)** Approve for 1 year if the patient meets BOTH of the following (i and ii):
- i.** Patient meets the standard *Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy* criteria; AND
 - ii.** Patient meets ONE of the following (a, b, c, d, e, or f):
 - a)** Patient has Ankylosing Spondylitis and has tried one of Enbrel or an adalimumab product; OR
Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts.
 - b)** Patient has Rheumatoid Arthritis and has tried one of Enbrel or an adalimumab product; OR
Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts.
 - c)** Patient has Juvenile Idiopathic Arthritis and has tried one of Enbrel or an adalimumab product; OR
Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars) or Simponi Aria also counts.
 - d)** Patient has Psoriatic Arthritis and has tried one of Enbrel or an adalimumab product; OR
Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts.
 - e)** Patient has Ulcerative Colitis and has tried one adalimumab product; OR
Note: Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of an infliximab product (e.g., Remicade, biosimilars; Zymfentra) or Simponi subcutaneous also counts.

	f) Patient has been established on Xeljanz/XR for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Xeljanz/XR was dispensed within the past 130 days</u> [verification in prescription claims history required], or if claims history is not available, according to the prescriber [verification by prescriber required]; OR <u>Note:</u> In cases when 130 days of the patient's prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Xeljanz/XR for at least 90 days AND the patient has been receiving Xeljanz/XR via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Xeljanz/XR). B) If the patient has met criterion 6Ai (the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria but criterion 6Aii is not met: a request for one of the following Products may be reviewed using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria: i. Ankylosing Spondylitis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, or Taltz.</u> ii. Rheumatoid Arthritis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi.</u> iii. Juvenile Idiopathic Arthritis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi.</u> iv. Psoriatic Arthritis: <u>Enbrel, Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Taltz, or Tremfya subcutaneous.</u> v. Ulcerative Colitis: <u>Humira (NDCs starting with 00074), adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, Yesintek subcutaneous, Tremfya subcutaneous, Velsipity, or Zymfentra.</u> 7. Other Conditions. Approve <u>Xeljanz/XR</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria.
Xeljanz oral solution	1. Juvenile Idiopathic Arthritis – Initial Therapy. A) Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND ii. Patient has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars) or Simponi Aria also counts.

	B) If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 2. Juvenile Idiopathic Arthritis – Patient is Currently Receiving Xeljanz. A) Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii): i. Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND ii. Patient meets ONE of the following (a <u>or</u> b): a) Patient has <u>Juvenile Idiopathic Arthritis</u> and has tried one of Enbrel or an adalimumab product; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi Aria also counts. b) Patient has been established on Xeljanz for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Xeljanz was dispensed within the past 130 days [verification in prescription claims history required]</u>, or if claims history is not available, according to the prescriber [verification by prescriber required]; OR <u>Note:</u> In cases when 130 days of the patient’s prescription claim history file is unavailable to be verified, an exception to this requirement is allowed if the prescriber has verified that the patient has been receiving Xeljanz for at least 90 days AND the patient has been receiving Xeljanz via paid claims (e.g., patient has <u>not</u> been receiving samples or coupons or other types of waivers in order to obtain access to Xeljanz). B) If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria but criterion 2Aii is not met: a request for a Preferred Product (<u>Enbrel, Humira [NDCs starting with 00074], adalimumab-adbm, Cyltezo, adalimumab-ryvk, or Simlandi</u>) may be reviewed using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria. 3. Other Conditions. Approve <u>Xeljanz oral solution</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria.
Phosphodiesterase Type 4 Inhibitor	
Otezla	All Conditions. Approve <u>Otezla</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Otezla Prior Authorization Policy</i> criteria.
Sphingosine 1-Phosphate Receptor Modulator	
Velsipity	1. All Conditions. Approve <u>Velsipity</u> if the patient meets the standard <i>Inflammatory Conditions – Velsipity Prior Authorization Policy</i> criteria.
Zeposia	All Conditions. Approve <u>Zeposia</u> if the patient meets the standard <i>Multiple Sclerosis and Ulcerative Colitis – Zeposia Preferred Specialty Management</i>

	<i>Policy: Standard/ Performance, Value/Advantage, Total Savings Prescription Drug Lists Policy</i> criteria.
Tyrosine Kinase 2 Inhibitor	
Sotyktu	All Conditions. Approve <u>Sotyktu</u> (initial therapy for a duration as directed or <u>1 year</u> for a patient continuing therapy) if the patient meets the standard <i>Inflammatory Conditions – Sotyktu Prior Authorization Policy</i> criteria.

