

Clinical Policy: Step Therapy

Reference Number: MCPB.ST.00

Effective Date: 01.01.22

Last Review Date: 08.22

Line of Business: Medicare Part B

See [Important Reminder](#) at the end of this policy for important regulatory and legal information.

Description

This policy provides a list of drugs that require step therapy. Step therapy is when we require the trial of a preferred therapeutic alternative prior to coverage of a non-preferred drug for a specific indication.

FDA Approved Indication(s)

Various.

Policy/Criteria

This policy does not replace existing Medicare rules and regulations for the applicable agent(s).

I. Approval Criteria (NEW STARTS ONLY – member has not received the drug for the past 365 days)

A. Step Therapy:

Drug Name	Part B Required Step-Through Agents* By Indication <i>*May require prior authorization</i>
Abatacept (Orencia®)	PART B STEP: <ul style="list-style-type: none"> All indications: a tumor necrosis factor (TNF) inhibitor (e.g., infliximab)* <i>(note credit may be given if another TNF inhibitor was tried)</i>
Ado-trastuzumab emtansine (Kadcyla®)	PART B STEP: <ul style="list-style-type: none"> Breast cancer: trastuzumab-based therapy* and a taxane* <i>(note some IV chemo may not require prior authorization)</i>
Aflibercept (Eylea®)	PART B STEP: <ul style="list-style-type: none"> Neovascular (wet) age-related macular degeneration (AMD), macular edema following retinal vein occlusion (RVO), diabetic macular edema (DME), or diabetic retinopathy (DR): intravitreal bevacizumab solution
Atezolizumab (Tecentriq®)	PART B STEP: <ul style="list-style-type: none"> Urothelial carcinoma: member is ineligible for platinum-containing chemotherapy as first-line systemic therapy* <i>(note some IV chemo may not require prior authorization)</i> Non-small cell lung cancer that is high-risk stage IIA with programmed death-ligand 1 (PD-L1)

Drug Name	Part B Required Step-Through Agents* By Indication <i>*May require prior authorization</i>
	<p>expression $\geq 1\%$ OR is recurrent, advanced, or metastatic and anaplastic lymphoma kinase (ALK) or epidermal growth factor receptor (EGFR) mutation negative or unknown: prior platinum-containing chemotherapy (<i>note some IV chemo may not require prior authorization</i>), UNLESS one of the following is met:</p> <ul style="list-style-type: none"> ○ Request is for use as a single agent, and disease is stage II to IIIA with previous resection ○ Request is for use as a single agent as first-line therapy for tumors that have high PD-L1 expression, defined as PD-L1 $\geq 50\%$ (tumor cells [TC] $\geq 50\%$) or tumor-infiltrating immune cells (IC) covering $\geq 10\%$ of the tumor area [IC $\geq 10\%$] ○ Disease is non-squamous, and Tecentriq is prescribed as combination therapy ○ No prior progression on a programmed death receptor-1 (PD-1) or PD-L1 inhibitor (e.g., Tecentriq, nivolumab, pembrolizumab, durvalumab), and Tecentriq is prescribed as single agent as subsequent therapy
<p>Axicabtagene ciloleucel (Yescarta[®])</p>	<p>PART B STEP:</p> <ul style="list-style-type: none"> ● Large B-cell lymphoma: one of the following: <ul style="list-style-type: none"> ○ 2 lines of systemic therapy that includes rituximab* and one anthracycline-containing regimen (e.g., doxorubicin) ○ First-line chemoimmunotherapy that includes an anti-CD20 monoclonal antibody (e.g., rituximab*) and anthracycline-containing regimen (e.g., doxorubicin), if disease was refractory (defined as no complete remission) to or relapsed (defined as complete remission followed by biopsy-proven disease relapse) no more than 12 months after chemoimmunotherapy ● Relapsed or refractory follicular lymphoma: 2 lines of systemic therapy that includes a combination of an anti-CD20 monoclonal antibody* (e.g., rituximab or Gazyva) and an alkylating agent (e.g., bendamustine, cyclophosphamide, chlorambucil) <p><i>Only for initial treatment dose; subsequent doses will not be covered</i></p>
<p>Bevacizumab (Avastin[®], Mvasi[®], Zirabev[™])</p>	<p>PART B STEP:</p>

