STEP THERAPY CRITERIA FOR MEDICARE PART B DRUGS

This list is current as of 12/01/2022 and pertains to the following Independent Health Medicare Advantage Plans for 2022:

Independent Health's Encompass 65® Basic (HMO)
Independent Health's Encompass 65® Core (HMO)
Independent Health's Encompass 65® Edge (HMO)
Independent Health's Encompass 65® Element (HMO)
Independent Health's Medicare Passport® Advantage (PPO)
Independent Health's Medicare Passport® Prime (PPO)
Independent Health's Medicare Family Choice® (HMO I-SNP)
Independent Health's Assure Advantage® (HMO C-SNP)

In some cases, we require that you first try certain drugs to treat your medical condition before we will cover another drug for that condition. For example, if Drug X and Drug Y both treat your medical condition, we may not cover Drug Y unless you try Drug X first. If Drug X does not work for you, we will then cover Drug Y.

The therapy outlined in this document may also involve a combination of Part B and Part D drugs. For example, we may not cover a Part B drug unless you try a Part D drug first. Or, we may not cover a Part D drug unless you try a Part B drug first. This is dependent on the therapy described to treat your medical condition. This document contains the Step Therapy protocols for Medicare Part B drugs that are associated with the Independent Health Medicare Advantage Plans mentioned above.

If you have any questions, please contact our Medicare Member Services Department at 1-800-665-1502 or, for TTY users 711, October 1st – March 31st: Monday through Sunday from 8 a.m. to 8 p.m., April 1st – September 30th: Monday through Friday from 8 a.m. to 8 p.m.

The formulary may change at any time. You will receive notice when necessary.

Apretude

Products Affected

• APRETUDE SUSPENSION EXTENDED RELEASE 600 MG/3ML INTRAMUSCULAR

| Criteria | For approval, patient must have tried and had an intolerance to or has a |
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| | contraindication to emtricitabine/tenofovir disoproxil fumarate. |

Asceniv

Products Affected

 ASCENIV SOLUTION 5 GM/50ML INTRAVENOUS

| Criteria | For approval, patient must have tried and failed to have a response to |
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| | another intravenous immunoglobulin (IVIG) product. |

Bendamustine

Products Affected

- TREANDA SOLUTION RECONSTITUTED 100 MG INTRAVENOUS
- TREANDA SOLUTION RECONSTITUTED 25 MG INTRAVENOUS

| Criteria | For approval of Treanda, the patient must have tried and failed to have an adequate response to Belrapzo or Bendeka. This specific requirement applies to new starts only. |
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Beovu

Products Affected

- BEOVU SOLUTION 6 MG/0.05ML INTRAVITREAL
- BEOVU SOLUTION PREFILLED SYRINGE 6 MG/0.05ML INTRAVITREAL

| | For approval for a diagnosis of neovascular (wet) age-related macular degeneration or diabetic macular edema, the patient must have tried and failed to have an adequate response to bevacizumab (Avastin). This specific requirement applies to new starts only. |
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Bevacizumab

Products Affected

- ALYMSYS SOLUTION 100 MG/4ML INTRAVENOUS
- ALYMSYS SOLUTION 400 MG/16ML INTRAVENOUS
- AVASTIN SOLUTION 100 MG/4ML INTRAVENOUS
- AVASTIN SOLUTION 400 MG/16ML INTRAVENOUS
- MVASI SOLUTION 100 MG/4ML INTRAVENOUS
- MVASI SOLUTION 400 MG/16ML INTRAVENOUS

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| Criteria | For approval of Alymsys, Avastin, or Mvasi for oncology (cancer) indications, the patient must have tried and failed to have an adequate response to Zirabev. This specific requirement applies to new starts only. This requirement does not apply to patients using Avastin for ophthalmic (eye) indications or for any indication not shared by Zirabev and Avastin or another bevacizumab biosimilar agent. |

Botulinum toxins

Products Affected

- DYSPORT SOLUTION RECONSTITUTED 300 UNIT INTRAMUSCULAR
- DYSPORT SOLUTION RECONSTITUTED 500 UNIT INTRAMUSCULAR
- MYOBLOC SOLUTION 10000 UNIT/2ML

INTRAMUSCULAR

- MYOBLOC SOLUTION 2500 UNIT/0.5ML INTRAMUSCULAR
- MYOBLOC SOLUTION 5000 UNIT/ML INTRAMUSCULAR

| | For approval, the patient must have tried and failed to have an adequate response to Botox and Xeomin. This specific requirement applies to new starts only. |
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Enjaymo

Products Affected

• ENJAYMO SOLUTION 1100 MG/22ML INTRAVENOUS

| For approval, the patient must have tried and failed to have an adequate response to Ruxience (or another rituximab product) or any other B-cell targeting therapy. This specific requirement applies to new starts only. |
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Erythropoietins

Products Affected

- EPOGEN SOLUTION 10000 UNIT/ML INJECTION
- EPOGEN SOLUTION 2000 UNIT/ML INJECTION
- EPOGEN SOLUTION 20000 UNIT/ML INJECTION
- EPOGEN SOLUTION 3000 UNIT/ML INJECTION
- EPOGEN SOLUTION 4000 UNIT/ML INJECTION
- PROCRIT SOLUTION 10000 UNIT/ML

INJECTION

- PROCRIT SOLUTION 2000 UNIT/ML INJECTION
- PROCRIT SOLUTION 20000 UNIT/ML INJECTION
- PROCRIT SOLUTION 3000 UNIT/ML INJECTION
- PROCRIT SOLUTION 4000 UNIT/ML INJECTION
- PROCRIT SOLUTION 40000 UNIT/ML INJECTION

| Criteria | For approval, the patient must have tried and failed to have an adequate |
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| | response to Retacrit. This specific requirement applies to new starts only. |