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Revision Details

Type of Revision	Summary of Changes	Date
New	New policy Legacy Drug List Plans criteria were relocated to a new policy. The criteria were previously located within Inflammatory Conditions Preferred Specialty Management Policy for Employer Plans (PSM001). The following updates were made to the previous criteria. Hyrimoz: Throughout the policy, NDCs starting with 61314 were removed from the Preferred Products. A previous trial of these NDCs counts towards a trial of an adalimumab product. Tremfya Subcutaneous: For Ulcerative Colitis, Tremfya subcutaneous was added as a Preferred Product. Cimzia: For Rheumatoid Arthritis, Ankylosing Spondylitis, Psoriatic Arthritis, Plaque Psoriasis, and Crohn's Disease, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. For Juvenile Idiopathic Arthritis, Cimzia was added to Step 3a. Documentation of a trial of two Step 1 or 2a Products is required. A trial of a tocilizumab intravenous product (Actemra intravenous, biosimilar), Kevzara, Orencia intravenous or subcutaneous, an infliximab product (e.g., Remicade, biosimilars), or Simponi Aria also counts. For Psoriatic Arthritis and Plaque Psoriasis, it was clarified that Tremfya is the subcutaneous formulation. Simponi Subcutaneous: For Rheumatoid Arthritis, Ankylosing Spondylitis, Psoriatic Arthritis, and Ulcerative Colitis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. For Psoriatic Arthritis, it was	01/01/2025

	clarified that Tremfya is the subcutaneous formulation. For Ulcerative Colitis, Tremfya subcutaneous was added as Preferred Products. Actemra Subcutaneous and Tyenne Subcutaneous: For Rheumatoid Arthritis and Polyarticular Juvenile Idiopathic Arthritis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. For Polyarticular Juvenile Idiopathic Arthritis, Cimzia was added as an agent that counts towards a trial of a Preferred Product. Kevzara: For Rheumatoid Arthritis and Juvenile Idiopathic Arthritis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. For Juvenile Idiopathic Arthritis, Cimzia was added as an agent that counts towards a trial of a Preferred Product. Bimzelx: For Ankylosing Spondylitis, Non-Radiographic Spondyloarthritis, and Psoriatic Arthritis, Bimzelx was added to Step 2a and requests are directed to a trial of one Step 1 Product. For Plaque Psoriasis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. Cosentyx Subcutaneous: For Ankylosing Spondylitis, Psoriatic Arthritis, and Plaque Psoriasis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. For Psoriatic Arthritis and Plaque Psoriasis, it was clarified that Tremfya is the subcutaneous formulation. Siliq: For Plaque Psoriasis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products, and it was clarified that Tremfya is the subcutaneous formulation. Ilumya: For Plaque Psoriasis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products, and it was clarified that Tremfya is the subcutaneous formulation. Entyvio Subcutaneous: For Crohn's Disease and Ulcerative Colitis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. For Ulcerative Colitis, Tremfya subcutaneous was added as a Preferred Product; a previous trial of Tremfya intravenous also counts. Kineret: For Rheumatoid Arthritis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. Orencia Subcutaneous: For Rheumatoid Arthritis, Juvenile Idiopathic Arthritis, and Psoriatic Arthritis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. For Juvenile Idiopathic Arthritis, Cimzia was	
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	added as an agent that counts towards a trial of a Preferred Product. For Psoriatic Arthritis, it was clarified that Tremfya is the subcutaneous formulation; for a patient ≥ 18 years of age, Cosentyx and Bimzelx were added as agents that count towards a trial of a Preferred Product. Olumiant: For Rheumatoid Arthritis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. Rinvoq: For Rheumatoid Arthritis, Ankylosing Spondylitis, Juvenile Idiopathic Arthritis, Psoriatic Arthritis, Crohn's Disease, and Ulcerative Colitis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. For Juvenile Idiopathic Arthritis, Cimzia was added as an agent that counts towards a trial of a Preferred Product. For Psoriatic Arthritis, it was clarified that Tremfya is the subcutaneous formulation. For Ulcerative Colitis, Tremfya subcutaneous was added as a Preferred Product. Rinvoq LQ: For Juvenile Idiopathic Arthritis and Psoriatic Arthritis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. For Juvenile Idiopathic Arthritis, Cimzia was added as an agent that counts towards a trial of a Preferred Product. For Psoriatic Arthritis, it was clarified that Tremfya is the subcutaneous formulation. Xeljanz/Xeljanz XR: For Rheumatoid Arthritis, Ankylosing Spondylitis, Juvenile Idiopathic Arthritis (Xeljanz tablets only), Psoriatic Arthritis, and Ulcerative Colitis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. For Juvenile Idiopathic Arthritis, Cimzia was added as an agent that counts towards a trial of a Preferred Product. For Psoriatic Arthritis, it was clarified that Tremfya is the subcutaneous formulation. For Ulcerative Colitis, Tremfya subcutaneous was added as a Preferred Product. Xeljanz Oral Solution: For Juvenile Idiopathic Arthritis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products. For Juvenile Idiopathic Arthritis, Cimzia was added as an agent that counts towards a trial of a Preferred Product. Velsipity: For Ulcerative Colitis, Hyrimoz (NDCs starting with 61314) was removed from the Preferred Products; Tremfya subcutaneous was added as a Preferred Product; a previous trial of Tremfya intravenous also counts.	
Selected Revision	Hidradenitis Suppurativa was added as a targeted indication in this policy. Adalimumab	02/01/2025

