# Pharmacy Management Drug Policy

#### SUBJECT: Step Therapy POLICY NUMBER: PHARMACY-72 EFFECTIVE DATE: 10/2011 LAST REVIEW DATE: 03/13/2025

If the member's subscriber contract excludes coverage for a specific service or prescription drug, it is not covered under that contract. In such cases, medical or drug policy criteria are not applied. This drug policy applies to the following line/s of business:

|           | Policy Application                                |                                     |  |  |
|-----------|---|-------------------------------------|--|--|
|           |   |                                     |  |  |
| Category: | ⊠ Commercial Group (e.g., EPO, HMO, POS, PPO)     | Medicare Advantage                  |  |  |
|           | ☑ On Exchange Qualified Health Plans (QHP)        | Medicare Part D                     |  |  |
|           | ☑ Off Exchange Direct Pay                         | ⊠ Essential Plan (EP)               |  |  |
|           | □ Medicaid & Health and Recovery Plans (MMC/HARP) | $\boxtimes$ Child Health Plus (CHP) |  |  |
|           | Federal Employee Program (FEP)                    | Ancillary Services                  |  |  |
|           | Dual Eligible Special Needs Plan (D-SNP)          |                                     |  |  |
|           |   |                                     |  |  |

#### DESCRIPTION:

Step Therapy encourages use of safe, cost-effective medications within different therapeutic drug categories. The entry of new generics and cost-effective therapeutic alternatives has provided an opportunity to promote these therapies as first-line.

#### POLICY:

Step Therapy requires members try certain first-line options before other medications will be considered medically necessary for treatment of a specific condition. Step therapy requirements may apply to both brands and generics. Typically, first-line medications are classified as generics, but there are instances where brand name medications may be preferred.

Based upon our review and assessment of the peer-reviewed literature, these medications have been medically proven to be effective and therefore **medically necessary** for medical treatment if the request meets the following criteria:

| ANTIBACTERIALS                                    |  |  |  |
|---|--|--|--|
| Drug  | Requirement  |  |  |
| Doryx, Doryx MPC                                  | Coverage requires documentation of serious side effects or drug failure with immediate-release doxycycline <b>AND</b> immediate-release                                    |  |  |
| Doxycycline hyclate DR                            | minocycline  |  |  |
| Clindagel 75 mL                                   | Coverage requires decompositation of conjugation side officials on drug  |  |  |
| Clindamycin 1% Gel 75 mL<br>(Oceanside & Solaris) | Coverage requires documentation of serious side effects or drug failure with generic clindamycin <b>AND</b> tretinoin  |  |  |
| Amzeeq  | Coverage requires serious side effects or drug failure with TWO topical treatments for acne (erythromycin, clindamycin, tretinoin, adapalene, dapsone, tazarotene)         |  |  |
| Zilxi 1.5%  | Coverage requires serious side effects or drug failure with topical metronidazole and one additional topical antibiotic (such as clindamycin, erythromycin, azelaic acid). |  |  |

