



Virginia's Medicaid Preferred Drug List (PDL)/ Common Core Formulary

4/1/26

Virginia's Medicaid Pharmacy Benefits Management System

Phone: 800-932-6648 Fax: 800-932-6651

General Information:

- Virginia's Medicaid Preferred Drug List (PDL)/ Common Core Formulary only includes select drug classes, other classes will pay such as but not limited to diuretics, many cardiac agents, many antibiotics etc.
- PDL preferred drugs do not require Service Authorizations (SA) unless subject to additional clinical criteria (e.g., long-acting opioids, hepatitis C therapies, growth hormone)
- Non-preferred drugs require a SA
- Drugs not on the PDL are subject to Virginia's mandatory generic substitution requirements.
- SAs may be submitted by fax, phone or WebPA, e-PA. For urgent requests, please call **800-932-6648**. Fax requests receive a response within 24 hours.

PDL drug coverage information can be found at <http://www.VirginiaMedicaidPharmacyServices.com>. The following "routine" PDL criteria guidelines will be applied to all non-preferred drugs.

1. Is there any reason the member cannot be changed to a preferred drug within the same class? Acceptable reasons include:
 - Allergy to preferred drug.
 - Contraindication to or drug-to-drug interaction with preferred drug.
 - History of unacceptable/toxic side effects to preferred drug.
 - Member's condition is clinically stable; changing to a preferred drug might cause deterioration of the member's condition.
2. The requested drug may be approved if both of the following are true:
 - There has been a therapeutic failure of at least **two** preferred drugs **within the same class as appropriate for diagnosis unless otherwise noted in the clinical criteria**. A therapeutic failure of only one preferred drug is required when there is only one preferred drug within a therapeutic class.
 - The requested drug's corresponding generic (if a generic is available **and** covered by the State) has been attempted and failed or is contraindicated.

All changes from the last post will be highlighted in yellow.

Drugs highlighted in blue indicate where brand is preferred over generic.

LEGEND

AGE = age edit

CE = clinical edit

ST = step edit

QL = quantity limit

cap = capsule

cr = cream

ER = extended release

inj = injection

IR = immediate release

ODT = oral disintegrating tab

oint = ointment

soln = solution

supp = suppository

susp = suspension

tab = tab

♦ = Clinical Utilization Edit

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Preferred Agents		Non-Preferred Agents	SA Criteria
Analgesics			
*♦ Opioids – Long Acting (LAO)			
Preferred (Sch III-VI)	Non-Preferred		*All Long-Acting Opioids (preferred and non-preferred) require submission of a Clinical SA. Refer to combined short/long-acting opioid SA form (Short & Long Acting Opioid SA Form) <u>LENGTH OF AUTHORIZATIONS</u> - Up to 3 months for (includes HIV/AIDS, Chronic back pain, Arthritis, Fibromyalgia, Diabetic neuropathy, Postherpetic Neuralgia). - Up to 6 months for chronic pain (includes Cancer pain, Sickle cell disease, Palliative care, End-of-Life Care, Hospice).
Butrans® (buprenorphine) Transdermal Patch	<i>Belbuca (buprenorphine buccal film)</i> <i>buprenorphine (gen Butrans®)</i> <i>buprenorphine (gen Belbuca)</i> <i>ConZip® (tramadol ER)</i> <i>tramadol ER</i> <i>Ultram ER® (tramadol ER)</i>		
Preferred (Sch II)	Non-Preferred		
fentanyl 12, 25, 50, 75 & 100 mcg patches morphine sulfate ER tab	<i>fentanyl 37.5 mcg, 62.5 mcg, and 87.5 mcg patches</i> <i>hydromorphone ER</i> <i>Hysingla ER™</i> <i>morphine ER cap (generic Avinza®)</i> <i>morphine ER cap (generic Kadian®)</i> <i>MS Contin®</i> <i>oxycodone-long acting</i> <i>OxyContin®</i> <i>oxymorphone ER</i>		Daily dose limits have been established for all LAO. Quantity limits can be found at: Daily Dose Limits for Short & Long Acting Opioids
*Methadone Drugs			
	<i>Methadose® oral soln & tab</i> <i>methadone oral soln & tab</i>		*Methadone requires the completion of the Clinical SA form (Methadone SA Form) unless prescribed for neonatal abstinence syndrome for an infant under the age of one.
*♦ Opioids – Short Acting			
Short-Acting Opioids			<u>LENGTH OF AUTHORIZATIONS:</u>
codeine/APAP	<i>codeine tab/soln</i>		Up to 3 months for (includes HIV/AIDS, Chronic back pain, Arthritis, Fibromyalgia, Diabetic neuropathy, Postherpetic Neuralgia).
hydrocodone/APAP	<i>butalbital comp with codeine</i>		

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hydrocodone/ibuprofen hydromorphone morphine IR oxycodone IR oxycodone/APAP tramadol HCl 50mg tramadol HCl/APAP		butalbital/caffeine/APAP w/codeine butorphanol tartrate nasal dihydrocodeine/APAP/caffeine hydromorphone liq/supp meperidine tab morphine supp oxycodone/APAP(gen PrimLev™) oxycodone conc oxycodone oral syringe oxycodone/ASA oxycodone/ibuprofen oxymorphone HCl pentazocine/naloxone RoxyBond™ Seglentsis® tapentadol tramadol 25mg, 75mg, 100 Mg tramadol soln (AG for Qdolo®) Xyvona		Up to 6 months for chronic pain (includes Cancer pain, Sickle cell disease, Palliative care, End-of-Life Care, Hospice). *All Short-Acting Opioids (preferred and non-preferred) require the submission of a Clinical SA if prescribed for > 7 days or if more than two 7-day supply prescriptions within 60 days. Refer to combined short/long-acting opioid SA form (Short & Long Acting Opioid SA Form)			
♦ Opioid Dependency		CLOSED CLASS					
*buprenorphine SL buprenorphine/naloxone tab SL Brixadi™ Suboxone® film Sublocade™ SQ Kloxxado™ Spray naloxone syringe & vial naloxone nasal spray		*buprenorphine/naloxone film SL *Zubsolv™ **Lucemyra® **lofexidine		*Buprenorphine Containing Drugs require submission of Clinical SA. Refer (Oral Buprenorphine SA Form) **Lucemyra/lofexidine (PDL criteria do not apply) -Member is 18 years of age or older AND -Medication used for the mitigation of opioid withdrawal symptoms to facilitate abrupt opioid discontinuation Limitation: No more than one (14 day) treatment course every 6 months Quantity Limits <table><tr><td>buprenorphine SL tab 2mg</td><td>3/day</td></tr></table>		buprenorphine SL tab 2mg	3/day
buprenorphine SL tab 2mg	3/day						

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naloxone nasal spray OTC Naloxone Carpuject naltrexone tab Narcan® Nasal Spray Rx/OTC Opvee® Rextovy™ Nasal Spray Vivitrol® Zurnai™			buprenorphine SL tab 8mg	2/day	
			buprenorphine/naloxone SL tab 2–0.5mg	3/day	
			buprenorphine/naloxone SL tab 8–2mg	3/day	
			buprenorphine/naloxone SL film 2–0.5mg	3/day	
			buprenorphine/naloxone SL film 4–1mg	1/day	
			buprenorphine/naloxone SL film 8–2mg	3/day	
			Suboxone® SL film 2–0.5mg	3/day	
			Suboxone® SL film 4–1mg	1/day	
			Suboxone® SL film 8–2mg	3/day	
			Suboxone® SL film 12–3mg	2/day	
			Zubsolv™ SL tab 0.7–0.18 mg	2/day	
			Zubsolv™ SL tab 1.4–0.36mg	2/day	
			Zubsolv™ SL tab 2.9–0.71mg	2/day	
			Zubsolv™ SL tab 5.7–1.4mg	2/day	
			Zubsolv™ SL tab 8.6–2.1mg	2/day	
			Zubsolv™ SL tab 11.4–2.9mg	2/day	
Non-Steroidal Anti-Inflammatory Drugs (NSAIDs)					
Oral NSAIDs		LENGTH OF AUTHORIZATIONS: 1 year			
Children’s Motrin® susp (OTC) diclofenac sodium ibuprofen cap ibuprofen tab 100mg, 200mg, 400mg, 600mg, 800mg (OTC & Rx) Infant’s ibuprofen drops meloxicam tab naproxen tab naproxen sodium (OTC) naproxen EC (Rx) sulindac	Anaprox® IR & DS® Advil® Aleve® Arthrotec® Cataflam® *Celebrex® & *celecoxib Coxanto diclofenac potassium diclofenac potassium gen Zipsor™ diclofenac sodium ER diclofenac sodium/misoprostol diflunisal etodolac IR & SR fenoprofen flurbiprofen	Routine PDL edits plus *Step edit required for Celebrex®, celecoxib and Vyscoxa™ - History of a trial of a minimum of two (2) different non-COX2 NSAIDs within the past year; OR - Concurrent use of anticoagulants (i.e., warfarin, heparin, etc.), methotrexate, oral corticosteroids; OR - History of previous GI bleed or conditions associated with GI toxicity risk factors (i.e., PUD, GERD, etc), OR - Specific indication for Celebrex®, celecoxib and Vyscoxa™ for which preferred drugs are not indicated.			



