

## CARE VALUE POLICY

**POLICY:** Inflammatory Conditions Care Value Policy for National Preferred, High Performance, and Basic Formularies – **Choice/Alternate**

<p><b>Tumor Necrosis Factor Inhibitors</b></p> <ul style="list-style-type: none"> <li>• Adalimumab Products* <ul style="list-style-type: none"> <li>○ adalimumab-adbm subcutaneous injection (Boehringer Ingelheim)</li> <li>○ adalimumab-adaz subcutaneous injection (Sandoz/Novartis)</li> <li>○ adalimumab-ryvk subcutaneous injection (Alvotect/Teva)</li> <li>○ Cyltezo® (adalimumab-adbm subcutaneous injection – Boehringer Ingelheim)</li> <li>○ Simlandi (adalimumab-rykv subcutaneous injection – Alvotect/Teva)</li> </ul> </li> <li>• Cimzia® (certolizumab pegol subcutaneous injection – UCB)</li> <li>• Enbrel® (etanercept subcutaneous injection – Amgen)</li> <li>• Simponi® (golimumab subcutaneous injection – Janssen Biotech/Johnson &amp; Johnson)</li> <li>• Zymfentra® (infliximab-dyyb subcutaneous injection – Celltrion)</li> </ul>
<p><b>Interleukin-6 Blockers</b></p> <ul style="list-style-type: none"> <li>• Tocilizumab Subcutaneous Products <ul style="list-style-type: none"> <li>○ Actemra® (tocilizumab subcutaneous injection – Genentech/Roche)</li> <li>○ Tyenne® (tocilizumab-aazg subcutaneous injection – Fresenius Kabi)</li> </ul> </li> <li>• Kevzara® (sarilumab subcutaneous injection – Regeneron)</li> </ul>
<p><b>Interleukin-17 Blockers</b></p> <ul style="list-style-type: none"> <li>• Bimzelx® (bimekizumab subcutaneous injection – UCB)</li> <li>• Cosentyx® (secukinumab subcutaneous injection – Novartis)</li> <li>• Siliq® (brodalumab subcutaneous injection – Valeant)</li> <li>• Taltz® (ixekizumab subcutaneous injection – Eli Lilly)</li> </ul>
<p><b>Interleukin-23 Blockers</b></p> <ul style="list-style-type: none"> <li>• Ilumya® (tildrakizumab-asmn subcutaneous injection – Sun/Merck)</li> <li>• Omvoh® (mirakizumab-mrkz subcutaneous injection – Eli Lilly)</li> <li>• Skyrizi® (risankizumab-rzaa subcutaneous injection – AbbVie)</li> <li>• Tremfya® (guselkumab subcutaneous injection – Janssen/Johnson &amp; Johnson)</li> </ul>
<p><b>Interleukin 12/23 Blocker</b></p> <ul style="list-style-type: none"> <li>• Stelara® (ustekinumab subcutaneous injection – Janssen Biotech/Johnson &amp; Johnson)</li> </ul>
<p><b>Interleukin-1 Blocker</b></p> <ul style="list-style-type: none"> <li>• Kineret® (anakinra subcutaneous injection – Swedish Orphan Biovitrim)</li> </ul>
<p><b>T-Cell Costimulation Modulator</b></p> <ul style="list-style-type: none"> <li>• Orencia® (abatacept subcutaneous injection – Bristol Myers Squibb)</li> </ul>
<p><b>Integrin Receptor Antagonist</b></p> <ul style="list-style-type: none"> <li>• Entyvio® (vedolizumab subcutaneous injection – Takeda)</li> </ul>
<p><b>Janus Kinases Inhibitors</b></p> <ul style="list-style-type: none"> <li>• Olumiant® (baricitinib tablets – Eli Lilly)</li> <li>• Rinvoq® (upadacitinib extended-release tablets – AbbVie)</li> <li>• Rinvoq® LQ (upadacitinib oral solution – AbbVie)</li> <li>• Xeljanz® (tofacitinib tablets, tofacitinib oral solution – Pfizer)</li> <li>• Xeljanz® XR (tofacitinib extended-release tablets – Pfizer)</li> </ul>
<p><b>Phosphodiesterase Type 4 Inhibitor</b></p> <ul style="list-style-type: none"> <li>• Otezla® (apremilast tablets – Amgen)</li> </ul>
<p><b>Sphingosine 1-Phosphate Receptor Modulator</b></p> <ul style="list-style-type: none"> <li>• Velsipity™ (etrasimod tablets – Pfizer)</li> <li>• Zeposia® (ozanimod capsules – Celgene)</li> </ul>
<p><b>Tyrosine Kinase 2 Inhibitor</b></p> <ul style="list-style-type: none"> <li>• Sotyktu™ (deucravacitinib tablets – Bristol Myers Squibb)</li> </ul>

\* For Non-Preferred adalimumab products, refer to the *Inflammatory Conditions – Adalimumab Products Care Value Policies for National Preferred, High Performance, and Basic Formularies – Alternate or Choice* version of this policy.

**REVIEW DATE:** 10/30/2024 (effective 01/01/2025); selected revision 11/20/2024, 12/04/2024

## 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.<sup>1-28</sup> 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*.

## 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.

**Automation:** None.

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**Preferred and Non-Preferred Products– Rheumatology Indications.<sup>‡</sup>**

	Rheumatology				
	RA	JIA	AS	nr-axSpA	PsA
<b>Step 1 Preferred</b>	<ul style="list-style-type: none"> <li>• Enbrel</li> <li>• Adalimumab Products<sup>^</sup> – Cyltezo/ adalimumab-adbm, adalimumab-adaz, Simlandi/ adalimumab-ryvk</li> </ul>	<ul style="list-style-type: none"> <li>• Enbrel</li> <li>• Adalimumab Products<sup>^</sup> – Cyltezo/ adalimumab-adbm, adalimumab-adaz, Simlandi/ adalimumab-ryvk</li> </ul>	<ul style="list-style-type: none"> <li>• Enbrel</li> <li>• Adalimumab Products<sup>^</sup> – Cyltezo/ adalimumab-adbm, adalimumab-adaz, Simlandi/ adalimumab-ryvk</li> <li>• Taltz</li> </ul>	<ul style="list-style-type: none"> <li>• Cimzia</li> <li>• Taltz</li> </ul>	<ul style="list-style-type: none"> <li>• Enbrel</li> <li>• Adalimumab Products<sup>^</sup> – Cyltezo/ adalimumab-adbm, adalimumab-adaz, Simlandi/ adalimumab-ryvk</li> <li>• Otezla</li> <li>• Skyrizi SC<sup>#</sup></li> <li>• Stelara SC</li> <li>• Taltz</li> <li>• Tremfya SC</li> </ul>
<b>Step 2a Non-Preferred</b> (directed to <b>ONE</b> Step 1 Product)	<ul style="list-style-type: none"> <li>• Tocilizumab SC Products – Actemra SC, Tyenne SC</li> <li>Directed to adalimumab specifically.</li> <li>• Rinvoq</li> <li>Xeljanz tablets/ Xeljanz XR tablets</li> </ul>	<ul style="list-style-type: none"> <li>• Tocilizumab SC Products – Actemra SC, Tyenne SC</li> <li>Directed to adalimumab specifically. JIA Step SC is for P/JIA.</li> <li>• Rinvoq/Rinvoq LQ</li> <li>• Xeljanz tablets/ Xeljanz oral solution</li> </ul>	<ul style="list-style-type: none"> <li>• Rinvoq</li> <li>Directed specifically to Enbrel or adalimumab.</li> <li>• Xeljanz tablets/ Xeljanz XR tablets</li> <li>Directed specifically to Enbrel or adalimumab.</li> </ul>	<ul style="list-style-type: none"> <li>• Rinvoq</li> <li>Directed specifically to Cimzia.</li> </ul>	<ul style="list-style-type: none"> <li>• Rinvoq/ Rinvoq LQ</li> <li>Directed specifically to Enbrel or adalimumab.</li> <li>• Xeljanz tablets/ Xeljanz XR tablets</li> <li>Directed specifically to Enbrel or adalimumab.</li> </ul>
<b>Step 2b Non-Preferred</b> (directed to <b>ONE</b> Step 1 Product)	--	--	• Bimzelx	• Bimzelx	• Bimzelx
<b>Step 3a Non-Preferred</b> (directed to <b>TWO</b> Step 1 or 2a Products) <b>[documentation required]*</b>	<ul style="list-style-type: none"> <li>• Cimzia</li> <li>• Kevzara</li> <li>• Kineret</li> <li>• Olumiant</li> <li>• Orencia SC</li> <li>• Simponi SC</li> </ul>	<ul style="list-style-type: none"> <li>• Cimzia</li> <li>• Kevzara</li> <li>• Orencia SC</li> </ul>	<ul style="list-style-type: none"> <li>• Cimzia</li> <li>• Cosentyx SC</li> <li>• Simponi SC</li> </ul>	<ul style="list-style-type: none"> <li>• Cosentyx SC</li> </ul>	<ul style="list-style-type: none"> <li>• Cimzia</li> <li>• Cosentyx SC</li> <li>• Orencia SC</li> <li>• Simponi SC</li> </ul>

<sup>‡</sup> For Non-Preferred Products, refer to the *Inflammatory Conditions – Adalimumab Products Care Value Policy for National Preferred, High Performance, and Basic Formularies – Alternate* or the *Choice* version of that policy; RA – Rheumatoid arthritis; <sup>^</sup> A trial of more than one adalimumab product counts as ONE Preferred Product; JIA – Juvenile idiopathic arthritis; AS – Ankylosing spondylitis; nr-axSpA – Nonradiographic axial spondyloarthritis; PsA – Psoriatic arthritis; SC – Subcutaneous; <sup>#</sup> Pen and syringe; P/JIA – 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.<sup>‡</sup>**