| ANTICOAGULANTS               |      |   |  |  |
|------------------------------|------|---|--|--|
|                              | Drug | Requirement   |  |  |
| Savaysa                      |      | Coverage requires documentation of serious side effects or drug failure with Xarelto or Eliquis   |  |  |
|                              |      | ANTIDEPRESSANTS   |  |  |
|                              | Drug | Requirement   |  |  |
| Emsam                        |      | Coverage requires documentation of serious side effects or drug failure<br>with at least <b>ONE</b> of the following first line agents: escitalopram,<br>fluoxetine, citalopram, sertraline, paroxetine, mirtazapine, bupropion or  |  |  |
| Forfivo XL                   | 450  | mg venlafaxine <b>immediate-release</b> tablets or venlafaxine extended-release capsules  |  |  |
| Venlafaxir<br><b>Tablets</b> |      | <ul> <li>Veniataxine ER 75 mg capsules, taken as 3 capsules once daily</li> <li>A daily dose of 112.5 mg veniafaxine ER may be obtained by ordering veniafaxine ER 37.5 mg capsules, taken as 3 capsules once daily</li> <li>The claims processing system will not read history for this edit therefore claims will not automatically pay, therefore a manual step therapy request must be made for coverage determination</li> </ul> |  |  |
| Drizalma Sprinkle            |      |   |  |  |
|                              |      | ANTIEMETICS   |  |  |
| Drug                         |      | Requirement   |  |  |
| Anzemet                      | Cov  | verage requires documentation of serious side effects or drug failure with ondansetron  |  |  |
| Sancuso                      |      | overage requires documentation of serious side effects or drug failure with ondansetron <b>ND</b> granisetron   |  |  |
| _                            |      | ANTIFUNGAL AGENTS   |  |  |
| Drug                         |      | Requirement   |  |  |
| Ecoza                        |      |   |  |  |
| Ertaczo                      |      | Coverage requires documentation of serious side effects or drug failure with TWO  |  |  |
| Luzu                         |      | of the following generic topical antifungals: ciclopirox, econazole, ketoconazole,  |  |  |
| Luliconazo                   | ble  | - nystatin  |  |  |
| Naftifine                    |      | _   |  |  |
| Xolegel                      |      |   |  |  |
| Oxistat Lo                   | tion |   |  |  |
| Naftin of the follo          |      | Coverage requires documentation of serious side effects or drug failure with TWO of the following generic topical antifungals: ciclopirox, econazole, ketoconazole, nystatin, <b>AND</b> generic naftifine  |  |  |
|                              |      |   |  |  |

|   |  |   | ANTIMIGRAINE AGENTS   |  |  |  |
|---|--|---|---|--|--|--|
| Drug                                      |  |   | Requirement   |  |  |  |
| Zomig Nasal                               |  | TWO   | erage requires documentation of serious side effects or drug failure with<br>D generic triptans:(Almotriptan, Eletriptan, Frovatriptan, Naratriptan,<br>triptan, Sumatriptan, Zolmitriptan)   |  |  |  |
| Tosymra Cov<br>gen<br>(Alm                |  | gene<br>(Almo   | erage requires documentation of serious side effects or drug failure with<br>ric sumatriptan nasal spray <b>AND</b> TWO generics oral triptans:<br>otriptan, Eletriptan, Frovatriptan, Naratriptan, Rizatriptan, Sumatriptan,<br>itriptan)  |  |  |  |
| Zembrace                                  |  |   | rage requires documentation of serious side effects or drug failure with able sumatriptan   |  |  |  |
|   |  |   | ANTIPSYCHOTICS  |  |  |  |
| Drug                                      | Diagno   | sis   | Requirement   |  |  |  |
| Caplyta                                   | Schizophr  | renia   | Coverage requires documentation of serious side effects or drug failure with TWO generic atypical antipsychotics  |  |  |  |
| Capiyia                                   | Bipolar<br>Depressic   | on  | Coverage requires documentation of serious side effects or drug failure with TWO alternative therapies for bipolar depression   |  |  |  |
| Fanapt                                    | Schizophr  | renia   | Coverage requires documentation of serious side effects or drug failure   |  |  |  |
| Γαπαρι                                    | Bipolar<br>Disorder  |   | with TWO generic atypical antipsychotics  |  |  |  |
|   | Schizophr  |   | Coverage requires documentation of serious side effects or drug failure with TWO generic atypical antipsychotics  |  |  |  |
| Major<br>Depressiv<br>Disorder<br>Rexulti | /e   | Coverage requires documentation of serious side effects or drug failure<br>with TWO different antidepressants (with different mechanisms of<br>action) used in combination <b>OR</b> ONE antidepressant in combination<br>with ONE other augmentation therapy (such as atypical antipsychotic,<br>lithium, buspirone) |   |  |  |  |
|   | Agitation<br>associated<br>with Dementia<br>due to<br>Alzheimer<br>disease |   | Requests for this diagnosis will be approved.   |  |  |  |
| Secuado                                   | Schizophrenia  |   | Coverage requires documentation of serious side effects or drug failure with TWO generic atypical antipsychotics  |  |  |  |
|   | Schizophre   |   | Coverage requires documentation of serious side effects or drug failure   |  |  |  |
| Vraylar                                   | Bipolar<br>disorder  |   | with TWO generic atypical antipsychotics  |  |  |  |
|   | Bipolar<br>Depression  |   | Coverage requires documentation of serious side effects or drug failure with TWO alternative therapies for bipolar depression   |  |  |  |
|   | Major<br>Depressive<br>Disorder  |   | Coverage requires documentation of serious side effects or drug failure<br>with TWO different antidepressants (with different mechanisms of<br>action) used in combination <b>OR</b> ONE antidepressant in combination<br>with ONE other augmentation therapy (such as atypical antipsychotic,<br>lithium, buspirone) |  |  |  |