Evkeeza

Products Affected

- EVKEEZA SOLUTION 1200 MG/8ML INTRAVENOUS
- EVKEEZA SOLUTION 345 MG/2.3ML INTRAVENOUS

| Criteria | For approval of Evkeeza, the patient must have tried and failed to have an adequate response to or have a contraindication to both a maximally-tolerated dose of a statin drug and a PCSK9 inhibitor. This specific requirement applies to new starts only. |
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| | requirement applies to new starts only. |

Eylea

Products Affected

- EYLEA SOLUTION 2 MG/0.05ML INTRAVITREAL
- EYLEA SOLUTION PREFILLED SYRINGE 2 MG/0.05ML INTRAVITREAL

| Criteria | For approval for a diagnosis of neovascular (wet) age-related macular degeneration, diabetic macular edema, or diabetic retinopathy in patients with diabetic macular edema, the patient must have tried and failed to have an adequate response to bevacizumab (Avastin). This specific requirement applies to new starts only. |
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Feiba

Products Affected

- FEIBA SOLUTION RECONSTITUTED 1000 UNIT INTRAVENOUS
- FEIBA SOLUTION RECONSTITUTED 2500

UNIT INTRAVENOUS

 FEIBA SOLUTION RECONSTITUTED 500 UNIT INTRAVENOUS

| Criteria | For approval of Feiba, the patient must have tried and failed to have an adequate response to Hemlibra. This specific requirement applies to new starts only. This requirement does not apply to treatment of hemophilia B. |
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Filgrastim

Products Affected

- GRANIX SOLUTION 300 MCG/ML SUBCUTANEOUS
- GRANIX SOLUTION 480 MCG/1.6ML SUBCUTANEOUS
- GRANIX SOLUTION PREFILLED SYRINGE 300 MCG/0.5ML SUBCUTANEOUS
- GRANIX SOLUTION PREFILLED SYRINGE 480 MCG/0.8ML SUBCUTANEOUS
- NEUPOGEN SOLUTION 300 MCG/ML INJECTION
- NEUPOGEN SOLUTION 480 MCG/1.6ML INJECTION
- NEUPOGEN SOLUTION PREFILLED SYRINGE 300 MCG/0.5ML INJECTION
- NEUPOGEN SOLUTION PREFILLED SYRINGE 480 MCG/0.8ML INJECTION

- NIVESTYM SOLUTION 300 MCG/ML INJECTION
- NIVESTYM SOLUTION 480 MCG/1.6ML INJECTION
- NIVESTYM SOLUTION PREFILLED SYRINGE 300 MCG/0.5ML INJECTION
- NIVESTYM SOLUTION PREFILLED SYRINGE 480 MCG/0.8ML INJECTION
- RELEUKO SOLUTION 300 MCG/ML INJECTION
- releuko solution 480 mcg/1.6ml injection
- releuko solution prefilled syringe 300 mcg/0.5ml subcutaneous
- releuko solution prefilled syringe 480 mcg/0.8ml subcutaneous

| Criteria | For approval of Neupogen for any indication other than Hematopoietic Syndrome of Acute Radiation Syndrome, Granix, Nivestym, or Releuko, the patient must have tried and failed to have an adequate response to |
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| | Zarxio. This specific requirement applies to new starts only. |

Growth hormone

Products Affected

- GENOTROPIN CARTRIDGE 12 MG SUBCUTANEOUS
- GENOTROPIN CARTRIDGE 5 MG SUBCUTANEOUS
- GENOTROPIN MINIQUICK PREFILLED SYRINGE 0.2 MG SUBCUTANEOUS
- GENOTROPIN MINIQUICK PREFILLED SYRINGE 0.4 MG SUBCUTANEOUS
- GENOTROPIN MINIQUICK PREFILLED SYRINGE 0.6 MG SUBCUTANEOUS
- GENOTROPIN MINIQUICK PREFILLED SYRINGE 0.8 MG SUBCUTANEOUS
- GENOTROPIN MINIQUICK PREFILLED SYRINGE 1 MG SUBCUTANEOUS
- GENOTROPIN MINIQUICK PREFILLED SYRINGE 1.2 MG SUBCUTANEOUS
- GENOTROPIN MINIQUICK PREFILLED SYRINGE 1.4 MG SUBCUTANEOUS
- GENOTROPIN MINIQUICK PREFILLED SYRINGE 1.6 MG SUBCUTANEOUS
- GENOTROPIN MINIQUICK PREFILLED SYRINGE 1.8 MG SUBCUTANEOUS
- GENOTROPIN MINIQUICK PREFILLED SYRINGE 2 MG SUBCUTANEOUS
- HUMATROPE CARTRIDGE 12 MG INJECTION
- HUMATROPE CARTRIDGE 24 MG INJECTION
- HUMATROPE CARTRIDGE 6 MG INJECTION