	Dermatology		Gastroenterology	
	HS	Psoriasis	CD	UC
<b>Step 1 Preferred</b>	<ul style="list-style-type: none"> <li>• <b>Adalimumab Products<sup>^</sup></b> – Cyltezo/ adalimumab-adbm, adalimumab-adaz, Simlandi/ adalimumab-ryvk</li> <li>• <b>Cosentyx SC</b></li> </ul>	<ul style="list-style-type: none"> <li>• <b>Enbrel</b></li> <li>• <b>Adalimumab Products<sup>^</sup></b> –Cyltezo/ adalimumab-adbm, adalimumab-adaz, Simlandi/ adalimumab-ryvk</li> <li>• <b>Otezla</b></li> <li>• <b>Skyrizi SC<sup>#</sup></b></li> <li>• <b>Sotyktu</b></li> <li>• <b>Stelara SC</b></li> <li>• <b>Taltz</b></li> <li>• <b>Tremfya SC</b></li> </ul>	<ul style="list-style-type: none"> <li>• <b>Adalimumab Products<sup>^</sup></b> –Cyltezo/ adalimumab-adbm, adalimumab-adaz, Simlandi/ adalimumab-ryvk</li> <li>• <b>Skyrizi SC</b> (on-body injector)</li> <li>• <b>Stelara SC</b></li> <li>• <b>Zymfentra</b></li> </ul>	<ul style="list-style-type: none"> <li>• <b>Adalimumab Products<sup>^</sup></b> –Cyltezo/ adalimumab-adbm, adalimumab-adaz, Simlandi/ adalimumab-ryvk</li> <li>• <b>OmvoH SC</b></li> <li>• <b>Skyrizi SC</b> (on-body injector)</li> <li>• <b>Stelara SC</b></li> <li>• <b>Tremfya SC</b></li> <li>• <b>Velsipity</b></li> <li>• <b>Zymfentra</b></li> </ul>
<b>Step 2a Non-Preferred</b> (directed to <b>ONE</b> Step 1 Product)	--	--	<ul style="list-style-type: none"> <li>• <b>Cimzia</b> Directed to adalimumab specifically.</li> <li>• <b>Rinvoq</b> Directed to adalimumab specifically.</li> </ul>	<ul style="list-style-type: none"> <li>• <b>Rinvoq</b> Directed to adalimumab specifically.</li> <li>• <b>Simponi SC</b> Directed to adalimumab specifically.</li> <li>• <b>Xeljanz tablets/ XR tablets</b> Directed to adalimumab specifically.</li> </ul>
<b>Step 2b Non-Preferred</b> (directed to <b>ONE</b> Step 1 Product)	• <b>Bimzelx</b>	• <b>Bimzelx</b>	--	--
<b>Step 3a Non-Preferred</b> (directed to <b>TWO</b> Step 1 or 2a Products) <b>[documentation required]*</b>	--	<ul style="list-style-type: none"> <li>• <b>Cimzia</b></li> <li>• <b>Cosentyx SC</b></li> <li>• <b>Humya</b></li> <li>• <b>Siliq</b></li> </ul>	• <b>Entyvio SC</b>	• <b>Entyvio SC</b>
<b>Step 3b Non-Preferred</b> (directed to <b>TWO</b> Step 1 Products)	--	--	--	• <b>Zeposia</b> Refer to MS and UC – Zeposia Care Value Policy

<sup>‡</sup> For Non-Preferred Products, refer to the *Inflammatory Conditions – Adalimumab Products Care Value Policy for for National Preferred, High Performance, and Basic Formularies – Alternate* or the *Choice* version of that policy; <sup>^</sup> A trial of more than one adalimumab product counts as ONE Preferred Product; HS – Hidradenitis suppurativa; CD – Crohn’s disease; UC – Ulcerative colitis; SC – Subcutaneous; <sup>#</sup> 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.

**RECOMMENDED EXCEPTION CRITERIA**

Non-Preferred Product	Exception Criteria
<b>Tumor Necrosis Factor Inhibitors</b>	
<b>Cimzia</b>	<p><b>1. <u>Rheumatoid Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR <b>[documentation required]</b>. <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product.</li> </ul> <p><b>B)</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: offer to review for a Step 1 or Step 2a Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>2. <u>Ankylosing Spondylitis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried TWO of Enbrel, an adalimumab product, Rinvoq, Taltz, or Xeljanz/XR <b>[documentation required]</b>. <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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product.</li> </ul> <p><b>B)</b> If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: offer to review for a Step 1 or Step 2 Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Taltz, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>3. <u>Juvenile Idiopathic Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria; AND</li> </ul>

	<p><b>ii.</b> Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq/Rinvoq LQ, and Xeljanz <b>[documentation required]</b>; OR</p> <p><u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of both tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz tablets and Xeljanz oral solution) collectively counts as <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> 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 <b>[documentation required]</b>.</p> <p><b>B)</b> If the patient has met criterion 3Ai (the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria), but criterion 3Aii is not met: offer to review for a Step 1 or Step 2a Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Rinvoq LQ, Xeljanz tablets, or Xeljanz oral solution</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>4. Psoriatic Arthritis – Initial Therapy.</b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <p><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria; AND</p> <p><b>ii.</b> Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, Stelara subcutaneous, Taltz, Tremfya subcutaneous, and Xeljanz/XR <b>[documentation required]</b>.</p> <p><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. A trial of multiple adalimumab products counts as <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product.</p> <p><b>B)</b> If the patient has met criterion 4Ai (the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria), but criterion 3Aii is not met: offer to review for a Step 1 or Step 2a Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>5. Plaque Psoriasis – Initial Therapy.</b></p> <p><b>A)</b> Approve for 3 months if the patient meets BOTH of the following (i and ii):</p>
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	<ul style="list-style-type: none"><li>i. Patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria; AND</li><li>ii. Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous <b>[documentation required]</b>. <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. A trial of multiple adalimumab products counts as <b>ONE</b> product.</li></ul> <p><b>B)</b> If the patient has met criterion 5Ai (the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria), but criterion 4Aii is not met: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>6. <u>Crohn’s Disease – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"><li>i. Patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria; AND</li><li>ii. Patient has tried one adalimumab product. <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.</li></ul> <p><b>B)</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: offer to review for a Preferred Product (<u>adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Skyrizi subcutaneous [on-body injector], Stelara subcutaneous, or Zymfentra</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>7. <u>Rheumatoid Arthritis, Ankylosing Spondylitis, Juvenile Idiopathic Arthritis, Psoriatic Arthritis, Plaque Psoriasis, or Crohn’s Disease – Patient is Currently Receiving Cimzia.</u></b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"><li>i. Patient meets the standard <i>Inflammatory Conditions – Cimzia Prior Authorization Policy</i> criteria; AND</li><li>ii. Patient meets ONE of the following (a, b, c, d, e, f, or g):<ul style="list-style-type: none"><li>a) Patient has <u>Rheumatoid Arthritis</u> and has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR <b>[documentation required]</b>; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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</li></ul></li></ul>
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	<p>adalimumab products counts as <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product.</p> <p>b) Patient has <u>Ankylosing Spondylitis</u> and has tried TWO of Enbrel, an adalimumab product, Rinvoq, Taltz, or Xeljanz/XR <b>[documentation required]</b>; 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product.</p> <p>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 <b>[documentation required]</b>; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of both tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz tablets and Xeljanz oral solution) collectively counts as <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> 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 <b>[documentation required]</b>.</p> <p>d) Patient has <u>Psoriatic Arthritis</u> and has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, Stelara subcutaneous, Taltz, Tremfya subcutaneous, or Xeljanz/XR <b>[documentation required]</b>; 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product.</p> <p>e) Patient has <u>Plaque Psoriasis</u> and has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous <b>[documentation required]</b>; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-fkjp, adalimumab-aaty, adalimumab-</p>
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	<p>ryvk, Simlandi, adalimumab-adbm, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as <b>ONE</b> product.</p> <p>f) Patient has <u>Crohn's Disease</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.</p> <p>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> [<b>verification in prescription claims history required</b>], or if claims history is not available, according to the prescriber [<b>verification by prescriber required</b>]. <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).</p> <p><b>B)</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: offer to review for one of the following Products using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria:</p> <ul style="list-style-type: none"><li>i. <b>Rheumatoid Arthritis:</b> <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR.</u></li><li>ii. <b>Juvenile Idiopathic Arthritis:</b> <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Rinvoq LQ, Xeljanz tablets, or Xeljanz oral solution.</u></li><li>iii. <b>Ankylosing Spondylitis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Taltz, Xeljanz tablets, or Xeljanz XR.</u></li><li>iv. <b>Psoriatic Arthritis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz tablets, or Xeljanz XR.</u></li><li>v. <b>Plaque Psoriasis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous (pen or syringe), Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous.</u></li><li>vi. <b>Crohn's Disease:</b> <u>adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, or Zymfentra.</u></li></ul> <p><b>8. Other Conditions.</b> 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.</p>
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<p><b>Simponi Subcutaneous</b></p>	<p><b>1. <u>Rheumatoid Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR <b>[documentation required]</b>;</li> </ul> <p><b>OR</b></p> <p><u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product.</p> <p><b>B)</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: offer to review for a Step 1 or Step 2 Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>2. <u>Ankylosing Spondylitis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried TWO of Enbrel, an adalimumab product, Rinvoq, Taltz, or Xeljanz/XR <b>[documentation required]</b>.</li> </ul> <p><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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product.</p> <p><b>B)</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: offer to review for a Step 1 or Step 2a Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Taltz, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>3. <u>Psoriatic Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, Stelara subcutaneous, Taltz, Tremfya subcutaneous, or Xeljanz/XR <b>[documentation required]</b>.</li> </ul>
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	<p><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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product.</p> <p><b>B)</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: offer to review for a Step 1 or Step 2 Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>4. <u>Ulcerative Colitis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ol style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> 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.</li></ol> <p><b>B)</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: offer to review for a Preferred Product (<u>adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Tremfya subcutaneous, Velsipity, or Zymfentra</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>5. <u>Rheumatoid Arthritis, Ankylosing Spondylitis, Psoriatic Arthritis, or Ulcerative Colitis – Patient is Currently Receiving Simponi Subcutaneous or Aria.</u></b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i and ii):</p> <ol style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Simponi Subcutaneous Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient meets ONE of the following (a, b, c, d, e, or f):<ol style="list-style-type: none"><li><b>a)</b> Patient has <u>Rheumatoid Arthritis</u> and has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [<b>documentation required</b>]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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</li></ol></li></ol>
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	<p>adalimumab products counts as <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product.</p> <p>b) Patient has <u>Ankylosing Spondylitis</u> and has tried TWO of Enbrel, an adalimumab product, Rinvoq, Taltz, or Xeljanz/XR [<b>documentation required</b>]; OR</p> <p><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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product.</p> <p>c) Patient has <u>Psoriatic Arthritis</u> and has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, Stelara subcutaneous, Taltz, Tremfya subcutaneous, or Xeljanz/XR [<b>documentation required</b>]; OR</p> <p><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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product.</p> <p>d) Patient has <u>Ulcerative Colitis</u> and has tried one adalimumab product; OR</p> <p><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.</p> <p>e) According to the prescriber, the patient has been established on Simponi Aria for at least 90 days; OR</p> <p>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> [<b>verification in prescription claims history required</b>], or if claims history is not available, according to the prescriber [<b>verification by prescriber required</b>].</p> <p><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).</p> <p><b>B)</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: offer to review for one of the following Products using the</p>
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	<p>respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria:</p> <ul style="list-style-type: none"> <li>i. <b>Rheumatoid Arthritis:</b> <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR.</u></li> <li>ii. <b>Ankylosing Spondylitis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Taltz, Xeljanz tablets, or Xeljanz XR.</u></li> <li>iii. <b>Psoriatic Arthritis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz tablets, or Xeljanz XR.</u></li> <li>iv. <b>Ulcerative Colitis:</b> <u>adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Tremfya subcutaneous, Velsipity, or Zymfentra.</u></li> </ul> <p>6. <b>Other Conditions.</b> 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.</p>
<b>Interleukin-6 Blockers</b>	
<p><b>Actemra Subcutaneous Tyenne Subcutaneous</b></p>	<p>1. <b><u>Polyarticular Juvenile Idiopathic Arthritis – Initial Therapy.</u></b></p> <p>A) Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li>i. Patient meets the standard <i>Inflammatory Conditions – Tocilizumab Subcutaneous Prior Authorization Policy</i> criteria; AND</li> <li>ii. Patient meets ONE of the following (a or b): <ul style="list-style-type: none"> <li>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.</li> <li>b) According to the prescriber, the patient has heart failure or a previously treated lymphoproliferative disorder.</li> </ul> </li> </ul> <p>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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p>2. <b><u>Rheumatoid Arthritis – Initial Therapy.</u></b></p> <p>A) Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li>i. Patient meets the standard <i>Inflammatory Conditions – Tocilizumab Subcutaneous Prior Authorization Policy</i> criteria; AND</li> <li>ii. Patient meets ONE of the following (a or b): <ul style="list-style-type: none"> <li>a) Patient has tried one adalimumab product; OR  <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp,</li> </ul> </li> </ul>