Drug Name	Part B Required Step-Through Agents* By Indication <i>*May require prior authorization</i>
	<ul style="list-style-type: none"> Oncology indications, if request is for Avastin: Mvasi and Zirabev
Brexucabtagene autoleucel (Tecartus™)	PART B STEP: <ul style="list-style-type: none"> Mantle cell lymphoma: 2 to 5 prior regimens that included all of the following: anthracycline (e.g., doxorubicin*) or bendamustine*-containing chemotherapy; anti-CD20 monoclonal antibody therapy (e.g., rituximab*) B-cell precursor acute lymphoblastic leukemia: at least two prior systemic therapies* <i>Only for initial treatment dose; subsequent doses will not be covered</i>
Brolucizumab-dblb (Beovu®)	PART B STEP: <ul style="list-style-type: none"> Neovascular (wet) AMD, DME: intravitreal bevacizumab solution
Cemiplimab-rwlc (Libtayo®)	PART B STEP: <ul style="list-style-type: none"> Cutaneous squamous cell carcinoma: cisplatin*, unless curative radiation therapy or surgery is not feasible
Certolizumab (Cimzia®)	PART B STEP: <ul style="list-style-type: none"> All indications: a different TNF inhibitor (e.g., infliximab)* (<i>note credit may be given if another TNF inhibitor was tried</i>)
Corticosteroid intravitreal implants: dexamethasone (Ozurdex®), fluocinolone acetonide (Iluvien®, Retisert®, Yutiq™)	PART B STEP: <ul style="list-style-type: none"> Macular edema following branch or central RVO (Ozurdex only): intravitreal bevacizumab solution and intravitreal corticosteroid injection (e.g., Triesence) DME (Ozurdex or Iluvien): intravitreal bevacizumab solution and intravitreal corticosteroid injection (e.g., Triesence) Non-infectious uveitis (Ozurdex, Retisert, or Yutiq): intravitreal corticosteroid injection (e.g., Triesence)
Corticotropin (H.P. Acthar®, Purified Cortrophin™ Gel)	PART B STEP: <ul style="list-style-type: none"> All indications, except infantile spasms, if request is for H.P. Acthar: Purified Cortrophin Gel IN ADDITION: <ul style="list-style-type: none"> Multiple sclerosis: corticosteroid
Daratumumab (Darzalex®), daratumumab/hyaluronidase-fihj (Darzalex Faspro™)	PART B STEP: <ul style="list-style-type: none"> Multiple myeloma: 1 prior systemic therapy (e.g., ixazomib*, bortezomib*, carfilzomib*) (<i>note some IV chemo may not require prior authorization</i>) if prescribed in combination with dexamethasone and either lenalidomide, bortezomib, or carfilzomib; OR 2

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	<p>prior systemic therapies (e.g., ixazomib*, bortezomib*, carfilzomib*) if prescribed as monotherapy or in combination with pomalidomide and dexamethasone; UNLESS Darzalex is prescribed as primary therapy in one of the following ways:</p> <ul style="list-style-type: none"> ○ In combination with lenalidomide and dexamethasone or bortezomib, melphalan, and prednisone, and member is ineligible for autologous stem cell transplant (ASCT); or ○ In combination with bortezomib, thalidomide, and dexamethasone, and member is eligible for ASCT <ul style="list-style-type: none"> ● Systemic light chain amyloidosis (<i>Darzalex only</i>): 1 prior systemic therapy (e.g., bortezomib*) (<i>note some IV chemo may not require prior authorization</i>)
Darbepoetin alfa (Aranesp [®])	<p>PART B STEP:</p> <ul style="list-style-type: none"> ● All indications: Retacrit <ul style="list-style-type: none"> ○ If Retacrit is unavailable due to shortage: Epogen
Denosumab (Xgeva [®])	<p>PART B STEP:</p> <ul style="list-style-type: none"> ● Systemic mastocytosis, hypercalcemia of malignancy: zoledronic acid (Zometa)* or pamidronate*
Durvalumab (Imfinzi [®])	<p>PART B STEP:</p> <ul style="list-style-type: none"> ● Non-small cell lung cancer: chemotherapy (e.g., platinum-containing chemotherapy)* (<i>note some IV chemo may not require prior authorization</i>)
Elotuzumab (Empliciti [®])	<p>PART B STEP:</p> <ul style="list-style-type: none"> ● Multiple myeloma: prior line of systemic therapy (e.g., bortezomib*) (<i>note some IV chemo may not require prior authorization</i>)
Emapalumab-lzsg (Gamifant [™])	<p>PART B STEP:</p> <ul style="list-style-type: none"> ● Primary hemophagocytic lymphohistiocytosis (HLH): conventional HLH therapy* (<i>note some IV chemo may not require prior authorization</i>)
Epoetin alfa (Epogen [®] , Procrit [®])	<p>PART B STEP:</p> <ul style="list-style-type: none"> ● All indications: Retacrit <ul style="list-style-type: none"> ○ If Retacrit is unavailable due to shortage: Epogen
Filgrastim (Neupogen [®] , Zarxio [®] , Nivestym [™] , Granix [®])	<p>PART B STEP:</p> <ul style="list-style-type: none"> ● All indications, if request is for an agent other than Zarxio: Zarxio <ul style="list-style-type: none"> ○ If unable to use Zarxio and request is for Neupogen: biosimilar filgrastim product (e.g., Nivestym, Granix)
Golimumab (Simponi [®] , Simponi Aria [®])	<p>PART B STEP:</p>