|                 | Paguiromont  |
|-----------------|--|
|                 | Requirement  |
| Coverage requir | es documentation of serious side effects or drug   |
|                 |  |
|                 |  |
| • ·             | es documentation of serious side effects or drug   |
|                 | lovir 5% ointment AND generic acyclovir 5% cream   |
| • .             | es documentation of serious side effects or drug failure   |
|                 | % ointment AND generic penciclovir 1% cream  |
|                 |  |
| (SELECT         | BENEFITS ONLY)   |
|                 | Requirement  |
|                 | Coverage requires documentation of serious side  |
|                 | effects or drug failure with Humalog, Humalog Mix  |
|                 | 75/25, or Insulin Lispro (Lilly authorized generic)  |
| Insulin Aspart  |  |
|                 | Coverage requires documentation of serious side  |
| volin R         | effects or drug failure with corresponding Humulin   |
|                 | product (N, R, 70-30)  |
|                 |  |
|                 |  |
|                 | Coverage requires documentation of serious side  |
|                 | effects or drug failure with Tradjenta or Jentadueto   |
|                 |  |
|                 |  |
|                 | Coverage requires documentation of serious side  |
|                 | effects or drug failure with generic immediate-release   |
| ortamet and     | metformin <b>AND</b> generic extended-release metformin  |
|                 | (generic equivalent of Glucophage XR)  |
|                 | Coverage of any non-preferred blood glucose meter  |
| est Strips      | or test strip requires either: a previous trial and failure <b>OR</b> the inability to use any Abbott (Freestyle |
| -               |  |
|                 | or Precision Xtra) or One Touch products   |
|                 | Coverage requires documentation of serious side effects <b>OR</b> drug failure with Glyxambi                     |
|                 | Coverage requires documentation of serious side  |
| gluromet        | effects or drug failure with Xigduo XR AND   |
| -               | Synjardy/Synjardy XR   |
|                 | Coverage requires documentation of serious side  |
|                 | effects or drug failure with Farxiga AND Jardiance   |
|                 | Coverage requires documentation of serious side  |
|                 | effects or drug failure with Tradjenta, Jentadueto, or   |
| )               | Jentadueto XR  |
|                 | Coverage requires documentation of serious side  |
| xin)            | effects or drug failure with Glyxambi  |
|                 |  |
|                 |  |
|                 |  |
|                 | failure with acyc<br>Coverage requir<br>failure with acyc<br>Coverage requir<br>with acyclovir 59<br>BLOOD GLU   |