	<p>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.</p> <p>b) According to the prescriber, the patient has heart failure or a previously treated lymphoproliferative disorder.</p> <p>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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p>3. <b><u>Polyarticular Juvenile Idiopathic Arthritis or Rheumatoid Arthritis – Patient is Currently Receiving Tocilizumab Subcutaneous or Intravenous.</u></b></p> <p>A) Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii):</p> <p>i. Patient meets the standard <i>Inflammatory Conditions – Tocilizumab Subcutaneous Policy</i> criteria; AND</p> <p>ii. Patient meets ONE of the following (a, b, c, d, <u>or</u> e):</p> <p>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.</p> <p>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.</p> <p>c) According to the prescriber, the patient has heart failure or a previously treated lymphoproliferative disorder; OR</p> <p>d) According to the prescriber, the patient has been established on tocilizumab intravenous for at least 90 days; OR</p> <p>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</u> <b>[verification in prescription claims history required]</b>, or if claims history is not available, according to the prescriber <b>[verification by prescriber required]</b>. <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 90 days AND the patient has been receiving tocilizumab subcutaneous via paid</p>
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	<p>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).</p> <p><b>B)</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: offer to review for a Preferred Product using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria:</p> <p><b>i. Polyarticular Juvenile Idiopathic Arthritis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi.</u></p> <p><b>ii. Rheumatoid Arthritis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi.</u></p> <p><b>4. <u>All Other Conditions</u></b> (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.</p>
<p><b>Kevzara</b></p>	<p><b>1. <u>Rheumatoid Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <p><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Kevzara Prior Authorization Policy</i> criteria; AND</p> <p><b>ii.</b> Patient meets ONE of the following (a or b):</p> <p><b>a)</b> Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR <b>[documentation required]</b>; OR</p> <p><u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> 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 <b>[documentation required]</b>.</p> <p><b>b)</b> According to the prescriber, the patient has heart failure or a previously treated lymphoproliferative disorder.</p> <p><b>B)</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: offer to review for a Step 1 or Step 2 Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>2. <u>Juvenile Idiopathic Arthritis/Juvenile Rheumatoid Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p>

	<ul style="list-style-type: none"><li>i. Patient meets the standard <i>Inflammatory Conditions – Kevzara Prior Authorization Policy</i> criteria; AND</li><li>ii. Patient meets ONE of the following conditions (a <u>or</u> b):<ul style="list-style-type: none"><li>a) Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq/Rinvoq LQ, or Xeljanz <b>[documentation required]</b>; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> 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 <b>[documentation required]</b>.</li><li>b) According to the prescriber, the patient has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, OR a demyelinating disorder.</li></ul></li></ul> <p><b>B)</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: offer to review for a Step 1 or Step 2a Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Rinvoq LQ, or Xeljanz tablets</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>3. <u>Juvenile Idiopathic Arthritis or Rheumatoid Arthritis – Patient is Currently Receiving Kevzara.</u></b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"><li>i. Patient meets the standard <i>Inflammatory Conditions – Kevzara Prior Authorization Policy</i> criteria; AND</li><li>ii. Patient meets ONE of the following (a, b, c, <u>or</u> d):<ul style="list-style-type: none"><li>a) Patient has <u>Rheumatoid Arthritis</u> and has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR <b>[documentation required]</b>; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of tocilizumab intravenous (Actemra intravenous, biosimilar), Cimzia, an infliximab product (e.g.,</li></ul></li></ul>
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	<p>Remicade, biosimilars), Orenzia (intravenous or subcutaneous), or Simponi (Aria or subcutaneous) also counts <b>[documentation required]</b>.</p> <p>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 <b>[documentation required]</b>; OR  <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product. A trial of a Cimzia, tocilizumab intravenous product (Actemra intravenous, biosimilar), Orenzia intravenous or subcutaneous, an infliximab product (e.g., Remicade, biosimilars), or Simponi Aria also counts <b>[documentation required]</b>.</p> <p>c) According to the prescriber, the patient has heart failure or a previously treated lymphoproliferative disorder; OR</p> <p>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> <b>[verification in prescription claims history required]</b>, or if claims history is not available, according to the prescriber <b>[verification by prescriber required]</b>.  <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).</p> <p><b>B)</b> If the patient has met criterion 3Ai (the standard <i>Inflammatory Conditions – Kevzara Prior Authorization Policy</i> criteria), but criterion 3Aii is not met: offer to review for one of the following Products using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p>i. <b>Rheumatoid Arthritis:</b> <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR.</u></p> <p>ii. <b>Juvenile Idiopathic Arthritis:</b> <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Rinvoq LQ, or Xeljanz tablets.</u></p> <p><b>3. Other Conditions.</b> 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 standard <i>Inflammatory Conditions – Kevzara Prior Authorization Policy</i> criteria.</p>
<b>Interleukin-17 Blockers</b>	
<b>Bimzelx</b>	<b>1. <u>Ankylosing Spondylitis – Initial Therapy.</u></b>