	products (Humira [NDCs starting with 00074], Cyltezo/adalimumab-adbm, adalimumab-adaz, Simlandi/adalimumab-ryvk) and Cosentyx subcutaneous are Preferred Products for Hidradenitis Suppurativa; Bimzelx was added to Step 2b and is directed to a trial of one Preferred Product.	
Selected Revision	Velsipity: For Ulcerative Colitis, Velsipity was added as a Preferred Product. Simponi Subcutaneous: For Ulcerative Colitis, Velsipity was added as a Preferred Product. Rinvoq: For Ulcerative Colitis, Velsipity was added as a Preferred Product. Xeljanz/XR: For Ulcerative Colitis, Velsipity was added as a Preferred Product. Omvoh Subcutaneous: For Ulcerative Colitis, Velsipity was added as a Preferred Product. For Crohn's Disease, Omvoh subcutaneous was added as a Step 2a Non-Preferred Product and is directed to a trial of one Step 1 Product. Entyvio Subcutaneous: For Ulcerative Colitis, Velsipity was added as a Preferred Product. For Crohn's Disease, Omvoh subcutaneous was added as a Step 2a Non-Preferred Product.	03/15/2025
Selected Revision	Omvoh Subcutaneous: For Crohn's Disease and Ulcerative Colitis, Omvoh subcutaneous was moved from Step 2a to Preferred (Step 1). Cimzia: For Crohn's Disease, Omvoh subcutaneous was added as a Preferred Product. Simponi Subcutaneous: For Ulcerative Colitis, Omvoh subcutaneous was added as a Preferred Product. Rinvoq: For Crohn's Disease and Ulcerative Colitis, Omvoh subcutaneous was added as a Preferred Product. Xeljanz/Xeljanz XR: For Ulcerative Colitis, Omvoh subcutaneous was added as a Preferred Product.	04/01/2025
Selected Revision	For Psoriatic Arthritis, Plaque Psoriasis, Crohn's Disease, and Ulcerative Colitis, Selarsdi subcutaneous, ustekinumab-ttwe subcutaneous, and Yesintek subcutaneous were added as Preferred ustekinumab subcutaneous products. The criteria for the following Non-Preferred Products were updated to include these Preferred ustekinumab products: Ilumya, Siliq, Entyvio subcutaneous, Rinvoq LQ, Rinvoq, Xeljanz, Bimzelx, Cimzia, Simponi subcutaneous, Cosentyx subcutaneous, and Orencia subcutaneous. Throughout the policy,	04/15/2025