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		ibuprofen 300mg tab ibuprofen tab chew OTC ibuprofen-famotidine (gen Duexis®) Indocin® supp indomethacin IR, SR & rectal ketoprofen IR & ER ketorolac ketorolac tromethamine (generic Sprix® nasal spray) meclofenamate mefenamic meloxicam susp meloxicam (generic Vivlodex™) Mobic® susp Motrin® nabumetone Nalfon® Naprelan® Naprosyn® naproxen CR (generic Naprelan®) naproxen-esomeprazole mag (generic Vimovo®)DR naproxen susp Orudis® oxaprozin piroxicam Sprix® nasal spray tolmetin sodium *Vyscoxa™ Zybic™	
Topical NSAIDs			LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus
	diclofenac sodium 1% gel	*diclofenac 1.5% topical soln *diclofenac 2% topical soln gen	

Preferred Agents		Non-Preferred Agents	SA Criteria
diclofenac sodium 1% gel (OTC)		*diclofenac epolamine 1.3% patch(QL) *Diclogen Kit *Lixofen 1.5% Kit	*Clinical Criteria for Non-Preferred Topical NSAIDs: - Approval is based on member failing the oral generic of the desired drug and at least one other preferred NSAID (to equal a total of at least two preferred). - diclofenac 1.5% and 2% topical sol, Lixofen 1.5% Kit and Diclogen Kit can only be approved for the FDA approved indication of osteoarthritis of the knee. <i>Quantity limit for diclofenac epolamine 1.3% patch = 30 patches per RX</i>
Antibiotic-Anti-Infective			
*♦ Antibiotics, Inhaled (QL, AGE) CLOSED CLASS			
Bethkis[®] Kitabis[™] Pak **Tobi Podhaler[®] (SE) tobramycin inhalation neb soln (generic Tobi[®] inh)		***Arikayce[®] (amikacin liposome) Cayston[®] Tobi[®] inh neb soln tobramycin (generic Bethkis[®]) tobramycin Pak (generic Kitabis[™] Pak)	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus **Tobi Podhaler[®] - Requires a clinical reason as to why one of the preferred tobramycin inhalation nebulizer solutions cannot be used ***Clinical Criteria for Arikayce[®] Duration of Approval: 12 months Initial Approval Criteria - Member is ≥ 18 years of age; AND - Diagnosis of Mycobacterium avium complex (MAC) lung disease as determined by the following: - chest radiography or high-resolution computed tomography (HRCT) scan; AND - at least 2 positive sputum cultures; AND - other conditions such as tuberculosis and lung malignancy have been ruled out; AND - Member has failed a multi-drug regimen with a macrolide (clarithromycin or azithromycin), rifampin, and ethambutol. (Failure is defined as continual positive sputum cultures for MAC while adhering to a multi-drug treatment regimen for a minimum duration of 6 months); AND - Member has documented failure or intolerance to aerosolized administration of amikacin solution for injection, including pretreatment with a bronchodilator; AND - Arikayce will be prescribed in conjunction with a multi-drug antimycobacterial regimen

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			*Minimum age for use is 6 years for all tobramycin inhalation nebulizer solution (Bethkis[®], Kitabis[™] Pak, Tobi[®] and Tobi Podhaler[®]) and 7 years for Cayston[®]. Quantity Limitations: Arikayce[®]= 590 mg/8.4 Ml (28 vials)/28 days Each carton contains a 28-day supply of medication (28 vials) Bethkis[®] = 224Ml (56 amps)/28 days Cayston[®] = 84 Ml (56 amps)/28 days Kitabis[™] Pak = 280Ml (56 amps)/28 days Tobi Podhaler[®] = 224 capsule/28 day Tobi[®] inhalation neb = 280Ml (56 amps)/28 days tobramycin nebs = 280Ml (56 amps)/28 days
	Antifungals, Oral		
	fluconazole tab/susp griseofulvin susp nystatin tab/susp terbinafine tab	<i>Brexafemme[®]</i> <i>clotrimazole (mucous mem)</i> <i>Cresemba[®]</i> <i>Diflucan[®] susp</i> <i>flucytosine</i> <i>griseofulvin tab</i> <i>griseofulvin ultramicrosize</i> <i>itraconazole</i> <i>itraconazole soln (generic for</i> <i>Sporanox[®] soln)</i> <i>ketoconazole</i> <i>Lamisil[®] tab/granules</i> <i>Noxafil[®]</i> <i>Noxafil[®] Powdermix</i> <i>posaconazole tab (generic for</i> <i>Noxafil)</i> <i>Sporanox[®] cap</i> <i>Tolsura[™]</i> <i>Vfend[®] susp</i>	<u>LENGTH OF AUTHORIZATIONS:</u> Duration of the prescription (up to 12 months) Routine PDL edits plus <u>Clinical Criteria for all Non-Preferred oral Antifungals.</u> Requires the submission of a Clinical SA. Refer to Antifungal Oral SA Form <u>*Clinical Criteria for Vivjoa:</u> Duration of approval: Date of service • Member is 10 years of age or older; AND • Documentation member has diagnosis of recurrent vulvovaginal candidiasis with ≥3 episodes of vulvovaginal candidiasis (VVC) in a 12-month period; AND • Member is a biological female who is postmenopausal or has another reason for permanent infertility (e.g., tubal ligation, hysterectomy, salpingo-oophorectomy); AND

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		*Vivjoa [®] voriconazole tab & powder for susp	Member has tried and failed or has a contraindication or intolerance to maintenance antifungal therapy with oral fluconazole. Quantity limit: 18 tablets per treatment course
Cephalosporins, Oral			
Second Generation Cephalosporins			LENGTH OF AUTHORIZATIONS: Date of service only; no refills.
cefaclor capsule cefprozil tab/susp cefuroxime tab	cefaclor ER cefaclor susp		Routine PDL edits plus
Third Generation Cephalosporins			Clinical Criteria for <u>Non-Preferred Cephalosporins</u>
cefdinir cap/susp	ceftibuten cefditoren pivoxil cefixime susp, tab, cap cefpodoxime proxetil cap/susp		- Infection caused by an organism resistant to preferred drugs, OR - A therapeutic failure to no less than a three-day trial of <u>one preferred cephalosporin</u>; OR - The member is completing a course of therapy with a non-preferred drug initiated in the hospital.
Macrolides, Oral			
Macrolides & Ketolides			LENGTH OF AUTHORIZATIONS: Date of service only; no refills
azithromycin pack/susp/tab clarithromycin tab/susp E.E.S. [®] 200 susp erythromycin base tab DR erythromycin ethylsuccinate 200mg susp erythromycin stearate	Biaxin [®] tab clarithromycin ER Eryped [®] 200 susp Eryped [®] 400 susp Ery-tab [®] E.E.S. [®] 400 tab Erythrocin [®] Stearate erythromycin base cap DR erythromycin base tab erythromycin ethylsuccinate 400mg tab(Generic E.E.S. [®] 400) PCE [®] Zithromax [®] pac/tab/susp		Routine PDL edits plus Clinical Criteria for <u>Non-Preferred Macrolides and Ketolides</u> - Infection caused by an organism resistant to preferred drugs, OR - A therapeutic failure to no less than a <u>three-day trial of one preferred drug within the same class</u>; OR - The member is completing a course of therapy with a non-preferred drug which was initiated in the hospital.