- NORDITROPIN FLEXPRO SOLUTION PEN-INJECTOR 10 MG/1.5ML SUBCUTANEOUS
- NORDITROPIN FLEXPRO SOLUTION PEN-INJECTOR 15 MG/1.5ML SUBCUTANEOUS
- NORDITROPIN FLEXPRO SOLUTION PEN-INJECTOR 30 MG/3ML SUBCUTANEOUS
- NORDITROPIN FLEXPRO SOLUTION PEN-INJECTOR 5 MG/1.5ML SUBCUTANEOUS
- NUTROPIN AQ NUSPIN 10 SOLUTION PEN-INJECTOR 10 MG/2ML SUBCUTANEOUS
- NUTROPIN AQ NUSPIN 20 SOLUTION PEN-INJECTOR 20 MG/2ML SUBCUTANEOUS
- NUTROPIN AQ NUSPIN 5 SOLUTION PEN-INJECTOR 5 MG/2ML SUBCUTANEOUS
- OMNITROPE SOLUTION CARTRIDGE 10 MG/1.5ML SUBCUTANEOUS
- OMNITROPE SOLUTION CARTRIDGE 5 MG/1.5ML SUBCUTANEOUS
- OMNITROPE SOLUTION RECONSTITUTED 5.8 MG SUBCUTANEOUS
- SAIZEN SOLUTION RECONSTITUTED 5 MG INJECTION
- SAIZEN SOLUTION RECONSTITUTED 8.8 MG
 INJECTION
- SAIZENPREP SOLUTION RECONSTITUTED 8.8 MG INJECTION

| Criteria | For Commercial and Essential plans, the patient must have tried and failed to have an adequate response to Genotropin. |
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| | For Medicaid plans, the patient must have tried and failed to have an adequate response to Norditropin. |

Ilumya

Products Affected

• ILUMYA SOLUTION PREFILLED SYRINGE 100 MG/ML SUBCUTANEOUS

| Criteria | For approval, the patient must have tried and failed to have an adequate response to or had an intolerance to at least the formulary-preferred IL-17 inhibitor (Cosentyx) and one other on-formulary biologic agent for the treatment of psoriasis (Cimzia, Enbrel, Humira, Otezla, Skyrizi, Stelara, Tremfya). |
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Infliximab

Products Affected

- AVSOLA SOLUTION RECONSTITUTED 100 MG REMICADE SOLUTION RECONSTITUTED 100 **INTRAVENOUS**
- infliximab solution reconstituted 100 mg intravenous
- MG INTRAVENOUS
- RENFLEXIS SOLUTION RECONSTITUTED 100 MG INTRAVENOUS

| Criteria | For approval of Avsola, generic infliximab, Remicade, or Renflexis, the patient must have tried and failed to have an adequate response to Inflectra. This specific requirement applies to new starts only. This requirement does not apply to patients using Avsola, generic infliximab, Remicade, Renflexis, or another infliximab biosimilar agent for any indication not shared with Inflectra. |
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Leucovorins

Products Affected

- FUSILEV SOLUTION RECONSTITUTED 50 MG INTRAVENOUS
- KHAPZORY SOLUTION RECONSTITUTED 175 MG INTRAVENOUS
- KHAPZORY SOLUTION RECONSTITUTED 300 MG INTRAVENOUS
- levoleucovorin calcium pf solution 175 mg/17.5ml intravenous
- levoleucovorin calcium pf solution 250 mg/25ml intravenous
- levoleucovorin calcium solution reconstituted 50 mg intravenous

| Criteria | For approval, the patient must have tried and failed to have an adequate response to generic leucovorin. This specific requirement applies to new starts only. |
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Long-acting hemophilia factors

Products Affected

- adynovate solution reconstituted 1000 unit intravenous
- adynovate solution reconstituted 1500 unit intravenous
- adynovate solution reconstituted 2000 unit intravenous
- adynovate solution reconstituted 250 unit intravenous
- adynovate solution reconstituted 3000 unit intravenous
- adynovate solution reconstituted 500 unit intravenous
- adynovate solution reconstituted 750 unit intravenous
- ELOCTATE SOLUTION RECONSTITUTED 1000 UNIT INTRAVENOUS
- ELOCTATE SOLUTION RECONSTITUTED 2000 UNIT INTRAVENOUS

- ELOCTATE SOLUTION RECONSTITUTED 250 UNIT INTRAVENOUS
- ELOCTATE SOLUTION RECONSTITUTED 4000 UNIT INTRAVENOUS
- ELOCTATE SOLUTION RECONSTITUTED 5000 UNIT INTRAVENOUS
- ELOCTATE SOLUTION RECONSTITUTED 6000 UNIT INTRAVENOUS
- ESPEROCT SOLUTION RECONSTITUTED 1000 UNIT INTRAVENOUS
- ESPEROCT SOLUTION RECONSTITUTED 1500 UNIT INTRAVENOUS
- ESPEROCT SOLUTION RECONSTITUTED 2000 UNIT INTRAVENOUS
- ESPEROCT SOLUTION RECONSTITUTED 3000 UNIT INTRAVENOUS
- ESPEROCT SOLUTION RECONSTITUTED 500 UNIT INTRAVENOUS

| tr | For approval of Adynovate, Eloctate, or Esperoct, the patient must have ried and failed to have an adequate response to Jivi. This specific requirement applies to new starts only. |
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Opdualag

Products Affected

 OPDUALAG SOLUTION 240-80 MG/20ML INTRAVENOUS

| Yervoy. This specific requirement applies to new starts only. | resp | opproval, the patient must have tried and failed to have an adequate onse to Opdivo plus Yervoy or has a contraindication to the use of oy. This specific requirement applies to new starts only. |
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Paclitaxel

Products Affected

- ABRAXANE SUSPENSION RECONSTITUTED 100 MG INTRAVENOUS
- paclitaxel protein-bound part suspension reconstituted 100 mg intravenous

| | For approval of Abraxane for any indication other than pancreatic cancer or small bowel carcinoma, the patient must have tried and failed to have an adequate response to generic paclitaxel. This specific requirement applies to new starts only. |
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Pegfilgrastim