	a note was added indicating that a trial of multiple ustekinumab products counts as one product and examples of ustekinumab products were added including Stelara, Wezlana, Otufli, Pyzchiva, ustekinumab-ttwe, Selarsdi, Steqeyma, and Yesintek. For Crohn's Disease and Ulcerative Colitis, the note that refers to a previous trial of Stelara intravenous was changed to more generally refer to an intravenous ustekinumab product.	
Selected Revision	Tremfya subcutaneous (SC) was added as a Preferred Product for Crohn's Disease. Criteria for Cimzia, Rinvoq, and Entyvio SC were updated to include Tremfya SC as a Preferred Product. For Entyvio SC, a previous trial of Tremfya intravenous also counts.	05/15/2025
Selected Revision	Adalimumab-adaz was removed as a Preferred Product throughout the policy. Throughout the policy, the note was updated for the examples of ustekinumab products which include Stelara, ustekinumab (unbranded Stelara), Imuldosa, Otufli, Pyzchiva, ustekinumab-ttwe, Selarsdi, ustekinumab-aekn, Starjemza, Steqeyma, Wezlana, and Yesintek.	09/01/2025
Annual Revision	Cimzia: For Crohn's Disease , a patient is directed to one Step 1 Product; previously, a patient was directed to adalimumab specifically. A previous trial of an infliximab intravenous product, Omvoh intravenous, Skyrizi intravenous, Tremfya intravenous, or an ustekinumab intravenous product also counts. Simponi Subcutaneous: For Ulcerative Colitis , a patient is directed to one Step 1 Product; previously, a patient was directed to adalimumab specifically. A previous trial of an infliximab intravenous product, Omvoh intravenous, Skyrizi intravenous, Tremfya intravenous, or an ustekinumab intravenous product also counts. Cosentyx SC: For Psoriatic Arthritis , Otezla was added as an agent that counts towards a trial of a Preferred Product for patients < 18 years of age. Orencia SC: For Psoriatic Arthritis , Otezla was added as an agent that counts towards a trial of a Preferred Product for patients < 18 years of age.	11/01/2025

The policy effective date is in force until updated or retired.

APPENDIX A

Table 1. Approved TNFis for Targeted Indications.*

	Rheumatology					Dermatology		Gastroenterology	
	RA	JIA	AS	nr-axSpA	PsA	HS	PsO	CD	UC
Tumor Necrosis Factor Inhibitors									
Cimzia	√	√	√	√	√	--	√	√	--
Enbrel	√	√	√	--	√	--	√	--	--
Adalimumab Products (Humira, biosimilars)	√	√	√	--	√	√	√	√	√
Infliximab Intravenous Products	√	--	√	--	√	--	√	√	√
Zymfentra	--	--	--	--	--	--	--	√^	√^
Simponi Subcutaneous	√	--	√	--	√	--	--	--	√
Simponi Aria	√	√	√	--	√	--	--	--	--

TNFis – Tumor necrosis factor inhibitors; * Refer to the selected standard *Inflammatory Conditions Prior Authorization Policies* for the specific patient population approved for each indication; RA – Rheumatoid arthritis; JIA – Juvenile idiopathic arthritis; AS – Ankylosing spondylitis; nr-axSpA – Non-radiographic spondyloarthritis; PsA – Psoriatic arthritis; HS – Hidradenitis suppurativa; PsO – Plaque psoriasis; CD – Crohn’s disease; UC – Ulcerative colitis; ^ Maintenance dosing only.