	<p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> 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.</li></ul> <p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, Cyltezo, adalimumab-adbm, adalimumab-adaz, Simlandi, adalimumab-ryvk, or Taltz</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>2. <u>Hidradenitis Suppurativa – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 3 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria for hidradenitis suppurativa; AND</li><li><b>ii.</b> 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.</li></ul> <p><b>B)</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>adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, or Cosentyx subcutaneous</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria</p> <p><b>3. <u>Non-Radiographic Spondyloarthritis (nr-axSpA) – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> 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.</li></ul> <p><b>B)</b> If the patient has met criterion 3Ai (the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria), but criterion 3Aii is not met: offer to review for a Preferred Product (<u>Cimzia or Taltz</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>4. <u>Plaque Psoriasis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 3 months if the patient meets BOTH of the following (i and ii):</p>
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	<ul style="list-style-type: none"><li>i. Patient meets the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria for plaque psoriasis; AND</li><li>ii. Patient has tried ONE of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, Stelara subcutaneous, Taltz, or Tremfya 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.</li></ul> <p><b>B)</b> If the patient has met criterion 4Ai (the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria), but criterion 4Aii is not met: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>5. <u>Psoriatic Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"><li>i. Patient meets the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria; AND</li><li>ii. Patient has tried one of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Stelara subcutaneous, 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, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts.</li></ul> <p><b>B)</b> If the patient has met criterion 5Ai (the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria), but criterion 5Aii is not met: offer to review for a Preferred Product (<u>Enbrel, Cyltezo, adalimumab-adbm, adalimumab-adaz, Simlandi, adalimumab-ryvk, Otezla, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Taltz, or Tremfya subcutaneous</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>6. <u>Ankylosing Spondylitis, Hidradenitis Suppurativa, nr-axSpA, Plaque Psoriasis, or Psoriatic Arthritis – Patient is Currently Receiving Bimzelx.</u></b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"><li>i. Patient meets the standard <i>Inflammatory Conditions – Bimzelx Prior Authorization Policy</i> criteria; AND</li><li>ii. Patient meets ONE of the following (a, b, c, d, e, or f):<ul style="list-style-type: none"><li>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.</li></ul></li></ul>
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	<p>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.</p> <p>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</p> <p>d) Patient has <u>Plaque Psoriasis</u> and has tried ONE of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, Stelara subcutaneous, 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.</p> <p>e) Patient has <u>Psoriatic Arthritis</u> and has tried one of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Stelara subcutaneous, 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, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts.</p> <p>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 [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 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).</p> <p><b>B)</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: offer to review for one of the following Preferred Products (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous,</u></p>
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	<p>Taltz, or Tremfya subcutaneous) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <ul style="list-style-type: none"> <li>i. <b>Ankylosing Spondylitis:</b> <u>Enbrel, Cyltezo, adalimumab-adbm, adalimumab-adaz, adalimumab-ryvk, Simlandi, or Taltz.</u></li> <li>ii. <b>Hidradenitis Suppurativa:</b> <u>Cyltezo, adalimumab-adbm, adalimumab-adaz, adalimumab-ryvk, Simlandi, or Cosentyx subcutaneous.</u></li> <li>iii. <b>nr-axSpA:</b> <u>Cimzia or Taltz.</u></li> <li>iv. <b>Plaque Psoriasis:</b> <u>Enbrel, Cyltezo, adalimumab-adbm, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous.</u></li> <li>v. <b>Psoriatic Arthritis:</b> <u>Enbrel, Cyltezo, adalimumab-adbm, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Taltz, or Tremfya subcutaneous</u></li> </ul> <p>7. <b>Other Conditions.</b> Approve <u>Bimzelx</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 – Bimzelx Prior Authorization Policy</i> criteria.</p>
<p><b>Cosentyx SC</b></p>	<ul style="list-style-type: none"> <li>1. <b>Ankylosing Spondylitis – Initial Therapy.</b> <ul style="list-style-type: none"> <li>A) Approve for 6 months if the patient meets BOTH of the following (i and ii): <ul style="list-style-type: none"> <li>i. Patient meets the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria; AND</li> <li>ii. Patient has tried TWO of Enbrel, an adalimumab product, Rinvoq, Taltz, or Xeljanz/XR <b>[documentation required]</b>. <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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of Cimzia, an infliximab product (e.g. Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts <b>[documentation required]</b>.</li> </ul> </li> <li>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: offer to review for a Step 1 or Step 2a Product (<u>adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Enbrel, Rinvoq, Taltz, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</li> </ul> </li> <li>2. <b>Non-Radiographic Spondyloarthritis (nr-axSpA) – Initial Therapy.</b> <ul style="list-style-type: none"> <li>A) Approve for 6 months if the patient meets BOTH of the following (i and ii): <ul style="list-style-type: none"> <li>i. Patient meets the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria; AND</li> <li>ii. Patient has tried TWO of Cimzia, Taltz, or Rinvoq <b>[documentation required]</b>. <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 <b>[documentation required]</b>. A trial of multiple adalimumab products counts as <b>ONE</b> product.</li> </ul> </li> </ul> </li> </ul>

	<p><b>B)</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: offer to review for a Step 1 or Step 2a Product (<u>Cimzia, Taltz, or Rinvoq</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>3. <u>Plaque Psoriasis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 3 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ol style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous <b>[documentation required]</b>.</li></ol> <p><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 <b>ONE</b> product.</p> <p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>4. <u>Psoriatic Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ol style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient meets ONE of the following (a <u>or</u> b):<ol style="list-style-type: none"><li><b>a)</b> Patient is <math>\geq 18</math> years of age AND has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, Stelara subcutaneous, Taltz, Tremfya subcutaneous, or Xeljanz/XR <b>[documentation required]</b>; OR</li><li><b>b)</b> Patient is <math>&lt; 18</math> years of age AND has tried ONE of Enbrel, Rinvoq/Rinvoq LQ, or Stelara SC <b>[documentation required]</b>.</li></ol></li></ol> <p><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 <b>ONE</b> 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 <b>[documentation required]</b>. For a patient <math>&lt; 18</math> years of age, a trial of another TNFi counts towards a trial of Enbrel <b>[documentation required]</b>. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product.</p>
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	<p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>5. <u>Ankylosing Spondylitis; nr-axSpA; Plaque Psoriasis; or Psoriatic Arthritis – Patient is Currently Receiving Cosentyx (Subcutaneous or Intravenous).</u></b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i and ii):</p> <ol style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient meets ONE of the following (a, b, c, d, e, f, or g):<ol style="list-style-type: none"><li><b>a)</b> Patient has <u>Ankylosing Spondylitis</u> and has tried TWO of Enbrel, an adalimumab product, Rinvoq, Taltz, or Xeljanz/XR <b>[documentation required]</b>; 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts <b>[documentation required]</b>.</li><li><b>b)</b> Patient has <u>nr-axSpA</u> and has tried TWO of Cimzia, Taltz, or Rinvoq <b>[documentation required]</b>; OR <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 <b>[documentation required]</b>. A trial of multiple adalimumab products counts as <b>ONE</b> product.</li><li><b>c)</b> Patient has <u>Plaque Psoriasis</u> and has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous <b>[documentation required]</b>; 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 <b>ONE</b> product.</li><li><b>d)</b> Patient is <math>\geq 18</math> years of age with <u>Psoriatic Arthritis</u> and has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, Stelara subcutaneous, Taltz, Tremfya subcutaneous, or Xeljanz/XR <b>[documentation required]</b>; OR <u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp,</li></ol></li></ol>
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	<p>adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo, Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of multiple adalimumab products counts as <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi (Aria or subcutaneous) also counts <b>[documentation required]</b>.</p> <p>e) Patient is &lt; 18 years of age with <u>Psoriatic Arthritis</u> and has tried ONE of Enbrel, Rinvoq/Rinvoq LQ, or Stelara subcutaneous <b>[documentation required]</b>; OR</p> <p><u>Note:</u> A trial of another TNFi counts towards a trial of Enbrel <b>[documentation required]</b>. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product.</p> <p>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</p> <p>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> <b>[verification in prescription claims history required]</b>, or if claims history is not available, according to the prescriber <b>[verification by prescriber required]</b>.</p> <p><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).</p> <p><b>B)</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: offer to review for one of the following Products using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria:</p> <p><b>i. Ankylosing Spondylitis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Taltz, Xeljanz tablets, or Xeljanz XR.</u></p> <p><b>ii. nr-axSpA:</b> <u>Cimzia, Taltz, or Rinvoq.</u></p> <p><b>iii. Plaque Psoriasis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous (pen or syringe), Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous.</u></p> <p><b>iv. Psoriatic Arthritis in a Patient ≥ 18 years of age:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz, or Xeljanz XR.</u></p> <p><b>v. Psoriatic Arthritis in a Patient &lt; 18 years of age:</b> <u>Enbrel, Rinvoq, Rinvoq LQ, or Stelara SC.</u></p>
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	<p><b>6. Other Conditions.</b> Approve <u>Cosentyx 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 – Cosentyx Subcutaneous Prior Authorization Policy</i> criteria.</p>
<p><b>Siliq</b></p>	<p><b>1. Plaque Psoriasis – Initial Therapy.</b></p> <p>A) Approve for 3 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li>i. Patient meets the standard <i>Inflammatory Conditions – Siliq Prior Authorization Policy</i> criteria for plaque psoriasis; AND</li> <li>ii. Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous <b>[documentation required]</b>.</li> </ul> <p><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 <b>ONE</b> product.</p> <p>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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>2. Plaque Psoriasis – Patient is Currently Receiving Siliq.</b></p> <p>A) Approve for 1 year if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li>i. Patient meets the standard <i>Inflammatory Conditions – Siliq Prior Authorization Policy</i> criteria; AND</li> <li>ii. Patient meets ONE of the following (a or b):</li> </ul> <ul style="list-style-type: none"> <li>a) Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous <b>[documentation required]</b>; OR</li> </ul> <p><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 <b>ONE</b> product.</p> <ul style="list-style-type: none"> <li>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> <b>[verification in prescription claims history required]</b>, or if claims history is not available, according to the prescriber <b>[verification by prescriber required]</b>.</li> </ul> <p><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).</p>

	<p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>3. <u>Other Conditions.</u></b> 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.</p>
<b>Interleukin-23 Blockers</b>	
<p><b>Ilumya</b></p>	<p><b>1. <u>Plaque Psoriasis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 3 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Ilumya Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous <b>[documentation required]</b>.</li> </ul> <p><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 <b>ONE</b> product.</p> <p><b>B)</b> If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Ilumya Prior Authorization Policy</i> criteria), but criterion 1Aii is not met: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>2. <u>Plaque Psoriasis – Patient is Currently Receiving Ilumya.</u></b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Ilumya Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient meets ONE of the following (a or b): <ul style="list-style-type: none"> <li><b>a)</b> Patient has plaque psoriasis and has tried TWO of Enbrel, an adalimumab product, Otezla, Skyrizi subcutaneous, Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous <b>[documentation required]</b>; OR</li> <li><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 <b>ONE</b> product.</li> <li><b>b)</b> Patient has been established on Ilumya for at least 90 days <u>and</u> prescription claims history indicates <u>at least a 90-day supply of Ilumya was dispensed within the past 130 days</u> <b>[verification in prescription claims history required]</b>, or if claims history is not</li> </ul> </li> </ul>