Products Affected

- FULPHILA SOLUTION PREFILLED SYRINGE 6 MG/0.6ML SUBCUTANEOUS
- NYVEPRIA SOLUTION PREFILLED SYRINGE 6
- MG/0.6ML SUBCUTANEOUS
- ZIEXTENZO SOLUTION PREFILLED SYRINGE 6 MG/0.6ML SUBCUTANEOUS

| Criteria For approval of Fulphila, Nyvepria, or Ziextenzo, the patient must have tried and failed to have an adequate response to both Udenyca and Neulasta. This specific requirement applies to new starts only. |
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Ranibizumab

Products Affected

- BYOOVIZ SOLUTION 0.5 MG/0.05ML INTRAVITREAL
- CIMERLI SOLUTION 0.3 MG/0.05ML INTRAVITREAL
- CIMERLI SOLUTION 0.5 MG/0.05ML INTRAVITREAL
- LUCENTIS SOLUTION 0.3 MG/0.05ML INTRAVITREAL
- LUCENTIS SOLUTION PREFILLED SYRINGE 0.3 MG/0.05ML INTRAVITREAL
- LUCENTIS SOLUTION PREFILLED SYRINGE 0.5 MG/0.05ML INTRAVITREAL
- SUSVIMO (IMPLANT 1ST FILL) SOLUTION 10 MG/0.1ML INTRAVITREAL
- SUSVIMO (IMPLANT REFILL) SOLUTION 10 MG/0.1ML INTRAVITREAL

Details

Criteria For approval of Cimerli for shared indications, the patient must have tried and failed to have an adequate response to both (Step 1) bevacizumab (Avastin) and (Step 2) either aflibercept (Eylea), brolucizumab (Beovu), or faricimab (Vabysmo). For approval of Byooviz, Lucentis, or Susvimo for shared indications, the patient must have tried and failed to have an adequate response to Steps 1 and 2 and Cimerli. This specific requirement applies to new starts only.

Rituximab

Products Affected

- RIABNI SOLUTION 100 MG/10ML INTRAVENOUS
- RIABNI SOLUTION 500 MG/50ML INTRAVENOUS
- RITUXAN HYCELA SOLUTION 1400-23400 MG -UT/11.7ML SUBCUTANEOUS
- RITUXAN HYCELA SOLUTION 1600-26800 MG -UT/13.4ML SUBCUTANEOUS
- RITUXAN SOLUTION 100 MG/10ML INTRAVENOUS
- RITUXAN SOLUTION 500 MG/50ML INTRAVENOUS
- TRUXIMA SOLUTION 100 MG/10ML INTRAVENOUS
- TRUXIMA SOLUTION 500 MG/50ML INTRAVENOUS

Details

Criteria For approval of Rituxan for all indications except pemphigus vulgaris (PV), Riabni, Rituxan Hycela, or Truxima, the patient must have tried and failed to have an adequate response to Ruxience. This specific requirement applies to new starts only. This requirement does not apply to patients using Riabni, Rituxan, Rituxan Hycela, or Truxima for any indication not shared by Ruxience or another rituximab biosimilar agent.

Saphnelo

Products Affected

• SAPHNELO SOLUTION 300 MG/2ML INTRAVENOUS

| For approval, the patient must have tried and failed to have an adequate |
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| response to or had an intolerance to Benlysta. |

Soliris

Products Affected

• SOLIRIS SOLUTION 300 MG/30ML INTRAVENOUS

| Criteria | For a diagnosis of neuromyelitis optica spectrum disorder (NMOSD), the patient must have tried and failed to have an adequate response to or had an intolerance to or contraindication to Enspryng and either Uplizna or rituximab. For a diagnosis of generalized myasthenia gravis, the |
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| | patient must have tried and failed to have an adequate response to or had an intolerance to or contraindication to Vyvgart. These requirements apply to new starts only. |

Spinraza

Products Affected

• SPINRAZA SOLUTION 12 MG/5ML INTRATHECAL

| response to or had an intolerance to or contraindication to Evrysdi. This specific requirement applies to new starts only. |
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Tezspire

Products Affected

• TEZSPIRE SOLUTION PREFILLED SYRINGE 210 MG/1.91ML SUBCUTANEOUS

| Criteria | For approval, the patient must have (1) tried and failed, was intolerant to, or had a contraindication to dupilumab plus one other biologic for severe asthma, including either an IL-5 antagonist, an IL-5 receptor antagonist, or omalizumab or (2) an eosinophil count below that required to use these other medications. This specific requirement applies to new starts only. |
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Trastuzumab

Products Affected

- HERCEPTIN HYLECTA SOLUTION 600-10000 MG-UNT/5ML SUBCUTANEOUS
- HERCEPTIN SOLUTION RECONSTITUTED 150 MG INTRAVENOUS
- HERZUMA SOLUTION RECONSTITUTED 150 MG INTRAVENOUS
- HERZUMA SOLUTION RECONSTITUTED 420 MG INTRAVENOUS
- KANJINTI SOLUTION RECONSTITUTED 150 MG INTRAVENOUS

- KANJINTI SOLUTION RECONSTITUTED 420 MG INTRAVENOUS
- OGIVRI SOLUTION RECONSTITUTED 150 MG INTRAVENOUS
- OGIVRI SOLUTION RECONSTITUTED 420 MG INTRAVENOUS
- ONTRUZANT SOLUTION RECONSTITUTED 150 MG INTRAVENOUS
- ONTRUZANT SOLUTION RECONSTITUTED 420 MG INTRAVENOUS

| Criteria | For approval of Herceptin, Herceptin Hylecta, Herzuma, Kanjinti, Ogivri, |
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| | or Ontruzant, the patient must have tried and failed to have an adequate response to Trazimera. This specific requirement applies to new starts only. This requirement does not apply to patients using a trastuzumab biosimilar agent for any indication not shared by Trazimera. |

