SUBJECT: Step Therapy POLICY NUMBER: PHARMACY-72 EFFECTIVE DATE: 10/2011 LAST REVIEW DATE: 03/13/2025					
	subscriber contract excludes coverage for a specific service or pract. In such cases, medical or drug policy criteria are not appliefollowing 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)	□ 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 AND immediate-release			
Doxycycline hyclate DR	minocycline			
Clindagel 75 mL	Covernment and accompanies of a prince aid affects on during			
Clindamycin 1% Gel 75 mL (Oceanside & Solaris)	Coverage requires documentation of serious side effects or drug failure with generic clindamycin AND 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
Τ	Drug		Requirement
Emsam			Coverage requires documentation of serious side effects or drug failure with at least ONE of the following first line agents: escitalopram, fluoxetine, citalopram, sertraline, paroxetine, mirtazapine, bupropion or
Forfivo XL	450	mg	venlafaxine immediate-release tablets or venlafaxine extended-release capsules
Venlafaxir Tablets Drizalma S Drug Anzemet	Sprink	Nenlafaxine ER 75 mg capsules, taken as 3 capsules once daily A daily dose of 112.5 mg venlafaxine ER may be obtained by ordering venlafaxine ER 37.5 mg capsules, taken as 3 capsules one daily 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 Coverage requires serious side effects or drug failure with duloxetine	
Sancuso	Coverage requires documentation of serious side effects or drug failure with ondansetron		
			ANTIFUNGAL AGENTS
Drug Requirement		Requirement	
1 1 1 1 7 1 1		of the follo	requires documentation of serious side effects or drug failure with TWO owing generic topical antifungals: ciclopirox, econazole, ketoconazole,
Naftin	Coverage requires documentation of serious side effects or drug failure with TWO		

			ANTIMIGRAINE AGENTS		
Dr	ug		Requirement		
Zomig Nasal TW		TWC	erage requires documentation of serious side effects or drug failure with D generic triptans:(Almotriptan, Eletriptan, Frovatriptan, Naratriptan, triptan, Sumatriptan, Zolmitriptan)		
Tosymra Cove gene (Alm		gene (Alm	erage requires documentation of serious side effects or drug failure with eric sumatriptan nasal spray AND TWO generics oral triptans: otriptan, Eletriptan, Frovatriptan, Naratriptan, Rizatriptan, Sumatriptan, itriptan)		
Zembrace)		erage requires documentation of serious side effects or drug failure with table sumatriptan		
			ANTIPSYCHOTICS		
Drug	Diagno	sis	Requirement		
	Schizophr		Coverage requires documentation of serious side effects or drug failure with TWO generic atypical antipsychotics		
Caplyta	Bipolar Depression	n	Coverage requires documentation of serious side effects or drug failure with TWO alternative therapies for bipolar depression		
Fanapt	Schizophr	enia	Coverage requires documentation of serious side effects or drug failure		
Тапарт	Bipolar Disorder		with TWO generic atypical antipsychotics		
	Schizophrenia		Coverage requires documentation of serious side effects or drug failure with TWO generic atypical antipsychotics		
Rexulti	Major Depressive Disorder		Coverage requires documentation of serious side effects or drug failure with TWO different antidepressants (with different mechanisms of action) used in combination OR ONE antidepressant in combination with ONE other augmentation therapy (such as atypical antipsychotic, lithium, buspirone)		
	Agitation associated with Dementia due to Alzheimer disease		Requests for this diagnosis will be approved.		
Secuado			Coverage requires documentation of serious side effects or drug failure with TWO generic atypical antipsychotics		
	Schizophrenia		Coverage requires documentation of serious side effects or drug failure		
	Bipolar disorder		with TWO generic atypical antipsychotics		
	Bipolar		Coverage requires documentation of serious side effects or drug failure		
Vraylar	Depression	n	with TWO alternative therapies for bipolar depression		
Major Depressive Disorder		⁄e	Coverage requires documentation of serious side effects or drug failure with TWO different antidepressants (with different mechanisms of action) used in combination OR ONE antidepressant in combination with ONE other augmentation therapy (such as atypical antipsychotic, lithium, buspirone)		