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Otic					
ciprofloxacin/dexamethasone (generic Ciprodex) ofloxacin neomycin/polymyxin/hc soln/sus		Cipro HC® ciprofloxacin 0.2% otic soln ciprofloxacin and fluocinolone acetonide (generic Otovel) ciprofloxacin-hydrocortisone (generic Cipro HC) Cortisporin TC	<u>LENGTH OF AUTHORIZATIONS:</u> Date of service only; no refills Routine PDL edits		
Quinolones, Oral					
Second Generation Quinolones			<u>LENGTH OF AUTHORIZATIONS:</u> Date of service only; no refills		
ciprofloxacin susp/tab		Cipro® IR, susp ciprofloxacin ER ofloxacin	Routine PDL edits plus: <u>Clinical Criteria for Non-Preferred Quinolones</u> • Infection caused by an organism resistant to preferred drugs; OR • A therapeutic failure to no less than a <u>three-day trial of one preferred quinolone</u>; OR • The member is completing a course of therapy with a non-preferred drug initiated in the hospital.		
Third Generation Quinolones					
levofloxacin tab		Baxdela™ tab levofloxacin susp moxifloxacin			
Topical Antibiotics					
mupirocin ointment		Centany® Centany AT® Kit mupirocin cream	<u>LENGTH OF AUTHORIZATIONS:</u> Date of service only; no refills Routine PDL edits		
Vaginal Antibiotics					
Cleocin® Ovules Clindesse® cr metronidazole gel		Cleocin® cr clindamycin cr Nuversa™ Vandazole™ gel Xaciato™ gel	<u>LENGTH OF AUTHORIZATIONS:</u> Date of Service Routine PDL edits		

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Preferred Agents		Non-Preferred Agents		SA Criteria	
Antivirals					
Hepatitis B Agents		CLOSED CLASS			
entecavir (generic for Baraclude®) tab		adefovir dipivoxil (generic for Hepsera®) tab Baraclude® (entecavir) soln Baraclude® (entecavir) tab Epivir HBV® (lamivudine) soln lamivudine HBV (generic for Epivir HBV®) tab *Vemlidy® (tenofovir alafenamide)		LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus *Vemlidy® - Member has a diagnosis of chronic hepatitis B infection; AND - Member has tried and failed or is intolerant to tenofovir disoproxil fumarate (generic Viread)	
*Hepatitis C Agents		CLOSED CLASS			
Interferon		LENGTH OF AUTHORIZATIONS:			
	Pegasys® syringe/vial	*Non-preferred agents require the submission of a Clinical SA. Preferred agents do not require submission of any SA. Refer to Hepatitis C Antivirals Non-Preferred SA Form			
*Nucleotide Analog NS5A & NS5B Polymerase Inhibitors & Combinations		* Hepatitis C Therapies may filled:			
sofosbuvir /velpatasvir (generic Epclusa®)		Epclusa® Epclusa® Pellet packs Sovaldi® Vosevi™		1. Monthly - Mavyret™ (28 days supply x 2 months) - Mavyret™ Pellet packs (28 days supply x 2 months) - sofosbuvir /velpatasvir (generic Epclusa®) (28 days supply x 3 months)	
*NS5A, NS3/4A Inhibitor Combinations		OR			
Mavyret™ Mavyret™ Pellet packs		Zepatier®		2. Full duration of the regimen: - Mavyret™ (56 days supply) - Mavyret™ Pellet packs (56 days supply) - sofosbuvir/velpatasvir (generic Epclusa®) (84 day supply)	
*NS5B & Protease Inhibitor Combinations					
	Harvoni® ledipasvir/sofosbuvir (generic Harvoni®)				
Herpes Oral					
acyclovir cap/tab/susp famciclovir		Valtrex®		LENGTH OF AUTHORIZATIONS: 1 year	



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valacyclovir		Routine PDL edits
Herpes Topical		
acyclovir oint docosanol	acyclovir cr (generic Zovirax cr) Denavir® penciclovir	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
HIV/AIDS CLOSED CLASS		
abacavir tab/soln abacavir-lamivudine tab abacavir-lamivudine-zidov tab Apretude atazanavir sulfate cap Biktarvy® tab Cabenuva Cimduo® tab Complera® tab darunavir Delstrigo™ tab Descovy® tab Dovato® tab Edurant® tab efavirenz tab efavir-emtri-tenof tab emtricitabine cap emtricitabine-tenofv Emtriva™ cap/soln Epivir® soln etravirine tab Evotaz® tab fosamprenavir tab/susp Genvoya® tab Intelence® tab	Aptivus® cap efavirenz-lamiv-tenof tab Epivir® tab emtricit-rilp-tenof Kaletra® tab Norvir® tab Prezista® tab/susp Retrovir® syrup Reyataz® cap rilpivirine tab Trogarzo® (intraven) Viracept® tab Viread® tab	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus Quantity limits see list here HIV Quantity Limits



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Isentress® chew tab/ tab/powder pack Juluca® tab Kaletra® soln lamivudine tab/soln lamivudine-zidovudine tab lopinavir-ritonavir tab Maraviroc nevirapine IR tab/ ER tab/susp Norvir® powder pack Odefsey® tab Pifeltro™ tab Prezcobix® tab Retrovir® cap/vial Reyataz® powder pack rilpivirine ER vial ritonavir tab/soln Rukobia™ ER tab Selzentry® tab/soln Stribild® tab Sunlenca® tab/vial Symfi® and Symfi Lo® tab Symtuza® tab tenofovir disoproxil fum tab Tivicay® tab/tab for susp Triumeq® tab Triumeq® PD Truvada® tab Tybost® tab Viramune® tab Viread® powder Vocabria Yeztugo® Ziagen® soln		



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zidovudine tab/cap/syrup		
Influenza		
oseltamivir susp/ cap	<i>Flumadine[®] tab</i> <i>rimantadine</i> <i>Relenza Disk[®]</i> <i>Tamiflu[®] susp/cap</i> <i>XofluzaTM</i>	LENGTH OF AUTHORIZATIONS: Date of service only Routine PDL edits
COVID-19 CLOSED CLASS		
Paxlovid		LENGTH OF AUTHORIZATIONS: Date of service only
Blood Modifiers		
Bile Salts		
ursodiol cap, tab	<i>*BylvayTM</i> <i>Chenodal[®]</i> <i>Cholbam[®]</i> <i>Ctexli</i> <i>Iqirvo[®]</i> <i>Livdelzi[®]</i> <i>**LivmarliTM</i> <i>Reltone[®]</i> <i>Urso[®] & Urso[®] Forte tab</i>	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits **LivmarliTM (PDL criteria do not apply) - Must have a confirmed diagnosis of cholestatic pruritus in: - Members 3 months of age and older with Alagille syndrome (ALGS) - Members 12 months of age and older with progressive familial intrahepatic cholestasis (PFIC). * BylvayTM (PDL criteria do not apply) - Must have a confirmed diagnosis of cholestatic pruritus in: - Members 12 months of age and older with Alagille syndrome (ALGS) - Members 3 months of age and older with progressive familial intrahepatic cholestasis (PFIC)
Colony Stimulating Factors CLOSED CLASS		



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<i>Preferred Agents</i>		<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
Fulphila Neupogen Disp Syrin Neupogen Vial		<i>Fylnetra</i> <i>Granix Syringe</i> <i>Granix Vial</i> <i>Leukine</i> <i>Neulasta Kit</i> <i>Neulasta Syringe</i> <i>Nivestym Syringe</i> <i>Nivestym Vial</i> <i>Nypozi</i> <i>Nyvepria</i> <i>Releuko Syringe</i> <i>Releuko Vial</i> <i>Rolvedon Syringe</i> <i>Ryzneuta Syringe</i> <i>Stimufend Syringe</i> <i>Udenyca</i> <i>Udenyca Autoinjector</i> <i>Udenyca Onbody</i> <i>Zarxio</i> <i>Ziextenzo Syringe</i>	<u>LENGTH OF AUTHORIZATIONS:</u> Routine PDL edits plus ♦All Non-Preferred Colony Stimulating Factors require submission of Clinical SA. Refer to: Non-Preferred Colony Stimulating Factors SA Form
Hemophilia Treatment CLOSED CLASS			
Factor VIII Products			<u>LENGTH OF AUTHORIZATIONS:</u> N/A All products are preferred without EDITS
Advate® Adynovate® Afstyla® Alphanate® Altuviiio™ Eloctate® Esperoct® Hemofil-M® Humate-P® Jivi® Koate-DVI®			