Ultomiris

Products Affected

- ULTOMIRIS SOLUTION 1100 MG/11ML INTRAVENOUS
- ULTOMIRIS SOLUTION 300 MG/3ML INTRAVENOUS

| Criteria | For a diagnosis of generalized myasthenia gravis, the patient must have tried and failed to have an adequate response to or had an intolerance to or contraindication to Vyvgart. These requirements apply to new starts only. |
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Uplizna

Products Affected

• UPLIZNA SOLUTION 100 MG/10ML INTRAVENOUS

| For approval, the patient must have tried and failed to have an adequate response to or had an intolerance to or contraindication to Enspryng. This specific requirement applies to new starts only. |
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Vabysmo

Products Affected

 VABYSMO SOLUTION 6 MG/0.05ML INTRAVITREAL

| Criteria | For approval for a diagnosis of neovascular (wet) age-related macular degeneration or diabetic macular edema, the patient must have tried and failed to have an adequate response to bevacizumab (Avastin). This specific requirement applies to new starts only. |
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Viscosupplements

Products Affected

- DUROLANE PREFILLED SYRINGE 60 MG/3ML INTRA-ARTICULAR
- GEL-ONE PREFILLED SYRINGE 30 MG/3ML INTRA-ARTICULAR
- GELSYN-3 SOLUTION PREFILLED SYRINGE 16.8 MG/2ML INTRA-ARTICULAR
- GENVISC 850 SOLUTION PREFILLED SYRINGE 25 MG/2.5ML INTRA-ARTICULAR
- HYALGAN SOLUTION 20 MG/2ML INTRA-ARTICULAR
- HYALGAN SOLUTION PREFILLED SYRINGE 20 MG/2ML INTRA-ARTICULAR
- HYMOVIS SOLUTION PREFILLED SYRINGE 24 MG/3ML INTRA-ARTICULAR
- MONOVISC SOLUTION PREFILLED SYRINGE

- 88 MG/4ML INTRA-ARTICULAR
- ORTHOVISC SOLUTION PREFILLED SYRINGE 30 MG/2ML INTRA-ARTICULAR
- SUPARTZ FX SOLUTION PREFILLED SYRINGE 25 MG/2.5ML INTRA-ARTICULAR
- SYNOJOYNT SOLUTION PREFILLED SYRINGE 20 MG/2ML INTRA-ARTICULAR
- SYNVISC SOLUTION PREFILLED SYRINGE 16 MG/2ML INTRA-ARTICULAR
- TRILURON SOLUTION PREFILLED SYRINGE 20 MG/2ML INTRA-ARTICULAR
- TRIVISC SOLUTION PREFILLED SYRINGE 25 MG/2.5ML INTRA-ARTICULAR
- VISCO-3 SOLUTION PREFILLED SYRINGE 25 MG/2.5ML INTRA-ARTICULAR

| Criteria | For approval of viscosupplements other than Euflexxa or Synvisc-One, the patient must have tried and failed to have an adequate response to or had an intolerance to both preferred agents (Euflexxa, Synvisc-One). All viscosupplements require medical prior authorization. Because all viscosupplements are considered medical devices and not drugs by the FDA, they can only be billed through Medicare Part B. |
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| | Viscosupplements are not covered on Commercial and State health plans. |

Vyepti

Products Affected

• VYEPTI SOLUTION 100 MG/ML INTRAVENOUS

| Criteria | For approval of Vyepti, the patient must have tried and failed to have an adequate response to any two oral agents used to prevent/reduce frequency of migraines and have had prior use of one injectable CGRP antagonist/receptor antagonist. This specific requirement applies to new starts only. |
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Vyvgart

Products Affected

 VYVGART SOLUTION 400 MG/20ML INTRAVENOUS

| Criteria | For approval, the patient must have failed to respond to therapy with at least two of the following drug groups: acetylcholinesterase inhibitors, corticosteroids, nonsteroidal immunosuppressive therapies (e.g., methotrexate, azathioprine, mycophenolate, cyclosporine). This specific requirement applies to new starts only. |
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Zilretta

Products Affected

• ZILRETTA SUSPENSION RECONSTITUTED ER 32 MG INTRA-ARTICULAR

| | For approval, the patient must have tried and failed to have an adequate response to at least one other injectable corticosteroid. This specific requirement applies to new starts only. |
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INDEX