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	<ul style="list-style-type: none"> • All indications: a different TNF inhibitor (e.g., infliximab)* (<i>note credit may be given if another TNF inhibitor was tried</i>)
<p>Hyaluronate derivatives: sodium hyaluronate (Euflexxa[®], Gelsyn-3[™], GenVisc[®] 850, Hyalgan[®], Supartz[™], Supartz FX[™], Synjoynt[™], Triluron[™], TriVisc[™], VISCO-3[™]), hyaluronic acid (Durolane[®]), cross-linked hyaluronate (Gel-One[®]), hyaluronan (Hymovis[®], Orthovisc[®], Monovisc[®]), hylan polymers A and B (Synvisc[®], Synvisc One[®])</p>	<p>PART B STEP:</p> <ul style="list-style-type: none"> • Osteoarthritis of the knee: intra-articular glucocorticoid injection*, and: <ul style="list-style-type: none"> ○ If request is for a product other than Synvisc/Synvisc One or Euflexxa: Synvisc*/Synvisc One* or Euflexxa*
<p>Idecabtagene vicleucel (Abecma[™])</p>	<p>PART B STEP:</p> <ul style="list-style-type: none"> • Multiple myeloma: 4 prior lines of therapy* that include all of the following: immunomodulatory agent (e.g., Revlimid, Pomalyst, Thalomid), proteasome inhibitor (e.g., bortezomib, Kyprolis), and anti-CD38 antibody (e.g., Darzalex/Darzalex Faspro, Sarclisa) <i>Only for initial treatment dose; subsequent doses will not be covered</i>
<p>Immune globulins (Asceniv[™], Bivigam[®], Carimune[®] NF, Cutaquig[®], Cuvitru[™], Flebogamma[®] DIF, GamaSTAN[®], GamaSTAN[®] S/D, Gammagard[®] liquid, Gammagard[®] S/D, Gammaked[™], Gammaplex[®], Gamunex[®]-C, Hizentra[®], HyQvia[®], Octagam[®], Panzyga[®], Privigen[®], Xembify[®])</p>	<p>PART B STEP:</p> <ul style="list-style-type: none"> • All indications except viral prophylaxis for hepatitis A, measles, varicella, or rubella viruses, if request is for an agent other than Gammagard: Gammagard* <p><u>IN ADDITION:</u></p> <ul style="list-style-type: none"> • Chronic idiopathic demyelinating polyneuropathy: a systemic corticosteroid, unless the member has pure motor symptoms • Polymyositis, myasthenia gravis, bullous pemphigoid, mucous membrane pemphigoid (a.k.a. cicatricial pemphigoid), epidermolysis bullosa acquisita: a systemic corticosteroid • Dermatomyositis: rituximab* • Idiopathic thrombocytopenic purpura: a systemic corticosteroid or Rho(D) immune globulin* • Pemphigus vulgaris, pemphigus foliaceus,: one corticosteroid and rituximab*