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Kogenate FS[®] Kovaltry[®] Novoeight[®] Nuwiq[®] Obizur[®] Recombinate[®] Xyntha[®] Kit & Solofuse Syr		
Factor IX Products		
AlphaNine SD[®] Alprolix[®] BeneFIX[®] Idelvion[®] Ixinity[®] Profilnine SD[®] Rebiny[®] Rixubis[®]		
Factor VIIa & Activated Prothrombin Complex Concentrate		
Feiba NF[®] NovoSeven RT[®] Sevenfact[®]		
Factor IXa and Factor X Directed Antibody		
Hemlibra[®]		
Factor X and Factor XIII Products		
Coagadex[®] Corifact[®] Kit Tretten[®]		
Antithrombin Lowering Agents		
Qfitlia		
Prothrombin Complex Concentrate		

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Preferred Agents		Non-Preferred Agents	SA Criteria	
Balfaxar®				
Tissue Factor Pathway Inhibitor Antagonist				
Alhemo®				
Hympavzi™				
Von Willebrand				
Vonvendi®				
Wilate®				
Phosphate Binders				
calcium acetate 667mg cap calcium acetate 668 mg sevelamer carbonate tab	Auryxia™ calcium acetate 667mg tab Calphron Eliphos Fosrenol® chewable tab lanthanum carbonate chewable tab Phoslyra® powder, tab Renvela® sevelamer carbonate powder packet sevelamer HCL (gen Renagel) Velphoro® chewable tab Xphozah™			LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
Sickle Cell CLOSED CLASS				
Droxia Endari **Siklos *Xromi	Adakveo IV l-glutamine(generic Endari)	*Non-preferred agents, **Siklos® (if 18 years of age and older), and *Xromi (if 2 years of age and older) require the submission of a Clinical SA. Preferred agents do not require submission of any SA. Refer to Sickle Cell SA Form **Clinical Criteria for Siklos (hydroxyurea): Member must be 2-17 years old		

Preferred Agents		Non-Preferred Agents	SA Criteria
			*Clinical Criteria for Xromi (hydroxyurea): Member must be < 2 years old
Bone Resorption Suppression and Related Agents			
Bisphosphonates			
alendronate tab ibandronate	<i>Actonel[®]</i> <i>alendronate soln</i> <i>Atelvia DR[®]</i> <i>BinostoTM</i> <i>etidronate</i> <i>Fosamax[®] tab & Fosamax[®] plusD</i> <i>risedronate DR</i>	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits	
Calcitonins			
calcitonin-salmon nasal		LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits	
Others			
raloxifene	<i>*Bonsity[®]</i> <i>Evista[®]</i> <i>*Forteo[®]</i> <i>*teriparatide</i> <i>*TymlosTM</i>	LENGTH OF AUTHORIZATIONS: Initial approval will be for 1 year Routine PDL edits for Evista[®] *♦ Clinical SA must be completed for (Bonsity, Forteo, teriparatide or Tymlos SA Form)	
♦ RANK Ligand Inhibitors			
Bildyos[®] Bilprevda[®]	<i>Bomyntra[®]</i> <i>Conexence[®]</i> <i>EnobyTM</i> <i>Jubbonti[®]</i> <i>Osenvelt[®]</i> <i>Prolia[®]</i> <i>Stoboclo[®]</i> <i>Wyost[®]</i> <i>Xgeva[®]</i> <i>XtrenboTM</i>	*♦ ALL RANK Ligand Inhibitors require the submission of a Clinical SA. Refer to denosumab SA Form	

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Preferred Agents		Non-Preferred Agents		SA Criteria	
Cardiac					
Anticoagulants					
CLOSED CLASS					
Low Molecular Weight Heparin includes FactorXA Inhibitor		LENGTH OF AUTHORIZATIONS: 1 year			
enoxaparin (generic Lovenox®)		Arixtra® fondaparinux Fragmin® syringe & vial Lovenox®		Routine PDL edits	
Oral Anticoagulants		*Clinicial Criteria for Savaysa™			
Eliquis™ tab, sprinkle cap, susp Eliquis™ Dose Pack Jantoven® Pradaxa® warfarin Xarelto® tab, susp & Starter Pack		dabigatran Pradaxa® Pellet Pack rivaroxaban *Savaysa™		• Diagnosis of: Non-valvular Atrial Fibrillation, OR • Deep vein thrombosis, OR • Pulmonary embolism; AND • Documentation that CrCl is NOT ≥ 95mL/min calculated by Cockcroft-Gault equation	
Antihypertensive Agents					
ACE Inhibitors		LENGTH OF AUTHORIZATIONS: 1 year			
benazepril enalapril lisinopril ramipril		Altace® captopril Epaned™ soln enalapril soln fosinopril Lotensin® moexipril perindopril Qbrelis™ quinapril trandolapril Zestril®		Routine PDL edits	

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Preferred Agents		Non-Preferred Agents	SA Criteria
ACE Inhibitors + Calcium Channel Blocker Combinations			
amlodipine/benazepril		Lotrel® trandolapril-verapamil ER	
ACE Inhibitors + Diuretic Combinations			
benazepril/HCTZ lisinopril/HCTZ enalapril/HCTZ		captopril/HCTZ fosinopril/HCTZ Lotensin HCT® moexipril/HCTZ quinapril /HCTZ quinapril-HCTZ (gen Accuretic) Zestoretic®	
Angiotensin Receptor Blockers			LENGTH OF AUTHORIZATIONS: 1 year
*Entresto™ (QL) irbesartan losartan olmesartan telmisartan valsartan		Arbli™ Atacand® Avapro® Benicar® candesartan Cozaar® Diovan® Edarbi® **Entresto Sprinkle™ eprosartan mesylate Micardis® sacubitril/valsartan (QL) valsartan solution	Routine PDL edits plus Quantity Limit = 2 per day for Entresto™ and sacubitril/valsartan generics **Entresto Sprinkle capsule: The member is intolerant to solid oral dosage forms
Angiotensin Receptor Blockers + Calcium Channel Blocker Combinations			
amlodipine/olmesartan amlodipine/valsartan		Azor® amlodipine/olmesartan/HCTZ amlodipine/valsartan/HCTZ	

Preferred Agents		Non-Preferred Agents	SA Criteria
		Exforge® & Exforge® HCT Tribenzor®	
Angiotensin Receptor Blockers + Diuretic Combinations			
irbesartan/HCTZ losartan/HCTZ olmesartan/HCTZ telmisartan/HCTZ valsartan/HCTZ		Atacand HCT® Avalide® Benicar HCT® candesartan/HCTZ Diovan HCT® Edarbyclor® Hyzaar® Micardis HCT® Teveten HCT®	
Beta Blockers			*Clinical Criteria for Hemangeol™ • Diagnosis of proliferating infantile hemangioma requiring systemic therapy
atenolol bisoprolol carvedilol labetalol metoprolol tartrate metoprolol succinate nadolol nebivolol propranolol tab & ER/soln sotalol AF sotalol HCL		acebutolol Betapace® IR & AF® betaxolol Carvedilol ER *Hemangeol™ Inderal® XL Innopran® XL Kapspargo™ Sprinkle Lopressor® pindolol propranolol LA Sotylize™ Tenormin® timolol maleate Toprol XL®	
Beta Blockers + Diuretic Combinations			
atenolol/chlorthalidone bisoprolol/HCTZ		Lopressor HCT® metoprolol/HCTZ	



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Preferred Agents		Non-Preferred Agents	SA Criteria
		nadolol/bendroflumethiazide propranolol/HCTZ Tenoretic®	
Calcium Channel Blockers –Dihydropyridine			
amlodipine nifedipine nifedipine ER		felodipine ER isradipine Katerzia™oral suspension levamlodipine (generic Conjupri®) nisoldipine nicardipine Norliqva® Norvasc® Procardia XL® Sdamlo Sular®	
Calcium Channel Blockers- Non-Dihydropyridine			
Cartia XT® diltiazem IR, ER q12hr & 24hr verapamil tab IR & ER		diltiazem LA Matzim LA verapamil 360 cap verapamil ER cap	
Direct Renin Inhibitors (includes combination)			
		aliskiren 150 & 300mg (generic for Tekturna) Tekturna® telmisartan/amlodipine	