	<p>available, according to the prescriber <b>[verification by prescriber required]</b>.</p> <p><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 Ilumya for at least 90 days AND the patient has been receiving Ilumya 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 Ilumya).</p> <p><b>B)</b> If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Ilumya Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Sotyktu, Stelara subcutaneous, Taltz, or Tremfya subcutaneous</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>3. <u>Other Conditions.</u></b> Approve <u>Ilumya</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 – Ilumya Prior Authorization Policy</i> criteria.</p>
<b>Integrin Receptor Antagonist</b>	
<p><b>Entyvio SC</b></p>	<p><b>1. <u>Crohn’s Disease – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Entyvio Subcutaneous Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient meets ONE of the following (a <u>or</u> b): <ul style="list-style-type: none"> <li><b>a)</b> Patient has tried TWO of an adalimumab product, Skyrizi subcutaneous, Stelara subcutaneous, Zymfentra, Cimzia, or Rinvoq <b>[documentation required]</b>; OR</li> </ul> </li> </ul> <p><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 multiple adalimumab products counts as <b>ONE</b> product. A trial of an infliximab intravenous product (e.g., Remicade, biosimilars), Skyrizi intravenous, or Stelara intravenous also counts <b>[documentation required]</b>.</p> <ul style="list-style-type: none"> <li><b>b)</b> According to the prescriber, the patient has already started on or is currently undergoing induction therapy with Entyvio IV.</li> </ul> <p><b>B)</b> If the patient has met criterion 1Ai (the standard <i>Inflammatory Conditions – Entyvio Subcutaneous Prior Authorization Policy</i> criteria), but criterion 1Aii is not met, offer to review for a Step 1 or Step 2a Product (<u>adalimumab-adaz, adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Rinvoq, Cimzia, or Zymfentra</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>2. <u>Ulcerative Colitis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Entyvio Subcutaneous Prior Authorization Policy</i> criteria; AND</li> </ul>

	<p>ii. Patient meets ONE of the following (a <u>or</u> b):</p> <p>a) Patient has tried TWO of an adalimumab product, Skyrizi subcutaneous, Stelara subcutaneous, Zymfentra, Omvoh subcutaneous, Rinvoq, Simponi subcutaneous, Tremfya subcutaneous, Velsipity, or Xeljanz/XR <b>[documentation required]</b>; OR</p> <p><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 multiple adalimumab products counts as <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of an infliximab intravenous product (e.g., Remicade, biosimilars), Omvoh intravenous, Skyrizi intravenous, Stelara intravenous, or Tremfya intravenous also counts <b>[documentation required]</b>.</p> <p>b) According to the prescriber, the patient has already started on or is currently undergoing induction therapy with Entyvio IV.</p> <p><b>B)</b> If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Entyvio Subcutaneous Prior Authorization Policy</i> criteria), but criterion 2Aii is not met, offer to review for a Step 1 or Step 2a Product (<u>adalimumab-adaz, adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Stelara subcutaneous, Omvoh subcutaneous, Rinvoq, Simponi SC, Skyrizi subcutaneous (on-body injector), Xeljanz/XR, Tremfya subcutaneous, Velsipity, or Zymfentra</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>3. <u>Crohn’s Disease and Ulcerative Colitis – Patient is Currently Receiving Entyvio Subcutaneous or Intravenous.</u></b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii):</p> <p>i. Patient meets the standard <i>Inflammatory Conditions – Entyvio Subcutaneous Prior Authorization Policy</i> criteria; AND</p> <p>ii. Patient meets ONE of the following conditions (a, b, c, <u>or</u> d):</p> <p>a) Patient has <u>Crohn’s Disease</u> and has tried TWO of an adalimumab product, Skyrizi subcutaneous, Stelara subcutaneous, Zymfentra, Cimzia, or Rinvoq <b>[documentation required]</b>; OR</p> <p><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 multiple adalimumab products counts as <b>ONE</b> product. A trial of an infliximab intravenous product (e.g., Remicade, biosimilars), Skyrizi intravenous, or Stelara intravenous also counts <b>[documentation required]</b>.</p> <p>b) Patient has <u>Ulcerative Colitis</u> and has tried TWO of an adalimumab product, Skyrizi subcutaneous, Stelara subcutaneous, Tremfya subcutaneous, Zymfentra, Omvoh subcutaneous, Rinvoq, Simponi subcutaneous, Velsipity, or Xeljanz/XR <b>[documentation required]</b>; OR</p>
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	<p><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 multiple adalimumab products counts as <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of an infliximab intravenous product (e.g., Remicade, biosimilars), Omvoh intravenous, Skyrizi intravenous, Stelara intravenous, or Tremfya intravenous also counts <b>[documentation required]</b>.</p> <p>c) According to the prescriber, the patient has been established on Entyvio intravenous for at least 90 days; OR</p> <p>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 [verification in prescription claims history required]</u>, or if claims history is not available, according to the prescriber <b>[verification by prescriber required]</b>.</p> <p><u>Note:</u> In cases where 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 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).</p> <p><b>B)</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, offer to review for one of the following Products using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p>i. <b>Crohn’s Disease:</b> <u>adalimumab-adaz, adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Rinvoq, Cimzia, or Zymfentra.</u></p> <p>ii. <b>Ulcerative Colitis:</b> <u>adalimumab-adaz, adalimumab-adbm, Cyltezo, adalimumab-ryvk, Simlandi, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Tremfya subcutaneous, Omvoh subcutaneous, Rinvoq, Simponi SC, Xeljanz/XR, Velsipity, or Zymfentra.</u></p> <p><b>4. Other Conditions.</b> 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.</p>
<b>Interleukin-1 Blocker</b>	
<b>Kineret</b>	<p><b>1. Rheumatoid Arthritis – Initial Therapy.</b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <p>i. Patient meets the standard <i>Inflammatory Conditions – Kineret Prior Authorization Policy</i> criteria; AND</p> <p>ii. Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR <b>[documentation required]</b>.</p>

	<p><u>Note</u>: Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of tocilizumab intravenous (Actemra intravenous, biosimilar), Cimzia, Orenzia (subcutaneous or intravenous), an infliximab product (e.g., Remicade, biosimilar), Kevzara, or Simponi (Aria or subcutaneous) also counts <b>[documentation required]</b>.</p> <p><b>B)</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: offer to review for a Step 1 or Step 2a Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>2. <u>Rheumatoid Arthritis – Patient is Currently Receiving Kineret.</u></b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Kineret Prior Authorization Policy</i> criteria; <b>AND</b></li><li><b>ii.</b> Patient meets ONE of the following (a <u>or</u> b):<ul style="list-style-type: none"><li><b>a)</b> Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR <b>[documentation required]</b>; <b>OR</b> <u>Note</u>: Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of tocilizumab intravenous (Actemra intravenous, biosimilar), Cimzia, Orenzia (subcutaneous or intravenous), an infliximab product (e.g., Remicade, biosimilar), Kevzara, or Simponi (Aria or subcutaneous) also counts <b>[documentation required]</b>.</li><li><b>b)</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> <b>[verification in prescription claims history required]</b>, or if claims history is not available, according to the prescriber <b>[verification by prescriber required]</b>.</li></ul></li></ul>
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	<p><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).</p> <p><b>B)</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: offer to review for a Step 1 or Step 2a Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>3. Other Conditions.</b> Approve <u>Kineret</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 – Kineret Prior Authorization Policy</i> criteria. <u>Note</u>: This includes Cryopyrin-Associated Periodic Syndromes (CAPS), Systemic Juvenile Idiopathic Arthritis.</p>
<b>T-Cell Costimulation Modulator</b>	
<p><b>Orencia Subcutaneous</b></p>	<p><b>1. Rheumatoid Arthritis – Initial Therapy.</b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Orencia Subcutaneous Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient meets ONE of the following (a <u>or</u> b): <ul style="list-style-type: none"> <li><b>a)</b> Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR [<b>documentation required</b>]; OR <u>Note</u>: Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> 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 [<b>documentation required</b>].</li> <li><b>b)</b> According to the prescriber, the patient has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, OR a demyelinating disorder.</li> </ul> </li> </ul> <p><b>B)</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: offer to review for a Step 1 or Step 2a Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or</u></p>