	AN	NTIVIRALS
Drug		Requirement
cyclovir 5% cream		res documentation of serious side effects or drug
Penciclovir 1% cream Coverage requir		clovir 5% ointment.
Xerese 5%-1% cream		
Zovirax 5% cream		res documentation of serious side effects or drug
2011ax 070 010a111		clovir 5% ointment AND generic acyclovir 5% cream
Denavir 1% cream		res documentation of serious side effects or drug failure
20114111 170 0104111		% ointment AND generic penciclovir 1% cream
		COSE REGULATORS
Drug	(SELECT	BENEFITS ONLY)
Drug		Requirement
Admelog		Coverage requires documentation of serious side
Apidra		effects or drug failure with Humalog, Humalog Mix
Fiasp Noveled Noveled Mix 70/20	Inoulin Assaut	75/25, or Insulin Lispro (Lilly authorized generic)
Novolog, Novolog Mix 70/30,	іпѕиіп Аѕрап	
Nevelie 70 00 Nevelie N. Ne		Coverage requires documentation of serious side
Novolin 70-30, Novolin N, No	volin K	effects or drug failure with corresponding Humulin
Nicologo		product (N, R, 70-30)
Nesina		
Alogliptin		
Kazano		Coverage requires documentation of serious side
Alogliptin/metformin		effects or drug failure with Tradjenta or Jentadueto
Oseni		
Alogliptin/pioglitazone		
Glumetza		Coverage requires documentation of serious side effects or drug failure with generic immediate-release metformin AND generic extended-release metformin
Fortamet		
Metformin ER (generics of Fo		
Glumetza), Metformin HCl 62	25 mg	(generic equivalent of Glucophage XR)
Blood Glucose Meters and Test Strips		Coverage of any non-preferred blood glucose meter or test strip requires either: a previous trial and failure OR the inability to use any Abbott (Freestyle or Precision Xtra) or One Touch products
Qtern		Coverage requires documentation of serious side
		effects OR drug failure with Glyxambi
		Coverage requires documentation of serious side
Invokamet, Invokamet Xr, Se	gluromet	effects or drug failure with Xigduo XR AND
invokamet, invokamet XI, Oegiaromet		Synjardy/Synjardy XR
		Coverage requires documentation of serious side
Invokana, Steglatro		effects or drug failure with Farxiga AND Jardiance
Januvia (sitagliptin), Janumet and Janumet XR (sitagliptin and metformin)		Coverage requires documentation of serious side effects or drug failure with Tradjenta, Jentadueto, or Jentadueto XR
Steglujan (ertugliflozin/sitagliptin)		Coverage requires documentation of serious side effects or drug failure with Glyxambi

		CARRIOVASCIII AR ACENTS		
Drug		CARDIOVASCULAR AGENTS		
Drug	Requirement			
Edarbi	Coverage requires documentation of serious side effects or drug failure with TWO of the following: losartan, irbesartan, valsartan			
Edarbyclor	<u> </u>	s documentation of serious side effects or drug failure with TWO of artan/hctz, irbesartan/hctz, valsartan/hctz		
	ŭ	s documentation of serious side effects or drug failure with generic		
Thalitone	chlorthalidone.			
_	CARI	DIOVASCULAR AGENTS, DYSLIPIDEMICS		
Drug		Requirement		
Livalo Pitavastatin Calcium Zypitamag		Documentation of serious side effects or drug failure with TWO of the following generic statins: atorvastatin, fluvastatin, lovastatin, pravastatin, rosuvastatin, simvastatin		
Praluent		uires documentation of serious side effects or drug failure Repatha for years and older.		
Nexletol, Nexlizet	Coverage requ	uires documentation of serious side effects or drug failure with one atorvastatin, fluvastatin, lovastatin, pitavastatin, pravastatin,		
	,	NEUROLOGICAL AGENTS		
Drug		Requirement		
Savella	Coverage requires documentation of serious side effects or drug failure with duloxetine			
AM300		es documentation of serious side effects or drug failure of pezil ODT, galantamine, OR rivastigmine		
Xadago	Coverage require generic selegiline	es documentation of serious side effects or drug failure with		
	<u> </u>	DERMATOLOGICAL AGENTS		
	Drug	Requirement		
Aczone 7.5%, Dapsone 7.5%		Coverage requires documentation of serious side effects or drug failure with a topical retinoid AND Dapsone 5%		
Adapalene 0.1% Lotion, Soln, Swab Differin 0.1% Lotion		Coverage requires documentation of serious side effects or drug failure with adapalene cream or gel AND tretinoin cream or gel		
Eucrisa Ointment		Coverage requires documentation of serious side effects or drug failure with ONE generic topical steroid (aclometasone, amcinonide, betamethasone, clobetasol, desonide, desoximetasone, diflorasone, fluocinolone, fluocinonide–E, fluticasone, halobetasol, hydrocortisone 2.5%, hydrocortisone valerate, mometasone, prednicarbate, triamcinolone) OR ONE of 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, Zyclara 3.75% Cream and Zyclara 3.75% Cream Pump Imiquimod 3.75% Cream, and Imiquimod 3.75% Cream Pump		Coverage requires documentation of serious side effects or drug failure with imiquimod 5% cream		