Table 2. Approved IL-17, IL-23, and IL-12/23 Blockers for Targeted Indications.*

	Rheumatology			Dermatology		Gastroenterology	
	Ankylosing Spondylitis	nr-axSpA	Psoriatic Arthritis	HS	Plaque Psoriasis	Crohn’s Disease	Ulcerative Colitis
Interleukin-17 Blockers							
Bimzelx	√	√	√	√	√	--	--
Cosentyx Subcutaneous	√	√	√	√	√	--	--
Cosentyx Intravenous	√	√	√	--	--	--	--
Siliq	--	--	--	--	√	--	--
Taltz	√	√	√	--	√	--	--
Interleukin-23 Blockers							
Ilumya	--	--	--	--	√	√	--
OmvoH Intravenous	--	--	--	--	--	√ [#]	√ [#]
OmvoH Subcutaneous	--	--	--	--	--	√^	√^
Skyrizi Intravenous	--	--	--	--	--	√ [#]	√ [#]
Skyrizi Subcutaneous	--	--	√	--	√	√^	√^
Tremfya Intravenous	--	--	--	--	--	√ [#]	√ [#]
Tremfya Subcutaneous	--	--	√	--	√	√ ^μ	√^
Interleukin-12/23 Blockers							
Stelara Subcutaneous	--	--	√	--	√	√^	√^

Stelara Intravenous	--	--	--	--	--	√ [#]	√ [#]
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IL – Interleukin; * Refer to the selected standard *Prior Authorization Policies* for the specific patient population approved for each indication; nr-axSpA – Non-radiographic spondyloarthritis; HS – Hidradenitis suppurativa; ^ Maintenance dosing only; # Induction dosing only; ^u Induction and maintenance dosing.

Table 3. Approved Oral tsDMARDs for Targeted Indications.*

	Rheumatology					Dermatology	Gastroenterology	
	RA	JIA	AS	nr-axSpA	PsA	PsO	CD	UC
Janus Kinases Inhibitors								
Olumiant	√	--	--	--	--	--	--	--
Rinvoq	√	√	√	√	√	--	√	√
Rinvoq LQ	--	√	--	√	--	--	--	--
Xeljanz tablets	√	√ [#]	√	--	√	--	--	√
Xeljanz oral solution	--	√ [#]	--	--	--	--	--	--
Xeljanz XR	√	--	√	--	√	--	--	√
Phosphodiesterase Type 4 Inhibitor								
Otezla	--	--	--	--	√	√	--	--
Sphingosine 1-Phosphate Receptor Modulator								
Velsipity	--	--	--	--	--	--	--	√
Zeposira	--	--	--	--	--	--	--	√
Tyrosine Kinase 2 Inhibitor								
Sotyktu	--	--	--	--	--	√	--	--

tsDMARDs – Targeted synthetic disease-modifying antirheumatic drugs; * Refer to the selected standard *Prior Authorization Policies* for the specific patient population approved for each indication; RA – Rheumatoid arthritis; JIA – Juvenile idiopathic arthritis; AS – Ankylosing spondylitis; nr-axSpA – Nonradiographic axial spondyloarthritis; PsA – Psoriatic arthritis; PsO – Plaque psoriasis; CD – Crohn’s disease; UC – Ulcerative colitis; # Indicated in polyarticular JIA.

Table 4. Other Approved Biologics for Targeted Indications.*

	Rheumatology			Gastroenterology	
	Rheumatoid Arthritis	Juvenile Idiopathic Arthritis	Psoriatic Arthritis	Crohn’s Disease	Ulcerative Colitis
Integrin Receptor Antagonist					
Entyvio Intravenous	--	--	--	√	√
Entyvio Subcutaneous	--	--	--	√ [#]	√ [#]
Interleukin-6 Blockers					
Actemra Intravenous	√	√ [^]	--	--	--
Actemra Subcutaneous	√	√ [^]	--	--	--
Kevzara	√	√ [#]	--	--	--
Interleukin-1 Blocker					
Kineret	√	--	--	--	--
T-Cell Costimulation Modulator					
Orencia Intravenous	√	√ [#]	√	--	--

Orencia Subcutaneous	√	√ [#]	√	--	--
CD20-Directed Cytolytic Antibody					
Rituximab Intravenous Products	√	--	--	--	--

* Refer to the selected standard *Prior Authorization Policies* for the specific patient population approved for each indication; ^ Indicated in polyarticular and systemic JIA; # Indicated in polyarticular JIA; ‡ Maintenance dosing only.

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