|   |                                     | CARDIOVASCULAR AGENTS  |  |  |
|---|-------------------------------------|--|--|--|
| Drug  |                                     | Requirement  |  |  |
| Edarbi  | the following: losa                 | Coverage requires documentation of serious side effects or drug failure with TWO of the following: losartan, irbesartan, valsartan   |  |  |
| Edarbyclor  | Coverage requires                   | documentation of serious side effects or drug failure with TWO of rtan/hctz, irbesartan/hctz, valsartan/hctz   |  |  |
| Thalitone   | Coverage requires chlorthalidone.   | documentation of serious side effects or drug failure with generic   |  |  |
|   | CARD                                | IOVASCULAR AGENTS, DYSLIPIDEMICS   |  |  |
| Drug  |                                     | Requirement  |  |  |
| Livalo<br>Pitavastatin<br>Calcium<br>Zypitamag  |                                     | n of serious side effects or drug failure with TWO of the following<br>atorvastatin, fluvastatin, lovastatin, pravastatin, rosuvastatin,   |  |  |
| Praluent  |                                     | ires documentation of serious side effects or drug failure Repatha for years and older.  |  |  |
| Nexletol,<br>Nexlizet   |                                     | ires documentation of serious side effects or drug failure with one atorvastatin, fluvastatin, lovastatin, pitavastatin, pravastatin, imvastatin   |  |  |
|   |                                     | NEUROLOGICAL AGENTS  |  |  |
| Drug  |                                     | Requirement  |  |  |
| Savella   | Coverage require duloxetine         | es documentation of serious side effects or drug failure with  |  |  |
| Adlarity  | <b>-</b> .                          | es documentation of serious side effects or drug failure of ezil ODT, galantamine, <b>OR</b> rivastigmine  |  |  |
| Xadago  | Coverage require generic selegiline | es documentation of serious side effects or drug failure with  |  |  |
|   |                                     | DERMATOLOGICAL AGENTS  |  |  |
|   | Drug                                | Requirement  |  |  |
| Aczone 7.5%, Dapsone 7.5%   |                                     | Coverage requires documentation of serious side effects or drug failure with a topical retinoid <b>AND</b> Dapsone 5%  |  |  |
| Adapalene 0.1% Lotion, Soln,<br>Swab<br>Differin 0.1% Lotion  |                                     | Coverage requires documentation of serious side effects or drug failure with adapalene cream or gel <b>AND</b> tretinoin cream or gel  |  |  |
| Eucrisa Ointment  |                                     | Coverage requires documentation of serious side effects or drug<br>failure with ONE generic topical steroid (aclometasone,<br>amcinonide, betamethasone, clobetasol, desonide,<br>desoximetasone, diflorasone, fluocinolone, fluocinonide–E,<br>fluticasone, halobetasol, hydrocortisone 2.5%, hydrocortisone<br>valerate, mometasone, prednicarbate, triamcinolone) <b>OR</b> ONE of<br>the following: tacrolimus ointment or pimecrolimus cream. |  |  |
| Noritate  |                                     | Coverage requires documentation of serious side effects or drug failure with generic metronidazole cream, gel, or lotion   |  |  |
| Zyclara 2.5% Cream Pump,<br>Zyclara 3.75% Cream and<br>Zyclara 3.75% Cream Pump<br>Imiquimod 3.75% Cream, and<br>Imiquimod 3.75% Cream Pump |                                     | Coverage requires documentation of serious side effects or drug failure with imiquimod 5% cream  |  |  |