| ABRAXANE SUSPENSION RECONSTITUTED | DYSPORT SOLUTION RECONSTITUTED 300 |
|--|-------------------------------------|
| 100 MG INTRAVENOUS19 | UNIT INTRAMUSCULAR6 |
| adynovate solution reconstituted 1000 unit | DYSPORT SOLUTION RECONSTITUTED 500 |
| intravenous17 | UNIT INTRAMUSCULAR6 |
| adynovate solution reconstituted 1500 unit | ELOCTATE SOLUTION RECONSTITUTED |
| intravenous17 | 1000 UNIT INTRAVENOUS17 |
| adynovate solution reconstituted 2000 unit | ELOCTATE SOLUTION RECONSTITUTED |
| intravenous17 | 2000 UNIT INTRAVENOUS17 |
| adynovate solution reconstituted 250 unit | ELOCTATE SOLUTION RECONSTITUTED 250 |
| intravenous17 | UNIT INTRAVENOUS17 |
| adynovate solution reconstituted 3000 unit | ELOCTATE SOLUTION RECONSTITUTED |
| intravenous17 | 4000 UNIT INTRAVENOUS17 |
| adynovate solution reconstituted 500 unit | ELOCTATE SOLUTION RECONSTITUTED |
| intravenous17 | 5000 UNIT INTRAVENOUS17 |
| adynovate solution reconstituted 750 unit | ELOCTATE SOLUTION RECONSTITUTED |
| intravenous17 | 6000 UNIT INTRAVENOUS17 |
| ALYMSYS SOLUTION 100 MG/4ML | ENJAYMO SOLUTION 1100 MG/22ML |
| INTRAVENOUS5 | INTRAVENOUS7 |
| ALYMSYS SOLUTION 400 MG/16ML | EPOGEN SOLUTION 10000 UNIT/ML |
| INTRAVENOUS5 | INJECTION8 |
| APRETUDE SUSPENSION EXTENDED | EPOGEN SOLUTION 2000 UNIT/ML |
| RELEASE 600 MG/3ML INTRAMUSCULAR 1 | INJECTION8 |
| ASCENIV SOLUTION 5 GM/50ML | EPOGEN SOLUTION 20000 UNIT/ML |
| INTRAVENOUS2 | INJECTION8 |
| AVASTIN SOLUTION 100 MG/4ML | EPOGEN SOLUTION 3000 UNIT/ML |
| INTRAVENOUS5 | INJECTION8 |
| AVASTIN SOLUTION 400 MG/16ML | EPOGEN SOLUTION 4000 UNIT/ML |
| INTRAVENOUS5 | INJECTION8 |
| AVSOLA SOLUTION RECONSTITUTED 100 | ESPEROCT SOLUTION RECONSTITUTED |
| MG INTRAVENOUS15 | 1000 UNIT INTRAVENOUS17 |
| BEOVU SOLUTION 6 MG/0.05ML | ESPEROCT SOLUTION RECONSTITUTED |
| INTRAVITREAL 4 | 1500 UNIT INTRAVENOUS17 |
| BEOVU SOLUTION PREFILLED SYRINGE 6 | ESPEROCT SOLUTION RECONSTITUTED |
| MG/0.05ML INTRAVITREAL4 | 2000 UNIT INTRAVENOUS17 |
| BYOOVIZ SOLUTION 0.5 MG/0.05ML | ESPEROCT SOLUTION RECONSTITUTED |
| INTRAVITREAL21 | 3000 UNIT INTRAVENOUS17 |
| CIMERLI SOLUTION 0.3 MG/0.05ML | ESPEROCT SOLUTION RECONSTITUTED 500 |
| INTRAVITREAL21 | UNIT INTRAVENOUS17 |
| CIMERLI SOLUTION 0.5 MG/0.05ML | EVKEEZA SOLUTION 1200 MG/8ML |
| INTRAVITREAL21 | INTRAVENOUS9 |
| DUROLANE PREFILLED SYRINGE 60 | EVKEEZA SOLUTION 345 MG/2.3ML |
| MG/3ML INTRA-ARTICULAR31 | INTRAVENOUS9 |

| EYLEA SOLUTION 2 MG/0.05ML | GRANIX SOLUTION 300 MCG/ML |
|--|--|
| INTRAVITREAL10 | SUBCUTANEOUS12 |
| EYLEA SOLUTION PREFILLED SYRINGE 2 | GRANIX SOLUTION 480 MCG/1.6ML |
| MG/0.05ML INTRAVITREAL10 | SUBCUTANEOUS12 |
| FEIBA SOLUTION RECONSTITUTED 1000 | GRANIX SOLUTION PREFILLED SYRINGE |
| UNIT INTRAVENOUS11 | 300 MCG/0.5ML SUBCUTANEOUS12 |
| FEIBA SOLUTION RECONSTITUTED 2500 | GRANIX SOLUTION PREFILLED SYRINGE |
| UNIT INTRAVENOUS11 | 480 MCG/0.8ML SUBCUTANEOUS12 |
| FEIBA SOLUTION RECONSTITUTED 500 | HERCEPTIN HYLECTA SOLUTION 600-10000 |
| UNIT INTRAVENOUS11 | MG-UNT/5ML SUBCUTANEOUS27 |
| FULPHILA SOLUTION PREFILLED SYRINGE 6 | HERCEPTIN SOLUTION RECONSTITUTED |
| MG/0.6ML SUBCUTANEOUS20 | 150 MG INTRAVENOUS27 |
| FUSILEV SOLUTION RECONSTITUTED 50 | HERZUMA SOLUTION RECONSTITUTED 150 |
| MG INTRAVENOUS16 | MG INTRAVENOUS27 |
| GEL-ONE PREFILLED SYRINGE 30 MG/3ML | HERZUMA SOLUTION RECONSTITUTED 420 |
| INTRA-ARTICULAR31 | MG INTRAVENOUS27 |
| GELSYN-3 SOLUTION PREFILLED SYRINGE | HUMATROPE CARTRIDGE 12 MG |
| 16.8 MG/2ML INTRA-ARTICULAR31 | INJECTION13 |
| GENOTROPIN CARTRIDGE 12 MG | HUMATROPE CARTRIDGE 24 MG |
| SUBCUTANEOUS13 | INJECTION13 |
| GENOTROPIN CARTRIDGE 5 MG | HUMATROPE CARTRIDGE 6 MG INJECTION. 13 |
| SUBCUTANEOUS13 | HYALGAN SOLUTION 20 MG/2ML INTRA- |
| GENOTROPIN MINIQUICK PREFILLED | ARTICULAR31 |
| SYRINGE 0.2 MG SUBCUTANEOUS13 | HYALGAN SOLUTION PREFILLED SYRINGE |
| GENOTROPIN MINIQUICK PREFILLED | 20 MG/2ML INTRA-ARTICULAR31 |
| SYRINGE 0.4 MG SUBCUTANEOUS13 | HYMOVIS SOLUTION PREFILLED SYRINGE |
| GENOTROPIN MINIQUICK PREFILLED | 24 MG/3ML INTRA-ARTICULAR31 |
| SYRINGE 0.6 MG SUBCUTANEOUS13 | ILUMYA SOLUTION PREFILLED SYRINGE |
| GENOTROPIN MINIQUICK PREFILLED | 100 MG/ML SUBCUTANEOUS14 |
| SYRINGE 0.8 MG SUBCUTANEOUS13 | infliximab solution reconstituted 100 mg |
| GENOTROPIN MINIQUICK PREFILLED | intravenous15 |
| SYRINGE 1 MG SUBCUTANEOUS13 | KANJINTI SOLUTION RECONSTITUTED 150 |
| GENOTROPIN MINIQUICK PREFILLED | MG INTRAVENOUS27 |
| SYRINGE 1.2 MG SUBCUTANEOUS13 | KANJINTI SOLUTION RECONSTITUTED 420 |
| GENOTROPIN MINIQUICK PREFILLED | MG INTRAVENOUS27 |
| SYRINGE 1.4 MG SUBCUTANEOUS13 | KHAPZORY SOLUTION RECONSTITUTED |
| GENOTROPIN MINIQUICK PREFILLED | 175 MG INTRAVENOUS16 |
| SYRINGE 1.6 MG SUBCUTANEOUS13 | KHAPZORY SOLUTION RECONSTITUTED |
| GENOTROPIN MINIQUICK PREFILLED | 300 MG INTRAVENOUS16 |
| SYRINGE 1.8 MG SUBCUTANEOUS13 | levoleucovorin calcium pf solution 175 |
| GENOTROPIN MINIQUICK PREFILLED | mg/17.5ml intravenous16 |
| SYRINGE 2 MG SUBCUTANEOUS13 | levoleucovorin calcium pf solution 250 |
| GENVISC 850 SOLUTION PREFILLED | mg/25ml intravenous16 |
| SYRINGE 25 MG/2.5ML INTRA-ARTICULAR 31 | |