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Preferred Agents		Non-Preferred Agents	SA Criteria
Lipotropics			LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits Routine PDL edits plus ♦ Clinical criteria NexletoTM or NexlizetTM Initial Approval Criteria - Member is ≥ 18 years of age; AND - Member has diagnosis of heterozygous familial hypercholesterolemia (HeFH) or established atherosclerotic cardiovascular disease (ASCVD); AND - Member has failed to achieve a target LDL-C despite physician attestation that the member is adherent to maximally tolerated doses of statins prior to the lipid panel demonstrating suboptimal reduction; AND - Member can be classified into ONE of the following risk factor groups: - Extremely high risk ASCVD: (defined as extensive or active burden of ASCVD, or ASCVD with extremely high burden of adverse or poorly controlled risk cardio-metabolic risk factors including HeFH or severe hypercholesterolemia [SH] LDL-C > 220 mg/dl) with an LDL-C ≥ 70 mg/dL; OR - Very high risk ASCVD: (defined as less extensive ASCVD and poorly controlled cardiometabolic risk factors) with an LDL-C ≥ 100 mg/dL; OR - High risk ASCVD: (defined as either less extensive ASCVD and well-controlled risk factors or primary prevention HeFH or SH >220 mg/dl with poorly controlled risk factors) with LDL-C ≥ 130 mg/dL; AND - Therapy will be used in conjunction with maximally tolerated doses of a statin; AND - Therapy will not be used with concurrent doses of simvastatin > 20 mg or pravastatin > 40mg;
Bile Acid Sequestrants			
cholestyramine powder reg & light colestipol tab Prevalite [®]		Colestid [®] packet/tab colesevelam tab and Pkt (generic Welchol) colestipol HCl granules Questran [®] powder/powder Light Welchol [®] pack, tab	
Cholesterol Absorption Inhibitor (CAI) and /or ACL Inhibitor (adenosine triphosphate citrate lyase)			
ezetimibe		Nexleto [®] (bempedoic acid) Nexlizet [®] (bempedoic acid/ezetimibe) Zetia [®]	

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Preferred Agents		Non-Preferred Agents	SA Criteria
			Renewal Criteria - Laboratory analyses demonstrate a reduction in LDL-C when compared to the baseline values (prior to initiating bempedoic acid or bempedoic acid/ezetimibe); AND - Member has shown continued adherence to maximally tolerated statin dosage
Fibric Acid Derivatives			
fenofibrate (generic Tricor® 48mg 145mg) gemfibrozil		<i>fenofibrate</i> (generics for Antara®, Fenoglide® & Lipofen®) <i>fenofibrate</i> (generics for Triglide®) <i>fenofibric acid</i> <i>Fibricor</i> ® <i>Lipofen</i> ® <i>Lopid</i> ® <i>Tricor</i> ®	
HMG CoA Reductase Inhibitors and Combo (High Potency Statins)			
atorvastatin rosuvastatin simvastatin		<i>Atorvaliq</i> ® <i>amlodipine/atorvastatin</i> <i>Caduet</i> ® <i>Crestor</i> ® <i>Lipitor</i> ® <i>Livalo</i> ® <i>pitavastatin calcium</i> (generic Livalo®) <i>rosuvastatin-ezetimibe</i> <i>simvastatin/ezetimibe</i> <i>Vytorin</i> ® <i>Zocor</i> ® <i>Zypitamag</i> ™	
HMG CoA Reductase Inhibitors and Combinations (Statins)			
lovastatin pravastatin		<i>fluvastatin</i> <i>Lescol XL</i> ®	

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Preferred Agents		Non-Preferred Agents	SA Criteria
Microsomal Triglyceride Transfer Protein Inhibitor			
		*Juxtapid™	*♦ Clinical Criteria for Juxtapid™. Refer to Juxtapid™ SA Fax Form
Niacin Derivatives			
niacin ER			
Omega 3 Fatty Acid Agent			
icosapent ethyl (gen Vascepa®)		***Lovaza®	***♦ <u>Clinical Criteria for Lovaza® and omega-3 acid ethyl esters</u>
***omega-3 acid ethyl esters			- Step edit requires trial and failure of any other lipotropic; OR - Documented high triglycerides of ≥ 500 mg/dL.
Omega-3 OTC			
*Proprotein Convertase Subtilisin Kexin Type 9 (PCSK9) Inhibitors			<u>LENGTH OF AUTHORIZATIONS:</u> Three months for initial approval; six months for renewal
		Leqvio® Praluent® Repatha®	*♦ ALL PCSK9 Inhibitors require the submission of a Clinical SA. Refer to Lipotropics, Other SA Form
Platelet Inhibitors			
Brilinta®		*ASA/dipyridamole	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year
clopidogrel		**ASA/omeprazole (generic)	Routine PDL edits plus
dipyridamole		Yosprala®)	<u>Clinical Criteria for Select Non-Preferred Platelet Inhibitors</u>
prasugrel (generic Effient®)		Effient®	* <u>ASA/dipyridamole</u>
		Plavix®	Aspirin and dipyridamole are covered as separate drugs without SA; clinical reason as to why the individual drugs cannot be used separately.
		ticagrelor	
*Pulmonary Arterial Hypertension Agents			
Activin Signaling Inhibitor			<u>LENGTH OF AUTHORIZATIONS:</u> 1 year
		Winrevair™	Routine PDL edits for all Pulmonary Arterial Hypertension Agents
Inhaled Prostacyclin Analogues			
		Tyvaso®	
		Tyvaso DPI™ (treprostinil)	

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Preferred Agents		Non-Preferred Agents	SA Criteria
		<i>Yutrepia™</i>	Routine PDL edits plus * ♦ Clinical Criteria for all preferred and non-preferred PDE-5 and PDE-5 inhibitor combinations • Diagnosis of pulmonary hypertension in members >18 years is required (≥1 years oral Revatio® only); AND • The prescriber must be a pulmonary specialist or cardiologist; AND • Must have a rationale for not taking the sildenafil tab to receive a SA for injectable Revatio®
Oral Endothelin Receptor Antagonist			
ambrisentan (generic Letairis) 5 & 10mg Tracleer® tab		<i>bosentan (generic Tracleer®)</i> <i>bosentan susp</i> <i>Letairis® 5 & 10 mg</i> <i>Opsumit®</i> <i>Tracleer® susp</i>	
*Phosphodiesterase 5 Inhibitors (PDE-5)			
Alyq (tadalafil) sildenafil tab/susp tadalafil(generic Adcirca®)		<i>Adcirca®</i> <i>Liqrev®</i> <i>Revatio® tab/inj</i>	
*Phosphodiesterase 5 Inhibitors (PDE-5) Combinations			
		<i>Opsynvi</i>	
Prostacyclin Vasodilator and Receptor Agonist			
		<i>Orenitram™</i> <i>Orenitram™ titration kit</i> <i>Uptravi®</i>	
Soluble Guanylate Cyclase Stimulators			
		<i>Adempas®</i>	
Central Nervous System			
Alzheimer's Agents			
Cholinesterase Inhibitors			LENGTH OF AUTHORIZATIONS: Length of prescription (up to 3 months)
donepezil OTD & tab rivastigmine (transderm)		<i>Adlarity®</i> <i>Aricept® ODT, tab</i> <i>Exelon® cap & transderm</i> <i>galantamine IR, ER tab/soln</i> <i>memantine/donepezil</i> <i>Namzaric® (memantine/donepezil)</i>	Routine PDL edits

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Preferred Agents		Non-Preferred Agents	SA Criteria
		Razadyne® IR, ER rivastigmine cap Zunveyl®	
NMDA Receptor Antagonist			
memantine tab		memantine Dose Pack, soln memantine ER	
Anticonvulsants		CLOSED CLASS	
Barbiturates			LENGTH OF AUTHORIZATIONS: 1 year
phenobarbital elixir/tab primidone Sezaby			Routine PDL
Benzodiazepines			
clobazam tab/susp clonazepam tab diazepam rectal & Device rectal Nayzilam® Valtoco® Nasal		clonazepam ODT clorazepate Klonopin Tab Onfi®susp/tab Sympazan™ (clobazam)	
Cannabidiol			
*Epidiolex® (cannabidiol)			*♦ Clinical Criteria for Epidiolex® (reviewed electronically, AutoPA) Duration of Approval: 1 year - Member must be ≥ 1 years of age; AND - Member has been diagnosed with: - Epilepsy and recurrent seizures including Lennox-Gastaut syndrome, - Dravet syndrome, or tuberous sclerosis complex
Carbamazepine Derivatives			
carbamazepine chewable tab/susp/tab		Aptiom® carbamazepine 200mg chewable tab carbamazepine ER	