| GASTROINTESTINAL AGENTS                               |  |         |   |  |
|---|--|---------|---|--|
|   | Drug                                     |         | Requirement   |  |
| Amitiza   | Chronic idiopathic constipation or IBS-C |         | Coverage requires documentation of serious side<br>effects or drug failure with lubiprostone <b>AND</b> <u>either</u><br>Linzess <b>OR</b> Trulance for a diagnosis of chronic<br>idiopathic constipation or irritable bowel syndrome with<br>constipation. |  |
|   | Opioid-induced constipati                | on      | Coverage requires documentation of drug failure or serious side effects with Movantik for a diagnosis of opioid induced constipation.   |  |
| Motegrity   |  |         | Coverage requires documentation of serious side<br>effects or drug failure with Linzess OR Trulance for a<br>diagnosis of chronic idiopathic constipation (CIC)   |  |
| Relistor Tab  | olet                                     |         | Coverage requires documentation of serious side   |  |
| Symproic  |  |         | effects or drug failure with Movantik for a diagnosis of opioid-induced constipation  |  |
| Ibsrela   |  |         | Coverage requires documentation of serious side<br>effects or drug failure with Linzess, lubiprostone, <b>AND</b><br>Trulance for a diagnosis of irritable bowel syndrome<br>with constipation  |  |
|   | Omeprazole/Sodium Bicarbonate Packets    |         | Coverage requires documentation of serious side<br>effects or drug failure with THREE of the following:   |  |
| Zegerid Pac   | ckets                                    |         | omeprazole, pantoprazole, lansoprazole, rabeprazole   |  |
| Pheburane   |  |         | Coverage requires documentation of serious side<br>effects or drug failure with generic sodium<br>phenylbutyrate  |  |
|   | GENITOURINARY                            | AGEN    | ITS; ANTISPASMODICS, URINARY  |  |
| Drug  |  |         | Requirement   |  |
|   |  |         | n of serious side effects or drug failure with TWO of the<br>ER, tolterodine, trospium, trospium XR   |  |
| Gelnique  | Gelnique does not require s              | sten th | Exception:<br>herapy for individuals 65 years of age or older   |  |
|   |  | -       | NT/REPLACEMENT/MODIFYING (ADRENAL)  |  |
|   | Drug                                     |         | Requirement   |  |
| Bryhali   |  | Cove    | erage requires documentation of a serious side effects  |  |
|   | ocortolone Pivalate                      |         | rug failure with TWO of the following generic topical   |  |
| Cordran (Cream, Lotion, Ointment)                     |  | sterc   | pids:   |  |
| Desonide 0.05% Gel                                    |  |         |   |  |
|   |  |         | metasone, amcinonide, betamethasone, clobetasol,  |  |
|   |  |         | onide, desoximetasone, diflorasone, fluocinolone,<br>cinonide–E, fluticasone, halobetasol (except foam),  |  |
| ппрекю  |  |         | ocortisone 2.5%, hydrocortisone valerate,   |  |
| Impoyz Cream (and generic<br>clobetasol 0.025% cream) |  | -       | ocortisone butyrate (except lotion), mometasone,  |  |
| Lexette   |  |         | nicarbate, triamcinolone  |  |
| Pandel  |  |         |   |  |
| Sernivo Loti  | on                                       |         |   |  |
| Ultravate Lo  |  | 1       |   |  |
| Verdeso   |  | ]       |   |  |

|  | IMMUNOLOGICAL AGENTS               |                       |  |  |  |
|--|------------------------------------|-----------------------|--|--|--|
| Drug                                       | Drug                               |                       |  | Requirement  |  |
| Prograf Grar                               | nules                              |                       | Must have documentation of serious side effects or drug failure with generic tacrolimus capsules <b>Exception:</b> age less than 9 years old |  |  |
|  |                                    |                       | Μ  | ULTIPLE SCLEROSIS AGENTS   |  |
| Drug                                       |                                    |                       |  | Requirement  |  |
| Bafiertam                                  |                                    | -                     |  | res documentation of serious side effects or drug failure with   |  |
| Ponvory                                    |                                    |                       |  | owing agents: Avonex, Copaxone 40mg, glatiramer, Glatopa,  |  |
| Vumerity                                   |                                    | fingolimo<br>or Zepos |  | ethyl fumarate, Mayzent, Plegridy, Rebif, teriflunomide, Kesimpta,   |  |
|  |                                    |                       | Γ  | OPHTHALMIC AGENTS  |  |
| D  | rug                                |                       |  | Requirement  |  |
| Zerviate                                   |                                    |                       | with T   | rage requires documentation of serious side effects or drug failure<br>WO of the following antihistamine eye drops: azelastine,<br>tadine, epinastine          |  |
| Xelpros                                    | Vyzu                               | lta                   |  |  |  |
| Zioptan                                    |                                    |                       | Cove   | rage requires documentation of serious side effects or drug failure  |  |
| lyuzeh                                     |                                    |                       | with L   | umigan AND either latanoprost or travoprost  |  |
| Tafluprost                                 |                                    |                       |  |  |  |
| Rhopressa,                                 | Rhopressa, Rocklatan with a        |                       | with a   | rage requires documentation of serious side effects or drug failure<br>ny covered prostaglandin analogue (such as bimatoprost,<br>prost, latanoprost, Lumigan) |  |
| Restasis 0.0                               | )5%                                |                       | Cove   | rage requires documentation of serious side effects or drug failure  |  |
| Restasis Mu                                | Itidos                             | e 0.05%               | of cyc   | losporine 0.05% eye emulsion AND Xiidra 5% eye drops   |  |
|  |                                    |                       | rage requires documentation of serious side effects or drug failure neric atropine 1% drops  |  |  |
|  | PANCREATIC ENZYMES                 |                       |  |  |  |
| Drug                                       |                                    |                       |  | Requirement  |  |
| PancreazeCoverage requires doPertzyeZenpep |                                    | ires do               | ocumentation of serious side effects or drug failure with Creon and  |  |  |
|  | RESPIRATORY TRACT/PULMONARY AGENTS |                       |  |  |  |
| Drug                                       |                                    |                       |  | Requirement  |  |
| Tudorza Pressair                           |                                    |                       | Coverage requires documentation of serious side effects or drug failure with ONE of the following: tiotropium bromide or Incruse.            |  |  |
| Alvesco                                    |                                    |                       | Coverage requires documentation of serious side effects or drug  |  |  |
| Pulmicort Flexhaler                        |                                    |                       | failure with ONE of the following: Arnuity Ellipta, Asmanex, or  |  |  |
|  | Armonair Digihaler                 |                       |  | Qvar Redihaler.  |  |
| AirDuo Respiclick                          |                                    |                       | Coverage requires documentation of severe intolerance or   |  |  |
| AirDuo Digihaler                           |                                    |                       | therapeutic failure with generic fluticasone/salmeterol inhaler  |  |  |
| Lonhala Magnair 25 mcg Starter             |                                    | arter                 | Coverage requires documentation of serious side effects or drug  |  |  |
|  | Lonhala Magnair 25 mcg Refill      |                       |  | failure with any TWO of the following long-acting muscarinic receptor antagonists (LAMA) containing inhalers: Anoro Ellipta,                                   |  |
| Yupelri                                    |                                    |                       | Bevespi Aerosphere, Incruse Ellipta, Neohaler, tiotropium bromide<br>Handihaler, Spiriva Respimat, Stiolto Respimat, or Utibron              |  |  |