| levoleucovorin calcium solution | NUTROPIN AQ NUSPIN 10 SOLUTION PEN- |
|-------------------------------------|--|
| reconstituted 50 mg intravenous16 | INJECTOR 10 MG/2ML SUBCUTANEOUS13 |
| LUCENTIS SOLUTION 0.3 MG/0.05ML | NUTROPIN AQ NUSPIN 20 SOLUTION PEN- |
| INTRAVITREAL | INJECTOR 20 MG/2ML SUBCUTANEOUS13 |
| LUCENTIS SOLUTION PREFILLED SYRINGE | NUTROPIN AQ NUSPIN 5 SOLUTION PEN- |
| 0.3 MG/0.05ML INTRAVITREAL21 | INJECTOR 5 MG/2ML SUBCUTANEOUS13 |
| LUCENTIS SOLUTION PREFILLED SYRINGE | NYVEPRIA SOLUTION PREFILLED SYRINGE 6 |
| 0.5 MG/0.05ML INTRAVITREAL | MG/0.6ML SUBCUTANEOUS |
| MONOVISC SOLUTION PREFILLED SYRINGE | OGIVRI SOLUTION RECONSTITUTED 150 |
| 88 MG/4ML INTRA-ARTICULAR31 | MG INTRAVENOUS27 |
| MVASI SOLUTION 100 MG/4ML | OGIVRI SOLUTION RECONSTITUTED 420 |
| INTRAVENOUS5 | MG INTRAVENOUS27 |
| MVASI SOLUTION 400 MG/16ML | OMNITROPE SOLUTION CARTRIDGE 10 |
| INTRAVENOUS5 | MG/1.5ML SUBCUTANEOUS13 |
| MYOBLOC SOLUTION 10000 UNIT/2ML | OMNITROPE SOLUTION CARTRIDGE 5 |
| INTRAMUSCULAR6 | MG/1.5ML SUBCUTANEOUS13 |
| MYOBLOC SOLUTION 2500 UNIT/0.5ML | OMNITROPE SOLUTION RECONSTITUTED |
| INTRAMUSCULAR6 | 5.8 MG SUBCUTANEOUS13 |
| MYOBLOC SOLUTION 5000 UNIT/ML | ONTRUZANT SOLUTION RECONSTITUTED |
| INTRAMUSCULAR6 | 150 MG INTRAVENOUS27 |
| NEUPOGEN SOLUTION 300 MCG/ML | ONTRUZANT SOLUTION RECONSTITUTED |
| INJECTION12 | 420 MG INTRAVENOUS27 |
| NEUPOGEN SOLUTION 480 MCG/1.6ML | OPDUALAG SOLUTION 240-80 MG/20ML |
| INJECTION12 | INTRAVENOUS18 |
| NEUPOGEN SOLUTION PREFILLED SYRINGE | ORTHOVISC SOLUTION PREFILLED SYRINGE |
| 300 MCG/0.5ML INJECTION12 | 30 MG/2ML INTRA-ARTICULAR31 |
| NEUPOGEN SOLUTION PREFILLED SYRINGE | paclitaxel protein-bound part suspension |
| 480 MCG/0.8ML INJECTION12 | reconstituted 100 mg intravenous19 |
| NIVESTYM SOLUTION 300 MCG/ML | PROCRIT SOLUTION 10000 UNIT/ML |
| INJECTION12 | INJECTION8 |
| NIVESTYM SOLUTION 480 MCG/1.6ML | PROCRIT SOLUTION 2000 UNIT/ML |
| INJECTION12 | INJECTION8 |
| NIVESTYM SOLUTION PREFILLED SYRINGE | PROCRIT SOLUTION 20000 UNIT/ML |
| 300 MCG/0.5ML INJECTION12 | INJECTION8 |
| NIVESTYM SOLUTION PREFILLED SYRINGE | PROCRIT SOLUTION 3000 UNIT/ML |
| 480 MCG/0.8ML INJECTION12 | INJECTION8 |
| NORDITROPIN FLEXPRO SOLUTION PEN- | PROCRIT SOLUTION 4000 UNIT/ML |
| INJECTOR 10 MG/1.5ML SUBCUTANEOUS13 | INJECTION8 |
| NORDITROPIN FLEXPRO SOLUTION PEN- | PROCRIT SOLUTION 40000 UNIT/ML |
| INJECTOR 15 MG/1.5ML SUBCUTANEOUS13 | INJECTION8 |
| NORDITROPIN FLEXPRO SOLUTION PEN- | RELEUKO SOLUTION 300 MCG/ML |
| INJECTOR 30 MG/3ML SUBCUTANEOUS13 | INJECTION12 |
| NORDITROPIN FLEXPRO SOLUTION PEN- | releuko solution 480 mcg/1.6ml injection12 |
| INJECTOR 5 MG/1.5ML SUBCUTANEOUS 13 | |