	<p><u>Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>2. <u>Juvenile Idiopathic Arthritis/Juvenile Rheumatoid Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Oencia Subcutaneous Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient meets ONE of the following (a <u>or</u> b):<ul style="list-style-type: none"><li><b>a)</b> Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq/Rinvoq LQ, or Xeljanz <b>[documentation required]</b>; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz tablets and Xeljanz oral solution) collectively counts as <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product. A trial of Cimzia, tocilizumab intravenous (Actemra intravenous, biosimilar), Kevzara, Oencia intravenous, an infliximab product (e.g., Remicade, biosimilar), or Simponi Aria also counts <b>[documentation required]</b>.</li><li><b>b)</b> According to the prescriber, the patient has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, OR a demyelinating disorder.</li></ul></li></ul> <p><b>C)</b> If the patient has met criterion 2Ai (the standard <i>Inflammatory Conditions – Oencia Subcutaneous Prior Authorization Policy</i> criteria), but criterion 2Aii is not met: offer to review for a Step 1 or Step 2a Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Rinvoq LQ, Xeljanz tablets, or Xeljanz oral solution</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>3. <u>Psoriatic Arthritis – Initial Therapy.</u></b></p> <p><b>B)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Oencia Subcutaneous Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient meets ONE of the following (a, b, <u>or</u> c):<ul style="list-style-type: none"><li><b>a)</b> Patient is <math>\geq 18</math> years of age AND has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, Stelara subcutaneous, Taltz, Tremfya subcutaneous, or Xeljanz/XR <b>[documentation required]</b>; 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</li></ul></li></ul>
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	<p>multiple adalimumab products counts as <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), Simponi (Aria or subcutaneous), Cosentyx, or Bimzelx also counts <b>[documentation required]</b>.</p> <p>b) Patient is &lt; 18 years of age AND has tried ONE of Enbrel, Rinvoq/Rinvoq LQ, or Stelara SC <b>[documentation required]</b>; OR <u>Note:</u> A trial of another TNFi counts towards a trial of Enbrel <b>[documentation required]</b>. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product.</p> <p>c) According to the prescriber, the patient has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, OR a demyelinating disorder.</p> <p>C) 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: offer to review for a Step 1 or Step 2a Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>4. <u>Rheumatoid Arthritis, Juvenile Idiopathic Arthritis, or Psoriatic Arthritis – Patient is Currently Receiving Orencia (Subcutaneous or Intravenous).</u></b></p> <p><b>B) Approve for 1 year if the patient meets BOTH of the following (i and ii):</b></p> <p>i. Patient meets the standard <i>Inflammatory Conditions – Orencia Subcutaneous Policy</i> criteria; AND</p> <p>ii. Patient meets ONE of the following (a, b, c, d, e, f, or g):</p> <p>a) Patient has <u>Rheumatoid Arthritis</u> and has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR <b>[documentation required]</b>; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> 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 <b>[documentation required]</b>.</p> <p>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 <b>[documentation required]</b>; OR</p>
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	<p><u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz tablets and Xeljanz oral solution) collectively counts as <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product. A trial of Cimzia, tocilizumab intravenous (Actemra intravenous, biosimilar), Kevzara, Orencia intravenous, an infliximab product (e.g., Remicade, biosimilars), or Simponi Aria also counts <b>[documentation required]</b>.</p> <p>c) Patient is <math>\geq 18</math> years of age with <u>Psoriatic Arthritis</u> AND has tried TWO of Enbrel, an adalimumab product, Otezla, Rinvoq/Rinvoq LQ, Skyrizi subcutaneous, Stelara subcutaneous, Taltz, Tremfya subcutaneous, or Xeljanz/XR <b>[documentation required]</b>; OR</p> <p><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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> product. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), Simponi (Aria or subcutaneous), Cosentyx, or Bimzelx also counts <b>[documentation required]</b>.</p> <p>d) Patient is <math>&lt; 18</math> years of age with <u>Psoriatic Arthritis</u> AND has tried ONE of Enbrel, Rinvoq/Rinvoq LQ, or Stelara subcutaneous<b>[documentation required]</b>; OR</p> <p><u>Note:</u> A trial of another TNFi counts towards a trial of Enbrel <b>[documentation required]</b>. A trial of either or both Rinvoq products (Rinvoq and Rinvoq LQ) collectively counts as <b>ONE</b> product.</p> <p>e) According to the prescriber, the patient has been established on Orencia intravenous for at least 90 days; OR</p> <p>f) According to the prescriber, the patient has heart failure, a previously treated lymphoproliferative disorder, a previous serious infection, OR a demyelinating disorder; OR</p> <p>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> <b>[verification in prescription claims history required]</b>, or if claims history is not available, according to the prescriber <b>[verification by prescriber required]</b>.</p> <p><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</p>
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	<p>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).</p> <p>C) If the patient has met criterion 4Ai (the standard <i>Inflammatory Conditions – Orencia Subcutaneous Prior Authorization Policy</i> criteria), but criterion 4Aii is not met, offer to review for one of the following Products using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p>i. <b>Rheumatoid Arthritis:</b> <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR.</u></p> <p>ii. <b>Juvenile Idiopathic Arthritis:</b> <u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Rinvoq LQ, Xeljanz tablets, or Xeljanz oral solution.</u></p> <p>iii. <b>Psoriatic Arthritis in a Patient ≥ 18 Years of Age:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Rinvoq, Rinvoq LQ, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Taltz, Tremfya subcutaneous, Xeljanz tablets, or Xeljanz XR.</u></p> <p>iv. <b>Psoriatic Arthritis in a Patient &lt; 18 Years of Age:</b> <u>Enbrel, Rinvoq, Rinvoq LQ, or Stelara subcutaneous.</u></p> <p>5. <b>Other Conditions.</b> 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.</p>
<b>Janus Kinases Inhibitors</b>	
<p><b>Olumiant</b></p>	<p>1. <b>Rheumatoid Arthritis – Initial Therapy.</b></p> <p>A) Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <p>i. Patient meets the standard <i>Inflammatory Conditions – Olumiant Prior Authorization Policy</i> criteria; AND</p> <p>ii. Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, or Xeljanz/XR <b>[documentation required]</b>.  <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> 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 <b>[documentation required]</b>.</p>

	<p><b>B)</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: offer to review for a Step 1 or Step 2a Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>2. <u>Rheumatoid Arthritis – Patient is Currently Receiving Olumiant.</u></b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Olumiant Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient meets ONE of the following (a <u>or</u> b):<ul style="list-style-type: none"><li><b>a)</b> Patient has tried TWO of a tocilizumab subcutaneous product, Enbrel, an adalimumab product, Rinvoq, and Xeljanz/XR [<b>documentation required</b>]; OR <u>Note:</u> Examples of tocilizumab subcutaneous products include Actemra subcutaneous and Tyenne subcutaneous. A trial of multiple tocilizumab products counts as <b>ONE</b> 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 <b>ONE</b> product. A trial of either or both Xeljanz products (Xeljanz and Xeljanz XR) collectively counts as <b>ONE</b> 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 [<b>documentation required</b>].</li><li><b>b)</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> [<b>verification in prescription claims history required</b>], or if claims history is not available, according to the prescriber [<b>verification by prescriber required</b>]. <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).</li></ul></li></ul> <p><b>B)</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: offer to review for a Step 1 or Step 2a Product (<u>Actemra subcutaneous, Tyenne subcutaneous, Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Rinvoq, Xeljanz tablets, or Xeljanz XR</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p>
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	<p><b>3. <u>Other Conditions.</u></b> 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.</p>
<p><b>Rinvoq</b></p>	<p><b>1. <u>Ankylosing Spondylitis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried one of Enbrel or an adalimumab product; OR</li> </ul> <p><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.</p> <p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, or Taltz</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>2. <u>Crohn’s Disease – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried one adalimumab product.</li> </ul> <p><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.</p> <p><b>B)</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: offer to review for a Preferred Product (<u>adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Skyrizi subcutaneous [on-body injector], Stelara subcutaneous, or Zymfentra</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>3. <u>Juvenile Idiopathic Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried one of Enbrel or an adalimumab product; OR</li> </ul> <p><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.</p> <p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo,</u></p>

	<p><u>adalimumab-adaz, adalimumab-ryvk, or Simlandi</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>4. <u>Non-Radiographic Spondyloarthritis (nr-axSpA) – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ol style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient has tried Cimzia.</li></ol> <p><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.</p> <p><b>B)</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: offer to review for a Preferred Product (<u>Cimzia or Taltz</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>5. <u>Rheumatoid Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ol style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient has tried one of Enbrel or an adalimumab product; OR</li></ol> <p><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.</p> <p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>6. <u>Psoriatic Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ol style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient has tried one of Enbrel or an adalimumab product; OR</li></ol> <p><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.</p> <p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Taltz, or Tremfya subcutaneous</u>) using</p>
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	<p>the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>7. <u>Ulcerative Colitis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> 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.</li></ul> <p><b>B)</b> If the patient has met criterion 7Ai (the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria), but criterion 7Aii is not met: offer to review for a Preferred Product (<u>adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Tremfya subcutaneous, Velsipity, or Zymfentra</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>8. <u>Ankylosing Spondylitis, Crohn’s Disease, Juvenile Idiopathic Arthritis, nr-axSpA, Rheumatoid Arthritis, Psoriatic Arthritis, or Ulcerative Colitis – Patient is Currently Receiving Rinvoq.</u></b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient meets ONE of the following (a, b, c, d, e, f, g, <u>or</u> h):<ul style="list-style-type: none"><li><b>a)</b> Patient has <u>Ankylosing Spondylitis</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.</li><li><b>b)</b> Patient has <u>Crohn’s Disease</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 Cimzia also counts.</li><li><b>c)</b> 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</li></ul></li></ul>
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	<p>Cimzia, an infliximab product (e.g., Remicade, biosimilars) or Simponi Aria also counts.</p> <p><b>d)</b> 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.</p> <p><b>e)</b> 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.</p> <p><b>f)</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.</p> <p><b>g)</b> 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.</p> <p><b>h)</b> 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> [<b>verification in prescription claims history required</b>], or if claims history is not available, according to the prescriber [<b>verification by prescriber required</b>]. <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).</p> <p><b>B)</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:</p>
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	<p>offer to review for one of the following Products using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria:</p> <ul style="list-style-type: none"> <li>i. <b>Ankylosing Spondylitis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, or Taltz.</u></li> <li>ii. <b>Crohn’s Disease:</b> <u>adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, or Zymfentra.</u></li> <li>iii. <b>Juvenile Idiopathic Arthritis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi.</u></li> <li>iv. <b>nr-axSpA:</b> <u>Cimzia or Taltz.</u></li> <li>v. <b>Rheumatoid Arthritis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi.</u></li> <li>vi. <b>Psoriatic Arthritis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Taltz, or Tremfya subcutaneous.</u></li> <li>vii. <b>Ulcerative Colitis:</b> <u>adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Tremfya subcutaneous, Velsipity, or Zymfentra.</u></li> </ul> <p>9. <b>All Other Conditions.</b> 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.</p>
<p><b>Rinvoq LQ</b></p>	<ul style="list-style-type: none"> <li>1. <b>Juvenile Idiopathic Arthritis – Initial Therapy.</b> <ul style="list-style-type: none"> <li>A) Approve for 6 months if the patient meets BOTH of the following (i and ii): <ul style="list-style-type: none"> <li>i. Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND</li> <li>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.</li> </ul> </li> <li>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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</li> </ul> </li> <li>2. <b>Psoriatic Arthritis – Initial Therapy.</b> <ul style="list-style-type: none"> <li>A) Approve for 6 months if the patient meets BOTH of the following (i and ii): <ul style="list-style-type: none"> <li>i. Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND</li> <li>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.</li> </ul> </li> </ul> </li> </ul>