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| Duaklir Pressair      |                           | Coverage requires serious side effects or drug failure with at<br>least TWO long-acting muscarinic receptor antagonist/long-<br>acting beta agonist (LAMA/LABA) agents. Agents include:<br>Anoro, Bevespi, Stiolto and Utibron. |  |  |
|-----------------------|---------------------------|---|--|--|
|                       | SE                        | LECTIVE ESTROGEN RECEPTOR MODIFYING AGENTS  |  |  |
| Drug                  |                           | Requirement   |  |  |
| Estring               | Coverage                  | requires documentation of serious side effects or drug failure with a topical   |  |  |
| Osphena               |                           | trogen product such as Premarin cream or estradiol vaginal cream.   |  |  |
|                       | SKELETAL MUSCLE RELAXANTS |   |  |  |
| Drug                  |                           | Requirement   |  |  |
| Norgesic F            | orte                      | overage requires documentation of serious side effects or drug failure with   |  |  |
| Orphenges             | sic Forte                 | THREE of the following (generic) agents: baclofen, carisoprodol,  |  |  |
|                       |                           | chlorzoxazone, cyclobenzaprine, methocarbamol, metaxalone, orphenadrine,  |  |  |
| Aspirin/Caffeine      |                           | tizanidine  |  |  |
| SLEEP DISORDER AGENTS |                           |   |  |  |
| Drug                  |                           | Requirement   |  |  |
| Edluar                |                           | Coverage requires documentation of serious side effects or drug failure with  |  |  |
| Zolpimist zo          |                           | zolpidem  |  |  |
|                       |                           | Coverage requires documentation of serious side effects or drug failure with  |  |  |
|                       |                           | TWO of the following: zolpidem, eszopiclone, zaleplon   |  |  |