	GAS	TROIN	NTESTINAL AGENTS
	Drug		Requirement
Amitiza	Chronic idiopathic constipation or IBS-C		Coverage requires documentation of serious side effects or drug failure with lubiprostone AND <u>either</u> Linzess OR Trulance for a diagnosis of chronic idiopathic constipation or irritable bowel syndrome with constipation.
	Opioid-induced constipation		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 effects or drug failure with Linzess OR Trulance for a diagnosis of chronic idiopathic constipation (CIC)
Relistor Ta	ablet		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 effects or drug failure with Linzess, lubiprostone, AND Trulance for a diagnosis of irritable bowel syndrome with constipation
	le/Sodium Bicarbonate Pacl	kets	Coverage requires documentation of serious side effects or drug failure with THREE of the following:
Zegerid Pa	ackets		omeprazole, pantoprazole, lansoprazole, rabeprazole
Pheburane	Pheburane		Coverage requires documentation of serious side effects or drug failure with generic sodium phenylbutyrate
	GENITOURINARY	AGEN	ITS; ANTISPASMODICS, URINARY
Drug			Requirement
Oxytrol			n of serious side effects or drug failure with TWO of the ER, tolterodine, trospium, trospium XR Exception:
Gelnique	Gelnique Golpique does not require str		erapy for individuals 65 years of age or older
H	·		NT/REPLACEMENT/MODIFYING (ADRENAL)
	Drug		Requirement
Bryhali	• 9	Cove	erage requires documentation of a serious side effects
Cloderm, (Clocortolone Pivalate	or dr	ug failure with TWO of the following generic topical
Cordran (Cream, Lotion, Ointment)		stero	ids:
Desonide 0.05% Gel		╡.	
7 10.10 9, 7 10.10 110.10			metasone, amcinonide, betamethasone, clobetasol,
Traiobetasor i Topioriate 0.0070 i dani			nide, desoximetasone, diflorasone, fluocinolone, inonide–E, fluticasone, halobetasol (except foam),
Пірекіо			ocortisone 2.5%, hydrocortisone valerate,
impoyz Cream (and generic			ocortisone butyrate (except lotion), mometasone,
clobetasol 0.025% cream)			nicarbate, triamcinolone
Lexette		-	
Pandel Sernivo Lo	ation	-	
Ultravate L		1	
Verdeso	-04011	1	
		1	

Step Therapy Policy

IMMUNOLOGICAL AGENTS					
Drug		Requirement			
Prograf Gra	nules			umentation of serious side effects or drug failure with generic sules Exception: age less than 9 years old	
			M	ULTIPLE SCLEROSIS AGENTS	
Drug				Requirement	
Bafiertam				ires documentation of serious side effects or drug failure with	
Ponvory				owing agents: Avonex, Copaxone 40mg, glatiramer, Glatopa,	
Vumerity		_		ethyl fumarate, Mayzent, Plegridy, Rebif, teriflunomide, Kesimpta,	
,		or Zepos	ola.	OPHTHALMIC AGENTS	
D	rug			Requirement	
Zerviate	<u>iug</u>		with T	rage requires documentation of serious side effects or drug failure WO of the following antihistamine eye drops: azelastine, tadine, epinastine	
Xelpros	Vyzu	lta			
Zioptan				rage requires documentation of serious side effects or drug failure	
lyuzeh			with L	umigan AND either latanoprost or travoprost	
Tafluprost					
Rhopressa, Rocklatan with a		with a	rage requires documentation of serious side effects or drug failure iny covered prostaglandin analogue (such as bimatoprost, brost, latanoprost, Lumigan)		
Restasis 0.05% Cover			rage requires documentation of serious side effects or drug failure closporine 0.05% eye emulsion AND Xiidra 5% eye drops		
Atropine Su	lfate/P	F		rage requires documentation of serious side effects or drug failure neric atropine 1% drops	
				PANCREATIC ENZYMES	
Drug				Requirement	
Pancreaze Pertzye	Pancreaze Coverage requires do		uires do	ocumentation of serious side effects or drug failure with Creon and	
. o.t.=yo		RE	SPIR/	ATORY TRACT/PULMONARY AGENTS	
	Dru			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		efill	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 Handihaler, Spiriva Respimat, Stiolto Respimat, or Utibron		

Step Therapy Policy

Duaklir Pressair		Coverage requires serious side effects or drug failure with at least TWO long-acting muscarinic receptor antagonist/long-acting beta agonist (LAMA/LABA) agents. Agents include: Anoro, Bevespi, Stiolto and Utibron.				
SELECTIVE ESTROGEN RECEPTOR MODIFYING AGENTS						
Drug		Requirement				
Estring	Coverage	requires documentation of serious side effects or drug failure with a topical				
Osphena	vaginal es	strogen product such as Premarin cream or estradiol vaginal cream.				
	SKELETAL MUSCLE RELAXANTS					
Drug		Requirement				
Norgesic F	orte	coverage requires documentation of serious side effects or drug failure with				
		THREE of the following (generic) agents: baclofen, carisoprodol,				
Orphenadi	rine/	chlorzoxazone, cyclobenzaprine, methocarbamol, metaxalone, orphenadrine,				
Aspirin/Caffeine tizanid		tizanidine				
	SLEEP DISORDER AGENTS					
Drug		Requirement				
Edluar		Coverage requires documentation of serious side effects or drug failure with				
Zolpimist		zolpidem				
		Coverage requires documentation of serious side effects or drug failure with				
Quviviq TW		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

Step Therapy Policy

- 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.

UPDATES:

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

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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
1710	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
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8/15	Revised
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11/14	Revised

10/14	Revised	
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1/14	Created	