	<p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Taltz, or Tremfya subcutaneous</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>3. <u>Juvenile Idiopathic Arthritis or Psoriatic Arthritis – Patient is Currently Receiving Rinvoq/LQ.</u></b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ul style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Rinvoq/LQ Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient meets ONE of the following conditions (a, b, <u>or</u> c):<ul style="list-style-type: none"><li><b>a)</b> 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.</li><li><b>b)</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.</li><li><b>c)</b> 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 [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 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).</li></ul></li></ul> <p><b>B)</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: offer to review for one of the following Products using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria:</p> <ul style="list-style-type: none"><li><b>i. <u>Juvenile Idiopathic Arthritis:</u></b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi.</u></li></ul>
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	<p><b>ii. Psoriatic Arthritis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Taltz, or Tremfya subcutaneous.</u></p> <p><b>4. Other Conditions.</b> 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.</p>
<p><b>Xeljanz tablets, Xeljanz XR tablets</b></p>	<p><b>1. Ankylosing Spondylitis – Initial Therapy.</b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried one of Enbrel or an adalimumab product; OR</li> </ul> <p><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.</p> <p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, or Taltz</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>2. Rheumatoid Arthritis – Initial Therapy.</b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried one of Enbrel or an adalimumab product; OR</li> </ul> <p><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.</p> <p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>3. Juvenile Idiopathic Arthritis – Initial Therapy.</b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried one of Enbrel or an adalimumab product; OR</li> </ul> <p><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.</p>

	<p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>4. <u>Psoriatic Arthritis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ol style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient has tried one of Enbrel or an adalimumab product; OR</li></ol> <p><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.</p> <p><b>B)</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: offer to review for a Step 1 Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous [pen or syringe], Stelara subcutaneous, Taltz, or Tremfya subcutaneous</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>5. <u>Ulcerative Colitis – Initial Therapy.</u></b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ol style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient has tried one adalimumab product.</li></ol> <p><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.</p> <p><b>B)</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: offer to review for a Preferred Product (<u>adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Tremfya subcutaneous, Velsipity, or Zymfentra</u>) using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria.</p> <p><b>6. <u>Ankylosing Spondylitis, Rheumatoid Arthritis, Juvenile Idiopathic Arthritis, Psoriatic Arthritis, or Ulcerative Colitis – Patient is Currently Receiving Xeljanz/XR.</u></b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i <u>and</u> ii):</p> <ol style="list-style-type: none"><li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND</li><li><b>ii.</b> Patient meets ONE of the following (a, b, c, d, e, <u>or</u> f):</li></ol>
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	<p>a) Patient has <u>Ankylosing Spondylitis</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.</p> <p>b) 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.</p> <p>c) 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.</p> <p>d) 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.</p> <p>e) 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.</p> <p>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 [verification in prescription claims history required]</u>, or if claims history is not available, according to the prescriber <u>[verification by prescriber required]</u>; 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</p>
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	<p>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).</p> <p><b>B)</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: offer to review for one of the following Products using the respective standard <i>Inflammatory Conditions Prior Authorization Policy</i> criteria:</p> <ul style="list-style-type: none"> <li><b>i. Ankylosing Spondylitis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, or Taltz.</u></li> <li><b>ii. Rheumatoid Arthritis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi.</u></li> <li><b>iii. Juvenile Idiopathic Arthritis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi.</u></li> <li><b>iv. Psoriatic Arthritis:</b> <u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Otezla, Skyrizi subcutaneous (pen or syringe), Stelara subcutaneous, Taltz, or Tremfya subcutaneous.</u></li> <li><b>v. Ulcerative Colitis:</b> <u>adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, Simlandi, Omvoh subcutaneous, Skyrizi subcutaneous (on-body injector), Stelara subcutaneous, Tremfya subcutaneous, Velsipity, or Zymfentra.</u></li> </ul> <p><b>7. Other Conditions.</b> 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.</p>
<p><b>Xeljanz oral solution</b></p>	<p><b>1. Juvenile Idiopathic Arthritis – Initial Therapy.</b></p> <p><b>A)</b> Approve for 6 months if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient has tried one of Enbrel or an adalimumab product; OR</li> </ul> <p><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.</p> <p><b>B)</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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p><b>2. Juvenile Idiopathic Arthritis – Patient is Currently Receiving Xeljanz.</b></p> <p><b>A)</b> Approve for 1 year if the patient meets BOTH of the following (i and ii):</p> <ul style="list-style-type: none"> <li><b>i.</b> Patient meets the standard <i>Inflammatory Conditions – Xeljanz/XR Prior Authorization Policy</i> criteria; AND</li> <li><b>ii.</b> Patient meets ONE of the following (a or b): <ul style="list-style-type: none"> <li><b>a)</b> Patient has <u>Juvenile Idiopathic Arthritis</u> and has tried one of Enbrel or an adalimumab product; OR</li> </ul> </li> </ul> <p><u>Note:</u> Examples of adalimumab products include Humira, Abrilada, adalimumab-adaz, adalimumab-adbm, adalimumab-fkjp, adalimumab-aaty, adalimumab-ryvk, Simlandi, Amjevita, Cyltezo,</p>

	<p>Hadlima, Hulio, Hyrimoz, Idacio, Yuflyma, and Yusimry. A trial of Cimzia, an infliximab product (e.g., Remicade, biosimilars), or Simponi Aria also counts.</p> <p>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 <u>[verification by prescriber required]</u>; OR</p> <p><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).</p> <p>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: offer to review for a Preferred Product (<u>Enbrel, adalimumab-adbm, Cyltezo, adalimumab-adaz, adalimumab-ryvk, or Simlandi</u>) using the respective standard <i>Inflammatory Conditions – Prior Authorization Policy</i> criteria.</p> <p>3. <b>Other Conditions.</b> 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.</p>
<b>Sphingosine 1-Phosphate Receptor Modulator</b>	
<b>Zeposia</b>	<b>All Conditions.</b> Approve <u>Zeposia</u> if the patient meets the standard <i>Multiple Sclerosis and Ulcerative Colitis – Zeposia Care Value Policy</i> criteria.

**REFERENCES**

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2. Cimzia<sup>®</sup> subcutaneous injection [prescribing information]. Smyrna, GA: UCB; March 2021.
3. Cosentyx<sup>®</sup> subcutaneous injection [prescribing information]. East Hanover, NJ: Novartis; June 2020.
4. Enbrel<sup>®</sup> subcutaneous injection [prescribing information]. Thousand Oaks, CA: Amgen; June 2023.
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7. Kevzara<sup>™</sup> subcutaneous injection [prescribing information]. Tarrytown, NY: Regeneron/sanofi Aventis; April 2018.
8. Kineret<sup>®</sup> subcutaneous injection [prescribing information]. Thousand Oaks, CA: Swedish Orphan Biovitrium; December 2020.
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12. Renflexis<sup>®</sup> intravenous injection [prescribing information]. Whitehouse Station, NJ: Merck/Samsung Bioepsis; March 2021.
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14. Siliq<sup>™</sup> subcutaneous injection [prescribing information]. Bridgewater, NJ: Valeant; June 2020.
15. Simponi<sup>®</sup> subcutaneous injection [prescribing information]. Horsham, PA: Janssen Biotech; September 2019.
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18. Taltz<sup>®</sup> subcutaneous injection [prescribing information]. Indianapolis, IN: Eli Lilly; March 2021.
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20. Xeljanz<sup>®</sup>/Xeljanz XR tablets/extended release tablets [prescribing information]. New York, NY: Pfizer; October 2020.
21. Ilumya<sup>™</sup> subcutaneous injection [prescribing information]. Whitehouse Station, NJ: Sun/Merck; April 2021.

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22. Rinvoq<sup>®</sup> tablets/Rinvoq LQ oral solution [prescribing information]. North Chicago, IL: AbbVie; April 2024.
  23. Zeposia<sup>®</sup> capsules [prescribing information]. Summit, NJ: Celgene; May 2021.
  24. Sotyktu<sup>™</sup> tablets [prescribing information]. Princeton, NJ: Bristol Myers Squibb; September 2022.
  25. Velsipity<sup>®</sup> tablets [prescribing information]. New York, NY: Pfizer; October 2023.
  26. Omvoh<sup>™</sup> intravenous infusion and subcutaneous injection [prescribing information]. Indianapolis, IN: Eli Lilly; October 2023.
  27. Entyvio<sup>®</sup> subcutaneous injection and intravenous infusion [prescribing information]. Lexington, MA: Takeda; September 2023.
  28. Zymfentra<sup>™</sup> subcutaneous injection [prescribing information]. Yeonsu-gu, Incheon: Celltrion; October 2023.
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**HISTORY**