#### POLICY GUIDELINES:

- 1. This policy is applicable to drugs that are included on a specific drug formulary. If a drug referenced in this policy is non-formulary, please reference the Coverage Exception Evaluation Policy for All Lines of Business Formularies policy for review guidelines.
- 2. Supportive documentation of previous drug use must be submitted for any criteria requiring trial of a preferred agent if the preferred drug is not found in claims history.
- 3. Approval for step therapy requirements may not bypass MAC penalty. Please see MAC penalty policy for detail of this benefit.
- 4. Utilization Management are contract dependent and coverage criteria may be dependent on the contract renewal date. Additionally, coverage of drugs listed in this policy are contract dependent. Refer to specific contract/benefit language for exclusions.
- 5. For contracts where Insurance Law § 4903(c-1), and Public Health Law § 4903(3-a) are applicable, if trial of preferred drug(s) is the only criterion that is not met for a given condition, and one of the following circumstances can be substantiated by the requesting provider, then trial of the preferred drug(s) will not be required.
  - a. The required prescription drug(s) is (are) contraindicated or will likely cause an adverse reaction or physical or mental harm to the member;
  - b. The required prescription drug is expected to be ineffective based on the known clinical history and conditions and concurrent drug regimen;
  - c. The required prescription drug(s) was (were) previously tried while under the current or a previous health plan, or another prescription drug or drugs in the same pharmacologic class or with the same mechanism of action was (were) previously tried and such prescription drug(s) was (were) discontinued due to lack of efficacy or effectiveness, diminished effect, or an adverse event;
  - d. The required prescription drug(s) is (are) not in the patient's best interest because it will likely cause a significant barrier to adherence to or compliance with the plan of care, will likely worsen

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a comorbid condition, or will likely decrease the ability to achieve or maintain reasonable functional ability in performing daily activities;

- e. The individual is stable on the requested prescription drug. The medical profile of the individual (age, disease state, comorbidities), along with the rational for deeming stability as it relates to standard medical practice and evidence-based practice protocols for the disease state will be taken into consideration.
- f. The above criteria are not applicable to requests for brand name medications that have an AB rated generic. We can require a trial of an AB-rated generic equivalent prior to providing coverage for the equivalent brand name prescription drug.
- 6. Initial approval will be granted for a period of 1 year.
  - Continued approval at time of recertification will require documentation that the drug is providing ongoing benefit to the patient in terms of improvement or stability in disease state or condition.
- 7. Clinical documentation must be submitted for each request (initial and recertification) unless otherwise specified (e.g., provider attestation required). Supporting documentation includes, but is not limited to, progress notes documenting previous treatments/treatment history, diagnostic testing, laboratory test results, genetic testing/biomarker results, imaging and other objective or subjective measures of benefit which support continued use of the requested product is medically necessary. Also, ongoing use of the requested product must continue to reflect the current policy's preferred formulary. Recertification reviews may result in the requirement to try more cost-effective treatment alternatives as they become available (i.e., generics, biosimilars, or other guideline supported treatment options). Requested dosing must continue to be consistent with FDA-approved or off-label/guideline-supported dosing recommendations.
- 8. In addition to the full prescribing information for each individual drug, the corresponding clinical guidelines (i.e., NCCN, DSM, etc.) are reviewed on an annual basis to determine the appropriateness of the medical necessity criteria that is applied.
- 9. All requests will be reviewed to ensure they are being used for an appropriate indication and may be subject to an off-label review in accordance with our Off-Label Use of FDA Approved Drugs Policy (Pharmacy-32)
- 10. All utilization management requirements outlined in this policy are compliant with applicable New York State insurance laws and regulations. Policies will be reviewed and updated as necessary to ensure ongoing compliance with all state and federally mandated coverage requirements.

| Date       | Revision                        |
|------------|---------------------------------|
| 03/13/2025 | Revised                         |
| 03/06/2025 | Revised                         |
| 01/01/2025 | Revised                         |
| 11/21/2024 | P&T Committee Review / Approval |
| 10/21/2024 | Revised                         |
| 09/23/2024 | Revised                         |
| 09/13/2024 | Revised                         |
| 08/13/2024 | Revised                         |
| 05/10/2024 | Revised                         |
| 04/09/2024 | Revised                         |
| 03/14/2024 | Revised                         |
| 02/08/2024 | Revised                         |
| 01/01/2024 | Revised                         |
| 12/06/2023 | Revised                         |