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Preferred Agents		Non-Preferred Agents	SA Criteria
carbamazepine XR Carbatrol® Epitol oxcarbazepine tab Tegretol®XR Trileptal® susp		<i>carbamazepine XR (authorized generic only)</i> <i>eslicarbazepine acetate</i> <i>Equetro® cap</i> <i>Oxtellar™ XR</i> oxcarbazepine susp <i>oxcarbazepine ER (generic)</i> <i>Oxtellar XR</i> <i>Tegretol® susp/tab</i> <i>Trileptal® tab</i>	* ♦ Clinical Criteria for Fintepla® - Member is at least two years of age or older; AND - Member must have a diagnosis of Dravet syndrome or Lennox-Gastaut syndrome ** ♦ Clinical Criteria for Ztalmy: - Initial: - Member is two years of age or older; AND
Hydantoins			
Dilantin 30mg phenytoin cap/chew tab/ susp phenytoin ext cap		<i>Dilantin® cap & Infatab, susp</i> <i>Phenytek®</i>	
Succinimides			
ethosuximide cap/syrup		<i>Celontin®</i> <i>Zarontin® cap/syrup</i>	
Valproic Acid and Derivatives			
Depakote® sprinkle divalproex tab divalproex ER valproic acid cap, soln		<i>Depakote® ER</i> divalproex sprinkle	
Other Anticonvulsants			
lacosamide soln/tab (gen Vimpat®) lamotrigine tab lamotrigine chew tab lamotrigine XR levetiracetam soln/tab levetiracetam ER roweepra (generic levetiracetam)		<i>Banze® susp/tab</i> <i>brivaracetam tab</i> <i>Briviact® tab/soln</i> <i>Diacomit® cap/powder pack</i> <i>Elepsia®XR tab</i> <i>Eprontia™</i> <i>felbamate susp/tab</i> <i>Felbatol® tab</i> <i>* Fintepla®</i> <i>Fycompa® susp/tab</i>	

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Preferred Agents		Non-Preferred Agents	SA Criteria
subvenite tab (generic lamotrigine) topiramate tab topiramate 15, 25 mg sprinkle cap zonisamide cap		<i>Keppra® soln/tab</i> <i>Keppra® XR</i> <i>lacosamide solution unit dose</i> <i>Lamictal® XR & XR dose pk</i> <i>Lamictal® tab/dose pk</i> <i>Lamictal® ODT</i> <i>Lamictal® ODT dose pk</i> <i>lamotrigine tab dose pk</i> <i>lamotrigine ODT</i> <i>lamotrigine ODT dose pk</i> <i>levetiracetam (generic Spritam®)</i> <i>Motpoly XR™</i> <i>perampanel</i> <i>rufinamide susp/tab (generic Banzel®)</i> <i>Sabril® powder pack/tab</i> <i>Spritam®</i> <i>subvenite dose pk/susp</i> <i>tiagabine</i> <i>Topamax® tab/sprinkle</i> <i>topiramate ER cap</i> <i>topiramate soln</i> <i>topiramate 50mg sprinkle cap</i> <i>Trokendi™ XR</i> <i>vigabatrin tab/ powder pack (generic Sabril®)</i> <i>Vigafyde™</i> <i>Vimpat® soln/tab</i> <i>Xcopri®</i> <i>Zonisade™ Solution</i> <i>**Ztalmu®</i>	○ Documented diagnosis of cyclin-dependent kinase-like 5 deficiency disorder AND ○ Documentation that seizures have been inadequately controlled by a trial of at least 2 antiepileptic drugs (e.g., clobazam, valproate, lamotrigine, levetiracetam, topiramate, felbamate, vigabatrin) or member has labeled contraindications to other antiepileptic drugs ● Renewal: ○ Documentation of positive clinical response as demonstrated by low disease activity and/or improvements in the condition's signs and symptoms (e.g., reduced seizure activity, frequency, and/or duration) ● Prescriber Requirements: Prescribed by or in consultation with a neurologist, geneticist, or physician who specialized in treatment of epileptic disorders
Antidepressants			
Other		LENGTH OF AUTHORIZATIONS:	
bupropion IR, SR & XL	<i>Auvelity™</i>	1 year	

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Preferred Agents		Non-Preferred Agents	SA Criteria
desvenlafaxine ER tab (generic for Pristiq®) mirtazapine ODT/tab trazodone tab venlafaxine IR tab & ER cap vilazodone tab		<i>bupropion XL (gen Forfivo® XL)</i> <i>desvenlafaxine ER tab (gen Khedezla™)</i> <i>Effexor® XR</i> <i>Emsam® transdermal</i> <i>Exxua™</i> <i>Fetzima®</i> <i>Khedezla™</i> <i>Marplan®</i> <i>Nardil®</i> <i>nefazodone</i> <i>phenelzine</i> <i>Pristiq®</i> <i>Raldesy™</i> <i>Remeron® ODT/tab</i> <i>tranylcypromine sulfate</i> <i>Trintellix</i> <i>venlafaxine ER tab</i> <i>Viibryd® dose pk</i> <i>Viibryd® tab</i> <i>Wellbutrin® SR</i> <i>*Zurzuva™</i>	Routine PDL edits *♦ Zurzuva criteria (PDL criteria do not apply) - Initial Criteria: - Member is ≥ 18 years of age; AND - Member has a diagnosis of postpartum depression (PPD) based on Diagnostic and Statistical Manual of Mental Disorders (DSM) criteria for a major depressive episode (DSM-5); AND - Member is not currently pregnant and is using effective contraception LENGTH OF AUTHORIZATIONS: 14 days Renewal Criteria: Not applicable Quantity limit: 28 capsules per 14 days
SSRI		<i>Brisdelle®</i> <i>Celexa® tab</i> <i>citalopram HBR 30mg</i> <i>escitalopram soln</i> <i>fluoxetine DR cap/tab</i> <i>fluvoxamine ER</i> <i>Lexapro® tab</i> <i>paroxetine CR</i> <i>paroxetine susp</i> <i>Paxil® tab/susp & Paxil® CR</i> <i>Prozac® cap/weekly</i>	



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Preferred Agents	Non-Preferred Agents	SA Criteria
	sertraline cap Zoloft® conc/tab	
Antimigraine Agents		
sumatriptan succinate tab cartridge/vial/pen rizatriptan tab/MLT	almotriptan Alsuma® Amerge® Axert® Cambia® eletriptan (generic Relpax®) Elyxyb® Frova® frovatriptan (generic Frova®) Imitrex® cartridge/pen/tab/vial Maxalt® tab & MLT Migranow™ Kit naratriptan Onzetra™ Xsail™ Relpax® sumatriptan KITS Sumavel® Dosepro sumatriptan/nap (gen Treximet®) Symbravo® Tosymra® Treximet® Zembrace™ SymTouch™ zolmitriptan nasal Zomig® tab/nasal spray/ZMT	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edit
Antimigraine Agents, Others CLOSED CLASS Calcitonin Gene-related Peptide Antagonist (CGRP)		Routine PDL edits plus *♦ All CGRPs require the submission of a Clinical SA. Refer to fax form Antimigraine Agents, Others SA Form **Vyepti SA Form

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Preferred Agents	Non-Preferred Agents	SA Criteria
Preventive Treatment		Preferred Agents *step edit required
Aimovig™ Ajovy® Autoinjector & Syringe Ajovy Autoinjector 3-Pk Emgality™ Syringe Emgality™ Pen Nurtec™ ODT Qulipta™	Emgality™ 100MG Syringe **Vyepti®	Quantity limit for preventive treatment Aimovig™ = 70 mg/mL autoinjector = 1 mL or 140 mg/mL autoinjector = 1 mL per 30 days Ajovy® = 1 Injection: 225 mg/1.5 mL single-dose prefilled autoinjector per 30 days. Ajovy 3 pack = one 3 pack per 90 days Emgality™ = 120 mg/mL pen and syringe = 1 mL per 30 days 100 mg/1 mL syringe = 3 mL per 30 days (Episodic cluster headache treatment) Nurtec® ODT = 16 tabs per 30 days Qulipta™ = 1 tab per day
Acute treatment of migraine		Quantity limit for acute treatment
Nurtec™ ODT Ubrely®	Reyvow® Zavzpret™	Nurtec® ODT = 16 tabs per 30 days Reyvow® = 50mg or 100mg - 8 tabs per strength per 30 days Ubrely® = 50mg or 100mg can have up to 16 tabs per strength per 30 days
*Antipsychotics (AGE) CLOSED CLASS		
Atypical		LENGTH OF AUTHORIZATIONS: 1 year or 6 months for members < 18 yrs
aripiprazole tab clozapine tab lurasidone olanzapine ODT, tab, IM paliperidone ER quetiapine fumarate ER quetiapine tab risperidone ODT/soln/tab ****Vraylar™ ziprasidone cap, vial	Abilify® tab/IM inj aripiprazole ODT, soln asenapine (generic for Saphris®) Caplyta Capsule™ Clozari® clozapine ODT Cobenfy™ Cobenfy™ Starter pk Fanapt® tab & titration pk Fazaclo® Geodon® tab, IM Invega® Latuda® Lybalvi® **Nuplazid™ tab, cap (QL) **olanzapine/fluoxetine	Routine PDL edits plus *♦ ALL antipsychotics for children 0 to 17 years of age (preferred and non-preferred) require the submission of a Clinical SA. Refer to (Antipsychotics In Children Less Than 18 Years SA Form) **Clinical Criteria Nuplazid™ Indicated for the treatment of hallucinations and delusions associated with Parkinson's disease psychosis. Quantity Limit Nuplazid™ = 2 per day