Drug Name	Part B Required Step-Through Agents* By Indication <i>*May require prior authorization</i>
	<ul style="list-style-type: none"> • Adenosine deaminase (ADA)-severe combined immunodeficiency disorders (SCID): Adagen* or Revcovi*
Infliximab (Remicade [®] , Avsola [™] , Renflexis [™] , Inflectra [®])	PART B STEP: <ul style="list-style-type: none"> • All indications: if request is Remicade: Avsola*, Inflectra*, and Renflexis*
Lisocabtagene maraleucel (Breyanzi [®])	PART B STEP: <ul style="list-style-type: none"> • Large B-cell lymphoma: one of the following: <ul style="list-style-type: none"> ○ 2 lines of systemic therapy that includes an anti-CD20 therapy (e.g., rituximab)* and one anthracycline-containing regimen (e.g., doxorubicin) ○ First-line chemoimmunotherapy that includes an anti-CD20 monoclonal antibody (e.g., rituximab*) and anthracycline-containing regimen (e.g., doxorubicin), if disease was refractory (defined as no complete remission) to or relapsed (defined as complete remission followed by biopsy-proven disease relapse) no more than 12 months after chemoimmunotherapy ○ If member is not eligible for hematopoietic stem cell transplantation due to comorbidities or age: First-line chemoimmunotherapy that includes an anti-CD20 monoclonal antibody (e.g., rituximab*) and anthracycline-containing regimen (e.g., doxorubicin), if disease was refractory (defined as no complete remission) to or has relapsed (defined as complete remission followed by biopsy-proven disease relapse) <p><i>Only for initial treatment dose; subsequent doses will not be covered</i></p>
Lurbinectedin (Zepzelca [™])	PART B STEP: <ul style="list-style-type: none"> • Small cell lung cancer: platinum-containing regimen (e.g., cisplatin, carboplatin)* (<i>note some IV chemo may not require prior authorization</i>)
Luspatercept-aamt (Reblozyl [®])	PART B STEP: <ul style="list-style-type: none"> • Myelodysplastic syndrome: erythropoiesis-stimulating agent used in combination with a granulocyte colony stimulating factor, unless current serum erythropoietin is greater than 500 mU/mL
Lutetium Lu 177 dotatate (Lutathera [®])	PART B STEP: <ul style="list-style-type: none"> • Neuroendocrine tumor: somatostatin analog (e.g., octreotide, lanreotide)
Natalizumab (Tysabri [®])	PART B STEP:

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	<ul style="list-style-type: none"> • Crohn's disease: a TNF inhibitor (e.g., infliximab*) (note credit may be given if another TNF inhibitor was tried)
Nivolumab (Opdivo [®])	<p>PART B STEP:</p> <ul style="list-style-type: none"> • Non-small cell lung cancer: prior systemic therapy*, UNLESS one of the following is met: <ul style="list-style-type: none"> ○ Tumor is positive for the tumor mutation burden (TMB) biomarker, or ○ Prescribed in combination with Yervoy for disease with RET rearrangement or unknown/negative mutation status for EGFR, ALK, ROS1, BRAF, MET exon 14 skipping, and NTRK gene fusion, or ○ Prescribed as neoadjuvant treatment • Malignant pleural mesothelioma: prior therapy*, unless prescribed in combination with Yervoy • Classical or pediatric Hodgkin lymphoma, anal carcinoma, vulvar cancer, extranodal NK/T-cell lymphoma - nasal type, small cell lung cancer, cervical cancer: prior therapy* • Squamous cell carcinoma of the head and neck: platinum-containing regimen* • Urothelial carcinoma: platinum-containing regimen*, unless prescribed as adjuvant treatment and member is at high risk of recurrence after undergoing resection, or member is at high risk of recurrence and did not previously receive a platinum-containing regimen • Esophageal squamous cell carcinoma: fluoropyrimidine-based (e.g., 5- fluorouracil, capecitabine) and platinum-based chemotherapy* • Gestational trophoblastic neoplasia: platinum/etoposide-containing regimen*, unless disease is methotrexate-resistant and high-risk (note some IV chemo may not require prior authorization)
OnabotulinumtoxinA (Botox [®])	<p>PART B STEP:</p> <ul style="list-style-type: none"> • Upper limb spasticity, cervical dystonia: Xeomin* and Dysport* • Blepharospasm: Xeomin* • Lower limb spasticity: Dysport*
Pegaptanib (Macugen [®])	<p>PART B STEP:</p> <ul style="list-style-type: none"> • Neovascular (wet) AMD: intravitreal bevacizumab solution