Type of Revision	Summary of Changes	Review Date
Annual Revision	<p><b>Effective 01/01/2025</b>                      A descriptor of Choice/Alternate was added to the policy name.</p> <p><b>Humira:</b> Throughout the policy, NDCs starting with 00074 were removed from the Preferred Products. A previous trial of these NDCs counts towards a trial of an adalimumab product.</p> <p><b>Hyrimoz:</b> 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.</p> <p><b>Tremfya Subcutaneous:</b> For <b>Ulcerative Colitis</b>, Tremfya subcutaneous was added as a Preferred Product.</p> <p><b>OmvoH Subcutaneous:</b> For <b>Ulcerative Colitis</b>, OmvoH subcutaneous was moved from Step 2a to Preferred (Step 1).</p> <p><b>Cimzia:</b> For <b>Rheumatoid Arthritis, Ankylosing Spondylitis, Psoriatic Arthritis, Plaque Psoriasis, and Crohn’s Disease</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products. For <b>Juvenile Idiopathic Arthritis</b>, 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 <b>Psoriatic Arthritis</b> and <b>Plaque Psoriasis</b>, it was clarified that Tremfya is the subcutaneous formulation.</p> <p><b>Simponi Subcutaneous:</b> For <b>Rheumatoid Arthritis, Ankylosing Spondylitis, Psoriatic Arthritis, and Ulcerative Colitis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products. For <b>Psoriatic Arthritis</b>, it was clarified that Tremfya is the subcutaneous formulation. For <b>Ulcerative Colitis</b>, Tremfya subcutaneous and OmvoH subcutaneous were added as Preferred Products.</p> <p><b>Actemra Subcutaneous and Tyenne Subcutaneous:</b> For <b>Rheumatoid Arthritis and Polyarticular Juvenile Idiopathic Arthritis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products. For <b>Polyarticular Juvenile Idiopathic Arthritis</b>, Cimzia was added as an agent that counts towards a trial of a Preferred Product.</p> <p><b>Kevzara:</b> For <b>Rheumatoid Arthritis and Juvenile Idiopathic Arthritis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products. For <b>Juvenile Idiopathic Arthritis</b>, Cimzia was added as an agent that counts towards a trial of a Preferred Product.</p> <p><b>Bimzelx:</b> For <b>Ankylosing Spondylitis, Non-Radiographic Spondyloarthritis, and Psoriatic Arthritis</b>, Bimzelx was added to Step 2a and requests are directed to a trial of one Step 1 Product. For <b>Plaque Psoriasis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products.</p> <p><b>Cosentyx Subcutaneous:</b> For <b>Ankylosing Spondylitis, Psoriatic Arthritis, and Plaque Psoriasis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products. For <b>Psoriatic Arthritis</b> and <b>Plaque Psoriasis</b>, it was clarified that Tremfya is the subcutaneous formulation.</p> <p><b>Siliq:</b> For <b>Plaque Psoriasis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products, and it was clarified that Tremfya is the subcutaneous formulation.</p> <p><b>Ilumya:</b> For <b>Plaque Psoriasis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products, and it was clarified that Tremfya is the subcutaneous formulation.</p> <p><b>Entyvio Subcutaneous:</b> For <b>Crohn’s Disease and Ulcerative Colitis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products. For <b>Ulcerative Colitis</b>, Tremfya subcutaneous was added as a Preferred Product; a previous trial of Tremfya intravenous also counts.</p> <p><b>Kineret:</b> For <b>Rheumatoid Arthritis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products.</p> <p><b>Orencia Subcutaneous:</b> For <b>Rheumatoid Arthritis, Juvenile Idiopathic Arthritis, and Psoriatic Arthritis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products. For <b>Juvenile Idiopathic Arthritis</b>, Cimzia was added as an agent that counts towards a trial of a Preferred Product. For <b>Psoriatic Arthritis</b>, it was clarified that Tremfya is the subcutaneous formulation; for a patient <math>\geq 18</math></p>	10/30/2024

	<p>years of age, Cosentyx and Bimzelx were added as agents that count towards a trial of a Preferred Product.</p> <p><b>Olumiant:</b> For <b>Rheumatoid Arthritis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products.</p> <p><b>Rinvoq:</b> For <b>Rheumatoid Arthritis, Ankylosing Spondylitis, Juvenile Idiopathic Arthritis, Psoriatic Arthritis, Crohn’s Disease, and Ulcerative Colitis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products. For <b>Juvenile Idiopathic Arthritis</b>, Cimzia was added as an agent that counts towards a trial of a Preferred Product. For <b>Psoriatic Arthritis</b>, it was clarified that Tremfya is the subcutaneous formulation. For <b>Ulcerative Colitis</b>, Tremfya subcutaneous and Omvoh subcutaneous were added as Preferred Products.</p> <p><b>Rinvoq LQ:</b> For <b>Juvenile Idiopathic Arthritis and Psoriatic Arthritis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products. For <b>Juvenile Idiopathic Arthritis</b>, Cimzia was added as an agent that counts towards a trial of a Preferred Product. For <b>Psoriatic Arthritis</b>, it was clarified that Tremfya is the subcutaneous formulation.</p> <p><b>Xeljanz/Xeljanz XR:</b> For <b>Rheumatoid Arthritis, Ankylosing Spondylitis, Juvenile Idiopathic Arthritis</b> (Xeljanz tablets only), <b>Psoriatic Arthritis, and Ulcerative Colitis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products. For <b>Juvenile Idiopathic Arthritis</b>, Cimzia was added as an agent that counts towards a trial of a Preferred Product. For <b>Psoriatic Arthritis</b>, it was clarified that Tremfya is the subcutaneous formulation. For <b>Ulcerative Colitis</b>, Tremfya subcutaneous and Omvoh subcutaneous were added as Preferred Products.</p> <p><b>Xeljanz Oral Solution:</b> For <b>Juvenile Idiopathic Arthritis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products. For <b>Juvenile Idiopathic Arthritis</b>, Cimzia was added as an agent that counts towards a trial of a Preferred Product.</p> <p><b>Velsipity:</b> For <b>Ulcerative Colitis</b>, Humira (NDCs starting with 00074) and Hyrimoz (NDCs starting with 61314) were removed from the Preferred Products; Tremfya subcutaneous was added as a Preferred Product; a previous trial of Tremfya intravenous also counts.</p>	
Selected Revision	<p><b>Effective 01/01/2025.</b></p> <p><b>Velsipity:</b> For <b>Ulcerative Colitis</b>, Velsipity was added as a Preferred Product.</p> <p><b>Simponi Subcutaneous:</b> For <b>Ulcerative Colitis</b>, Velsipity was added as a Preferred Product.</p> <p><b>Rinvoq:</b> For <b>Ulcerative Colitis</b>, Velsipity was added as a Preferred Product.</p> <p><b>Xeljanz/XR:</b> For <b>Ulcerative Colitis</b>, Velsipity was added as a Preferred Product.</p> <p><b>Entyvio Subcutaneous:</b> For <b>Ulcerative Colitis</b>, Velsipity was added as a Preferred Product.</p>	11/20/2024
Selected Revision	<p><b>Effective 01/01/2025.</b></p> <p><b>Hidradenitis Suppurativa</b> was added as a targeted indication in this policy. Adalimumab products (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.</p>	12/04/2025

**APPENDIX A**

**Table 1. Approved TNFis for Targeted Indications.\***

	Rheumatology					Dermatology		Gastroenterology	
	RA	JIA	AS	nr-axSpA	PsA	HS	PsO	CD	UC
<b>Tumor Necrosis Factor Inhibitors</b>									
Cimzia	√	√	√	√	√	--	√	√	--
Enbrel	√	√	√	--	√	--	√	--	--
Adalimumab Products (Humira, biosimilars)	√	√	√	--	√	√	√	√	√
Infliximab Intravenous Products	√	--	√	--	√	--	√	√	√
Zymfentra	--	--	--	--	--	--	--	√ <sup>^</sup>	√ <sup>^</sup>
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; <sup>^</sup> 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
<b>Interleukin-17 Blockers</b>							
Bimzelx	√	√	√	√	√	--	--
Cosentyx Subcutaneous	√	√	√	√	√	--	--
Cosentyx Intravenous	√	√	√	--	--	--	--
Siliq	--	--	--	--	√	--	--
Taltz	√	√	√	--	√	--	--
<b>Interleukin-23 Blockers</b>							
Ilumya	--	--	--	--	√	√	--
OmvoH Intravenous	--	--	--	--	--	--	√ <sup>#</sup>
OmvoH Subcutaneous	--	--	--	--	--	--	√ <sup>^</sup>
Skyrizi Intravenous	--	--	--	--	--	√ <sup>#</sup>	√ <sup>#</sup>
Skyrizi Subcutaneous	--	--	√	--	√	√ <sup>^</sup>	√ <sup>^</sup>
Tremfya Intravenous	--	--	--	--	--	--	√ <sup>#</sup>
Tremfya Subcutaneous	--	--	√	--	√	--	√ <sup>^</sup>
<b>Interleukin-12/23 Blockers</b>							
Stelara Subcutaneous	--	--	√	--	√	√ <sup>^</sup>	√ <sup>^</sup>
Stelara Intravenous	--	--	--	--	--	√ <sup>#</sup>	√ <sup>#</sup>

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; <sup>^</sup> Maintenance dosing only; <sup>#</sup> Induction dosing only.

**Table 3. Approved Oral tsDMARDs for Targeted Indications.\***

	Rheumatology					Dermatology	Gastroenterology	
	RA	JIA	AS	nr-axSpA	PsA	PsO	CD	UC
<b>Janus Kinases Inhibitors</b>								
Olumiant	√	--	--	--	--	--	--	--
Rinvoq	√	√	√	√	√	--	√	√
Rinvoq LQ	--	√	--	√	--	--	--	--
Xeljanz tablets	√	√ <sup>#</sup>	√	--	√	--	--	√
Xeljanz oral solution	--	√ <sup>#</sup>	--	--	--	--	--	--
Xeljanz XR	√	--	√	--	√	--	--	√
<b>Phosphodiesterase Type 4 Inhibitor</b>								
Otezla	--	--	--	--	√	√	--	--
<b>Sphingosine 1-Phosphate Receptor Modulator</b>								
Velsipity	--	--	--	--	--	--	--	√
Zeposia	--	--	--	--	--	--	--	√
<b>Tyrosine Kinase 2 Inhibitor</b>								
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
<b>Integrin Receptor Antagonist</b>					
Entyvio Intravenous	--	--	--	√	√
Entyvio Subcutaneous	--	--	--	√ <sup>‡</sup>	√ <sup>‡</sup>
<b>Interleukin-6 Blockers</b>					
Tocilizumab Intravenous Products (Actemra, biosimilar)	√	√ <sup>^</sup>	--	--	--
Tocilizumab Subcutaneous Products (Actemra, biosimilar)	√	√ <sup>^</sup>	--	--	--
Kevzara	√	√	--	--	--
<b>Interleukin-1 Blocker</b>					
Kineret	√	--	--	--	--
<b>T-Cell Costimulation Modulator</b>					
Orencia Intravenous	√	√ <sup>#</sup>	√	--	--
Orencia Subcutaneous	√	√ <sup>#</sup>	√	--	--
<b>CD20-Directed Cytolytic Antibody</b>					
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.