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Preferred Agents		Non-Preferred Agents	SA Criteria
		<i>OpipzaTM</i> <i>quetiapine fumarate ER</i> <i>(authorized generic only)</i> <i>Rexulti[®] tab</i> <i>Risperdal[®] ODT/soln/tab</i> <i>Saphris[®]</i> <i>Secuado[®] Patch</i> <i>Seroquel[®] IR</i> <i>Seroquel[®] XR</i> <i>VersaclozTM</i>	****Clinical Criteria for Vraylar as adjunct treatment for Major Depressive Disorder (reviewed electronically, AutoPA) - Member must be ≥ 18 years of age; AND - Member must have a diagnosis of Major Depressive Disorder; AND - Member must be adherent to complementary therapy with an antidepressant (e.g. selective serotonin reuptake inhibitor (SSRI) or serotonin/norepinephrine reuptake inhibitor (SNRI)) (80% of fills in the last 90 days) - Length of Authorization: 1 year
Atypical, Long-Acting Injectable			LENGTH OF AUTHORIZATIONS: 1 year
Abilify Asimtufii [®] Abilify Maintena [®] Aristada [®] Aristada [®] Initio Erzofri [®] Invega Hafyera TM , Sustenna [®] &Trinza [®] Perseris TM Risperdal Consta [®] Uzedi TM		<i>RykindoTM</i> <i>Zyprexa[®] RelprevvTM</i>	Routine PDL edits
Typical			LENGTH OF AUTHORIZATIONS: 1 year
amitriptyline/perphenazine chlorpromazine fluphenazine elixir/soln/tab fluphenazine decantate haloperidol decantate haloperidol lactate conc haloperidol tab loxapine perphenazine trifluoperazine thiothixene		<i>Adasuve[®]</i> <i>Haldol decanoate (injection)</i> <i>pimozide</i> <i>molindone</i>	Routine PDL edits

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Preferred Agents	Non-Preferred Agents	SA Criteria
thioridazine		
Duchenne Muscular Dystrophy CLOSED CLASS		
**Emflaza	**Agamree Amondys-45 **deflazacort **Duvyzat Exondys-51 **Jaythari **Kymbee **Pyquvi Vilepso Vyondys-53	LENGTH OF AUTHORIZATIONS: 1 year **Clinical Criteria for oral Duchenne Muscular Dystrophy agents - Member is 2 years of age or older for Agamree®, Emflaza™ - Member is 5 years of age or older for Jaythari, Pyquvi, Kymbee and deflazacort - Member is 6 years of age or older for Duvyzat™ - Member has a diagnosis of Duchenne Muscular Dystrophy (DMD) - For all oral agents, the member has tried and failed or is intolerant to prednisone or prednisolone - For non-preferred agents, the member has tried and failed or is intolerant to Emflaza™ Routine PDL edits plus All Antisense Oligonucleotides require submission of Clinical SA. Refer to: Antisense Oligonucleotides SA Form
*Movement Disorders CLOSED CLASS		
Austedo® tab Austedo XR® tab Austedo XR® titration pack Ingrezza® cap Ingrezza® Sprinkle Ingrezza® Initiation Pack tetrabenazine tab (generic Xenazine®)	Xenazine® tab	LENGTH OF AUTHORIZATIONS: 1 year *Clinical Criteria for Movement Disorders - Diagnoses of Tardive Dyskinesia or Huntington's disease - Prescribed by or in consult with a neurologist or psychiatrist - Quantity limit 4 tabs/day Austedo® 1 tab per day Austedo XR® 42 tablets per 365 Austedo XR® titration pack 1 cap/day Ingrezza®/Ingrezza Sprinkle®

Virginia's Medicaid Preferred Drug List (PDL)/ Common Core Formulary

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Preferred Agents	Non-Preferred Agents	SA Criteria
		4 tabs/day Xenazine®
Neuropathic Pain and Select Agents		
capsaicin OTC topical duloxetine 20, 30 & 60 mg (generic for Cymbalta®) gabapentin cap/tab/soln lidocaine 5% patch pregabalin cap	<i>Drizalma™ Sprinkle</i> <i>duloxetine 40 mg (generic</i> <i>Irenka™)</i> <i>gabapentin ER</i> <i>Gralise™ (gabapentin, ER)</i> <i>Horizant™ (gabapentin</i> <i>enacarbil ER)</i> <i>Journavx™</i> <i>Lidoderm® patch</i> <i>Lyrica® / CR/ soln</i> <i>Neurontin® cap/tab/soln</i> <i>pregabalin soln, ER</i> <i>Qutenza Kit® (capsaicin)</i> <i>Savella™ & Savella™ Dose Pak</i> <i>Tonmya™</i> <i>Zilido™ 1.8% (lidocaine topical</i> <i>system)</i>	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus Journavx SA Form Criteria for Lyrica solution: – If diagnosis is epilepsy/seizures, member must have a problem swallowing tabs/capsules AND clinical reason why at least TWO preferred seizure medications cannot be used. – For other diagnoses, member must have a problem swallowing tabs/capsules AND routine PDL • If Lyrica CR is requested, routine PDL applies. Note – This is NOT indicated for epilepsy/seizures.
Non-Ergot Dopamine Receptor Agonist		
pramipexole ropinirole HCl	<i>Neupro®</i> <i>pramipexole ER</i> <i>ropinirole HCl ER</i>	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
Sedatives / Hypnotics		
temazepam 15 & 30 mg	<i>estazolam</i> <i>flurazepam</i> <i>Halcion®</i> <i>quazepam 15 mg</i> <i>Restoril®</i> <i>temazepam 7.5 mg & 22.5 mg</i> <i>triazolam</i>	LENGTH OF AUTHORIZATIONS: Length of the prescription (up to 3 months) Routine PDL edits

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Preferred Agents	Non-Preferred Agents	SA Criteria
Sedatives / Hypnotics (Non-Benzodiazepine)		
eszopiclone zaleplon zolpidem	Ambien [®] IR & CR Belsomra [®] Dayvigo [™] doxepin (generic for Silenor [®]) Eduar [™] *Hetlioz [®] **Hetlioz [™] LQ Intermezzo [®] Lunesta [®] Quviviq [™] Rozerem [®] Silenor [®] Sonata [®] tasimelteon zolpidem CR zolpidem (generic Intermezzo [®]) zolpidem tartrate capsule	LENGTH OF AUTHORIZATIONS: 6 months. For Renewal - must document therapeutic benefit and confirm compliance Routine PDL edits plus *♦ Clinical Criteria for Hetlioz[®] and tasimelteon - For the treatment of Non-24-Hour Sleep-Wake Disorder (Non-24), AND - Member must be age 16 years of age or older. - Quantity limit = 1 capsule per day. **♦ Clinical Criteria for Hetlioz[™] LQ oral suspension - For the treatment Nighttime sleep disturbances in SMS in pediatric members AND - Member must be 3 years to 15 years of age.
Skeletal Muscle Relaxants		
baclofen cyclobenzaprine HCL methocarbamol tizanidine tab	Amrix [®] baclofen soln baclofen susp (gen Fleqsuvy [™]) *carisoprodol *carisoprodol/ASA *carisoprodol/ASA/codeine chlorzoxazone cyclobenzaprine ER dantrolene sodium Dantrium [®] Flexmid [®] Fleqsuvy [™] metaxalone orphenadrine citrate orphenadrine/ASA/caffeine	LENGTH OF AUTHORIZATIONS: 1 year for chronic conditions - Duration of prescription (up to 3 months) for acute conditions Routine PDL edits plus *♦ Clinical Criteria for Carisoprodol Drugs - Member is at least 16 years old - Only approve for ACUTE, painful musculoskeletal conditions. - Length of Authorization: One month per every 6 months for carisoprodol drugs - Quantity limit = 4 tablets per day