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Pegfilgrastim (Neulasta [®] , Fulphila [™] , Nyvepria [™] , Udenyca [™] , Ziextenzo [™])	PART B STEP: <ul style="list-style-type: none"> • All indications: Zarxio*, unless member requires \geq 10 doses of Zarxio, member is unable to self-administer Zarxio due to lack of caregiver or support system for assistance with administration and inadequate access to healthcare facility or home care interventions <ul style="list-style-type: none"> ○ If unable to use Zarxio for any of the reasons listed above and request is for an agent other than Ziextenzo: Ziextenzo* <ul style="list-style-type: none"> ▪ If unable to use Ziextenzo and request is for Neulasta: biosimilar pegfilgrastim product (e.g., Fulphila, Nyvepria, Udenyca)*
Pembrolizumab (Keytruda [®])	PART B STEP: <ul style="list-style-type: none"> • Head and neck squamous cell carcinoma: platinum-containing chemotherapy*, unless prescribed as part of combination therapy or prescribed as a single agent for a tumor that expresses PD-L1 with a combined positive score (CPS) \geq 1 • Classical Hodgkin lymphoma, primary mediastinal large B-cell lymphoma, esophageal squamous cell carcinoma, endometrial carcinoma, anal carcinoma, gestational trophoblastic neoplasia, extranodal NK/T-cell lymphoma, vulvar carcinoma, anaplastic large cell lymphoma, small cell lung cancer: at least 1 prior therapy* • Tumor mutational burden-high cancer: at least 1 prior therapy*, unless member has ampullary adenocarcinoma or pancreatic adenocarcinoma • Cervical cancer: at least 1 prior therapy*, unless prescribed in combination with chemotherapy (e.g., paclitaxel/cisplatin, paclitaxel/carboplatin) • Microsatellite instability-high/mismatch repair deficient cancer: at least 1 prior therapy*, unless member has colorectal cancer, ampullary adenocarcinoma, gallbladder cancer, intrahepatic/extrahepatic cholangiocarcinoma, non-nasopharyngeal head and neck cancer, occult primary tumor, pancreatic adenocarcinoma, or small bowel adenocarcinoma • Urothelial carcinoma: platinum-containing chemotherapy*, unless ineligible • Non-muscle invasive bladder cancer: Bacillus Calmette-Guerin*

Drug Name	Part B Required Step-Through Agents* By Indication <i>*May require prior authorization</i>
	<ul style="list-style-type: none"> • Gastric, esophagogastric junction, or esophageal adenocarcinoma: at least 1 prior therapy (if CPS \geq 10) or 2 prior therapies*, unless prescribed in combination with fluoropyrimidine- and platinum-containing chemotherapy <i>(note some IV chemo may not require prior authorization)</i>
Polatuzumab vedotin-piiq (Polivy [™])	PART B STEP: <ul style="list-style-type: none"> • Diffuse large B-cell lymphoma, high-grade B-cell lymphoma, mantle cell lymphoma, monomorphic post-transplant lymphoproliferative disorder (B-cell type), AIDS-related B-cell lymphoma: 2 prior therapies* • Follicular lymphoma: 1 prior therapy* <i>(note some IV chemo may not require prior authorization)</i>
Ramucirumab (Cyramza [®])	PART B STEP: <ul style="list-style-type: none"> • Esophageal, esophagogastric junction, and gastric cancer: prior lines of systemic therapy* <i>(note some IV chemo may not require prior authorization)</i>
Ranibizumab (Lucentis [®])	PART B STEP: <ul style="list-style-type: none"> • Neovascular (wet) AMD, macular edema following RVO, DME, DR, or myopic choroidal neovascularization (mCNV): intravitreal bevacizumab solution
Rituximab (Rituxan [®] , Riabni [™] , Ruxience [™] , Truxima [®]), rituximab/hyaluronidase (Rituxan Hycela [™])	PART B STEP: <ul style="list-style-type: none"> • All indications, if request is for Rituxan: Ruxience, Truxima, and Riabni[†] • All indications, if request is for Riabni: Ruxience* and Truxima* • All oncology indications, if request is for Rituxan Hycela: member has received at least one full dose of Rituxan, Riabni, Ruxience, or Truxima IN ADDITION: <ul style="list-style-type: none"> • Rheumatoid arthritis, if request is for Rituxan or Riabni: infliximab*
Romiplostim (Nplate [®])	PART B STEP: <ul style="list-style-type: none"> • Immune thrombocytopenia: systemic corticosteroid (if intolerant or contraindicated to systemic corticosteroids, then immune globulin*) • Myelodysplastic syndrome: hypomethylating agent (e.g., azacitadine*, decitabine*) or immunosuppressive therapy (e.g., Atgam*)