| releuko solution prefilled syringe 300 | TREANDA SOLUTION RECONSTITUTED 100 |
|---|---------------------------------------|
| mcg/0.5ml subcutaneous12 | MG INTRAVENOUS3 |
| releuko solution prefilled syringe 480 | TREANDA SOLUTION RECONSTITUTED 25 |
| mcg/0.8ml subcutaneous12 | MG INTRAVENOUS3 |
| REMICADE SOLUTION RECONSTITUTED | TRILURON SOLUTION PREFILLED SYRINGE |
| 100 MG INTRAVENOUS | 20 MG/2ML INTRA-ARTICULAR31 |
| RENFLEXIS SOLUTION RECONSTITUTED | TRIVISC SOLUTION PREFILLED SYRINGE 25 |
| 100 MG INTRAVENOUS | MG/2.5ML INTRA-ARTICULAR31 |
| RIABNI SOLUTION 100 MG/10ML | TRUXIMA SOLUTION 100 MG/10ML |
| INTRAVENOUS22 | INTRAVENOUS22 |
| RIABNI SOLUTION 500 MG/50ML | TRUXIMA SOLUTION 500 MG/50ML |
| INTRAVENOUS22 | INTRAVENOUS22 |
| RITUXAN HYCELA SOLUTION 1400-23400 | ULTOMIRIS SOLUTION 1100 MG/11ML |
| MG -UT/11.7ML SUBCUTANEOUS22 | INTRAVENOUS28 |
| RITUXAN HYCELA SOLUTION 1600-26800 | ULTOMIRIS SOLUTION 300 MG/3ML |
| MG -UT/13.4ML SUBCUTANEOUS22 | INTRAVENOUS28 |
| RITUXAN SOLUTION 100 MG/10ML | UPLIZNA SOLUTION 100 MG/10ML |
| INTRAVENOUS22 | INTRAVENOUS29 |
| RITUXAN SOLUTION 500 MG/50ML | VABYSMO SOLUTION 6 MG/0.05ML |
| INTRAVENOUS22 | INTRAVITREAL30 |
| SAIZEN SOLUTION RECONSTITUTED 5 MG | VISCO-3 SOLUTION PREFILLED SYRINGE 25 |
| INJECTION13 | MG/2.5ML INTRA-ARTICULAR31 |
| SAIZEN SOLUTION RECONSTITUTED 8.8 | VYEPTI SOLUTION 100 MG/ML |
| MG INJECTION13 | INTRAVENOUS32 |
| SAIZENPREP SOLUTION RECONSTITUTED | VYVGART SOLUTION 400 MG/20ML |
| 8.8 MG INJECTION13 | INTRAVENOUS33 |
| SAPHNELO SOLUTION 300 MG/2ML | ZIEXTENZO SOLUTION PREFILLED SYRINGE |
| INTRAVENOUS23 | 6 MG/0.6ML SUBCUTANEOUS20 |
| SOLIRIS SOLUTION 300 MG/30ML | ZILRETTA SUSPENSION RECONSTITUTED ER |
| INTRAVENOUS24 | 32 MG INTRA-ARTICULAR 34 |
| SPINRAZA SOLUTION 12 MG/5ML | |
| INTRATHECAL25 | |
| SUPARTZ FX SOLUTION PREFILLED | |
| SYRINGE 25 MG/2.5ML INTRA-ARTICULAR 31 | |
| SUSVIMO (IMPLANT 1ST FILL) SOLUTION | |
| 10 MG/0.1ML INTRAVITREAL21 | |
| SUSVIMO (IMPLANT REFILL) SOLUTION 10 | |
| MG/0.1ML INTRAVITREAL21 | |
| SYNOJOYNT SOLUTION PREFILLED SYRINGE | |
| 20 MG/2ML INTRA-ARTICULAR31 | |
| SYNVISC SOLUTION PREFILLED SYRINGE 16 | |
| MG/2ML INTRA-ARTICULAR31 | |
| TEZSPIRE SOLUTION PREFILLED SYRINGE | |
| 210 MG/1.91ML SUBCUTANEOUS | |
| 210 11.3/ 1.5 11112 50 500 17 (142 50 51 11111 20 | |