#### **UPDATES**:

| 11/30/2023 | P&T Committee Approval           |
|------------|----------------------------------|
| 11/10/2023 | Revised                          |
| 9/7/2023   | Revised                          |
| 8/10/2023  | Revised                          |
| 7/7/2023   | Revised                          |
| 6/8/2023   | Revised                          |
| 4/24/2023  | Revised                          |
| 4/5/2023   | Revised                          |
| 3/31/2023  | Revised                          |
| 3/16/2023  | Revised                          |
| 2/9/2023   | Revised                          |
| 2/3/2023   | Revised                          |
| 12/20/2022 | Revised                          |
| 12/15/2022 | Revised                          |
| 12/2/22    | Revised                          |
| 11/17/2022 | P&T Committee Approval           |
| 11/3/22    | Revised                          |
| 10/3/22    | Revised                          |
| 8/29/22    | Revised                          |
| 8/25/22    | Revised                          |
| 7/28/22    | Revised                          |
| 6/30/22    | Revised                          |
| 6/3/22     | Revised                          |
| 5/12/22    | Revised                          |
| 5/9/2022   | Revised                          |
| 05/05/2022 | P&T Committee Approval           |
| 5/1/2022   | Revised                          |
| 3/29/22    | Revised                          |
| 3/18/22    | Revised                          |
| 2/18/22    | Revised                          |
| 2/8/22     | Revised / P&T Committee Approval |
| 1/22       | Revised                          |
| 12/21      | Revised                          |
| 11/21      | Revised                          |
| 10/21      | Revised                          |
| 9/21       | Revised                          |
| 8/21       | Revised                          |
| 5/21       | Revised                          |
| 4/21       | Revised                          |
| 3/21       | Revised                          |
| 2/11/2021  | P&T Committee Approval           |
| 1/21       | Revised                          |
| 12/20      | Revised                          |
| 10/20      | Revised                          |
| 8/2020     | Revised                          |
| 7/2020     | Revised                          |
| 6/2020     | i të nësa                        |
| 0/2020     | Revised                          |

| 3/20          | Revised  |
|---------------|--|
| 2/20          | Revised  |
| 1/20          | Revised  |
| 12/19         | Revised  |
| 11/19         | Revised  |
| 10/19         | Revised  |
| 8/19          | Revised  |
| 7/19          | Revised  |
| 5/19          | P&T Committee Approval   |
| 4/19          | Revised  |
| 3/19          | Revised  |
| 2/19          | Revised  |
| 1/19          | Revised  |
| 11/18         | Revised  |
| 10/18         | Revised  |
| 9/18          | Revised  |
| 5/18          | Revised  |
| 4/18          | Revised  |
| 3/18          | Revised  |
| 2/18          | Revised  |
| 1/18          | Revised- Both STEP Policies combined to one policy The Commercial Open step therapy and  |
|               | Exchange Closed/CHP policies have been merged. The policy has also been changed into a table format with headers that match the web formularies (derived from RxFlex). |
| 12/17         | Revised  |
| 11/2017       | P&T Committee Approval   |
| 9/17          | Revised  |
| 7/17          | Revised  |
| 5/17          | Revised  |
| 4/17          | Revised  |
| 1/17          | Revised  |
| 10/16         | Revised  |
| 9/16          | Revised  |
| 8/16          | Revised  |
| 7/16          | Revised  |
| 6/16          | Revised  |
| 5/16          | Revised  |
| 4/16          | Revised  |
| 3/16          | Revised  |
| 1/16          | Revised  |
| 12/15         | Revised  |
| 11/15         | Revised  |
| 8/15          | Revised  |
| 7/15          | Revised  |
| 6/15          | Revised  |
| 5/15          | Revised  |
| 4/15          | Revised  |
| 3/15          |  |
|               | Revised  |
| 1/15<br>11/14 | Revised<br>Revised   |

| 10/14 | Revised |  |
|-------|---------|--|
| 8/14  | Revised |  |
| 7/14  | Revised |  |
| 5/14  | Revised |  |
| 3/14  | Revised |  |
| 1/14  | Created |  |