Drug Name	Part B Required Step-Through Agents* By Indication <i>*May require prior authorization</i>
	<ul style="list-style-type: none"> • Chemotherapy-induced thrombocytopenia: prior chemotherapy* (<i>note some IV chemo may not require prior authorization</i>)
Ramosuzumab-aqqg (Evenity™)	PART B STEP: <ul style="list-style-type: none"> • Postmenopausal osteoporosis: bisphosphonate, unless member is very high risk for fracture (recent osteoporotic fracture within the past 12 months, BMD T-score at hip or spine ≤ -3.0, OR BMD T-score at hip or spine ≤ -2.5 and major osteoporotic fracture [i.e., hip, spine, forearm, wrist, humerus])
Sargramostim (Leukine®)	PART B STEP: <ul style="list-style-type: none"> • All indications: Zarxio
Sipuleucel-T (Provenge®)	PART B STEP: <ul style="list-style-type: none"> • Prostate cancer: androgen deprivation therapy (e.g., Zoladex*, Vantas*, leuprolide*, Trelstar*, Firmagon*)
Teprotumumab-trbw (Tepezza™)	PART B STEP: <ul style="list-style-type: none"> • Thyroid eye disease: a systemic corticosteroid
Tisagenlecleucel (Kymriah®)	PART B STEP: <ul style="list-style-type: none"> • B-cell precursor acute lymphoblastic leukemia: at least two prior systemic therapies* <i>Only for initial treatment dose; subsequent doses will not be covered</i> • Large B-cell lymphoma: 2 lines of systemic therapy that includes rituximab* and one anthracycline-containing regimen (e.g., doxorubicin*) <i>Only for initial treatment dose; subsequent doses will not be covered</i> • Relapsed or refractory follicular lymphoma: 2 lines of systemic therapy that includes a combination of an anti-CD20 monoclonal antibody (e.g., rituximab or Gazyva)* and an alkylating agent (e.g., bendamustine, cyclophosphamide, chlorambucil) <i>Only for initial treatment dose; subsequent doses will not be covered</i>
Tocilizumab (Actemra®)	PART B STEP: <ul style="list-style-type: none"> • Polyarticular juvenile idiopathic arthritis, systemic juvenile idiopathic arthritis, and rheumatoid arthritis: a TNF inhibitor (e.g., infliximab)* (<i>note credit may be given if another TNF inhibitor was tried</i>)
Trastuzumab (Herceptin®), Ontruzant®, Herzuma®, Ogivri™, Trazimera™	PART B STEP:

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Kanjinti™), trastuzumab/hyaluronidase (Herceptin Hylecta™)	<ul style="list-style-type: none"> • All indications, if request is for an agent other than Trazimera: Trazimera* <ul style="list-style-type: none"> ○ If unable to use Trazimera and request is for Herceptin or Herceptin Hylecta: biosimilar trastuzumab product (e.g., Ogivri, Kanjinti)
Triamcinolone ER injection (Zilretta®)	PART B STEP: <ul style="list-style-type: none"> • Osteoarthritis of the knee: intra-articular glucocorticoid injection
Triamcinolone acetonide suprachoroidal injection (Xipere™)	PART B STEP: <ul style="list-style-type: none"> • All indications: Triesence (triamcinolone) intravitreal injection
Ustekinumab (Stelara®)	PART B STEP: <ul style="list-style-type: none"> • All indications: a TNF inhibitor (e.g., infliximab)* <i>(note credit may be given if another TNF inhibitor was tried)</i>
Vedolizumab (Entyvio®)	PART B STEP: <ul style="list-style-type: none"> • All indications: a TNF inhibitor (e.g., infliximab)* <i>(note credit may be given if another TNF inhibitor was tried)</i>
Verteporfin (Visudyne®)	PART B STEP: <ul style="list-style-type: none"> • Classic subfoveal CNV due to AMD, pathologic myopia, or presumed ocular histoplasmosis: intravitreal bevacizumab solution

For questions, please reach out to your provider relations.

Important Reminder

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on FDA recommendation(s), peer-reviewed medical literature and evidence-based clinical practice guidelines.

The purpose of this clinical policy is to provide a guide to medical necessity, which is a component of the guidelines used to assist in making coverage decisions and administering benefits. It does not constitute a contract or guarantee regarding payment or results. Coverage decisions and the administration of benefits are subject to all terms, conditions, exclusions and limitations of the coverage documents (e.g., evidence of coverage, certificate of coverage, policy, contract of insurance, etc.), as well as to state and federal requirements and applicable Health Plan-level administrative policies and procedures.

This clinical policy is effective as of the date determined by the Health Plan or responsible business unit. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall

govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

This clinical policy does not constitute medical advice, medical treatment or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members. This clinical policy is not intended to recommend treatment for members. Members should consult with their treating physician in connection with diagnosis and treatment decisions.

Providers referred to in this clinical policy are independent contractors who exercise independent judgment and over whom the Health Plan has no control or right of control. Providers are not agents or employees of the Health Plan.

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