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Preferred Agents		Non-Preferred Agents	SA Criteria
		Ozobax [®] DS *Soma [®] tizanidine cap Zanaflex [®]	
Smoking Cessation			
bupropion SR Chantix [®] Chantix [®] DS PK nicotine gum/lozenge/patch varenicline		Nicoderm CQ [®] Patch Nicorette [®] Gum/Lozenges Nicotrol [®] Inhaler & NS	LENGTH OF AUTHORIZATIONS: 6 months Routine PDL edits
*Stimulants/ADHD Medications (AGE) CLOSED CLASS			
Amphetamine Drugs			LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
amphetamine salts combo (generic for Adderall IR) amphetamine salts combo XR (generic for Adderall XR) dextroamphetamine tab (generic for Dexedrine) dextroamphetamine cap SR (generic for Dexedrine Spansule) Vyvanse [®] capsule (lisdexamfetamine)		Adderall [®] IR (amphetamine salts combo) Adderall [®] XR Adzenys XR ODT [™] amphetamine sulfate (generic) Evekeo [®] amphetamine susp (generic) Adzenys ER [™] susp) Dexedrine [®] dextroamphetamine soln Dyanavel [®] XR susp Evekeo [®] lisdexamfetamine(generic) Vyvanse [®] methamphetamine Mydayis ER [™] Procentra [®] soln Vyvanse [®] chewable tab Xelstry [™] Zenzedi [™]	*♦ All stimulants (preferred and non-preferred) require the submission of Clinical SA if prescribed for a child less than four or an adult eighteen years and older. Member must meet the minimum FDA approved age. Refer to Stimulant SA form (Stimulant/ADHD Medications SA Form) This does not include nonstimulant agents such as atomoxetine, clonidine ER, guanfacine ER or others *Clinical Criteria for Vyvanse[®] chewable tab Member must have tried and failed methylphenidate solution

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Preferred Agents		Non-Preferred Agents	SA Criteria
Methylphenidate Drugs			
All methylphenidate IR generic methylphenidate soln methylphenidate ER tab (Metadate ER tab) methylphenidate ER (Concerta) dexmethylphenidate IR dexmethylphenidate XR		<i>AptensioTM XR</i> <i>AzstarysTM</i> <i>Concerta[®]</i> <i>Cotempla XR-ODTTM</i> <i>Focalin[®] IR & Focalin[®] XR</i> <i>Jornay PM</i> <i>methylphenidate CD cap (generic)</i> <i>Metadate CD)</i> <i>Methylin ER[®], soln IR</i> <i>methylphenidate chew</i> <i>methylphenidate LA, SR</i> <i>methylphenidate ER(generic</i> <i>Relexxii[®])</i> <i>methylphenidate ER(generic</i> <i>AptensioTM XR)</i> <i>methylphenidate (generic</i> <i>Daytrana[®])</i> <i>Ritalin[®] IR</i> <i>Relexxii[®]</i> <i>QuilliChew ERTM</i> <i>QuillivantTM XR susp</i>	
Miscellaneous Drugs			
atomoxetine (generic for Strattera[®]) clonidine ER guanfacine ER		<i>Intuniv[®]</i> <i>OnydaTM XR</i> <i>Qelbree[®]</i>	
♦ Narcolepsy Drugs			
armodafinil modafinil SunosiTM		<i>NuvigilTM</i> <i>Provigil[®]</i> <i>Wakix[®]</i>	Routine PDL edits plus

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Preferred Agents		Non-Preferred Agents		SA Criteria	
				♦All Narcolepsy Drugs require submission of Clinical SA. Refer to: Narcolepsy Medications SA Form	
Dermatologic					
*Acne Agents, Topical (AGE)					
Combo Benzoyl Peroxide, Clindamycin, Erythromycin & other Top			LENGTH OF AUTHORIZATIONS: 1 year		
Acne Medication gel, lot benzoyl peroxide wash, cr, gel, lot (OTC) clindacin ETZ 1% pledget clindamycin ph 1% solution, pledget, swab, gel, lotion clindamycin/benzoyl peroxide (Duac®) erythromycin solution		Acne Clearing System® (OTC) Aklief® Avar Cleanser Avar-E Avar LS Cleanser Azelex® BP 10-1 benzoyl peroxide wash/cr/gel/ lotion/foam/towelette (RX) BPO Cleocin T® Clindacin™ Pac Kit clindamycin phosphate (generic for Clindagel®) clindamycin/benzoyl peroxide (generic for Benzaclin®) clindamycin phosphate foam, el, med swab clindamycin/tretinoin (generic Veltin®) dapsone 5% gel erythromycin gel/med. swab Evoclin™ Neuac™ topical/kit Ovace® Wash		Routine PDL edits plus *♦ <u>Clinical Criteria for Dermatologic Acne Agents</u> - Prescriptions for members over the age of 18 years will require the submission of a SA to evaluate treatment diagnosis; AND - Drugs are intended for acne only. SA for a cosmetic indication cannot be approved.	

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Preferred Agents		Non-Preferred Agents	SA Criteria
		<i>Ovace® Plus shampoo/cr/lotion/foam</i> <i>Sulfacetamide Cleanser ER</i> <i>Sulfacetamide Cleanser, Shampoo, Susp</i> <i>Sulfacetamide Sodium/Sulfur Cr, Susp, Sunscreen</i> <i>SSS 10-5 Foam</i> <i>Sulfacetamide/Sulfur/Cleanser, Cleanser Kit, Lotion Med. Pad</i> <i>Sulfacetamide/Sulfur/Urea Cleanser</i> <i>Sumadan Wash, Kit</i> <i>Sumadan XLT</i> <i>Sumaxin CP Kit</i> <i>Winlevi®</i>	
Retinoids/Combinations, Topical			
adapalene gel OTC tretinoin 0.025, 0.05, 0.1% cream	<i>adapalene 0.1% cr/gel/lot</i> <i>adapalene 0.3% gel/gel w/pump</i> <i>adapalene-benzoyl peroxide (gen Epiduo® and Epiduo® Forte)</i> <i>Differin® 0.1% cr/gel/lot RX</i> <i>Differin® 0.3% gel pump</i> <i>Differin 0.1% gel (OTC)</i> <i>Epiduo® & Epiduo® Forte Gel</i> <i>*Fabior™ 0.1% Foam</i> <i>Tazorac® cr/gel</i> <i>tazarotene 0.1% cr, foam</i> <i>tretinoin 0.01, 0.025% gel</i> <i>tretinoin microsphere 0.04%, 0.08%, & 0.1% gel</i> <i>Twynéo®</i>	*♦ <u>Age Edit for Fabior™ Foam</u> Members must be between the ages of 12 and 18 years of age	

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Preferred Agents		Non-Preferred Agents		SA Criteria	
Antifungal Topical					
antifungal 1% cr, powder antifungal 2% cr ciclopirox soln ciclodan 8% soln clotrimazole cr (OTC & Rx) clotrimazole soln (Rx) clotrimazole-betamethasone cr ketoconazole shampoo ketoconazole cr miconazole cr/spray (OTC) nystatin cr/oint/ powder terbinafine cr (OTC) tolnaftate cr/powder(OTC)		Alevazol® OTC Ciclodan® Kit ciclopirox cr/shampoo/gel ciclopirox kit ciclopirox suspension clotrimazole soln (OTC) clotrimazole-betamethasone lot econazole Ertaczo® Extina® ketoconazole foam Lasolex™ AG Loprox® Lotrimin AF® cr (OTC) miconazole Oint/ powder (OTC) Naftifine CR nystatin-triamcinolone cr/oint oxiconazole cr (generic Oxistat®) tavaborole (generic Kerydin®) tolnaftate aero powde/spray (OTC) tolnaftate soln (OTC) Vusion®		LENGTH OF AUTHORIZATIONS: 6 months Routine PDL edits	
♦Immunomodulators, Atopic Dermatitis					
Adbry™ **Dupixent® **Eucrisa™ *pimecrolimus *tacrolimus		***Anzupgo® *** Ebglyss™ *** Nemludio® ***Opzelura™ Protopic® *****Vtama® ***Zoryve cream, 0.15%		LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus ♦ Clinical Criteria for Atopic Dermatitis-all drugs • Member must have an FDA approved diagnosis: Atopic dermatitis AND • Age and severity for drug as follows below:	

Virginia's Medicaid Preferred Drug List (PDL)/ Common Core Formulary

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	Preferred Agents	Non-Preferred Agents	SA Criteria
		***Zoryve cream, 0.05%	○ Adbry™: moderate to severe for ages ≥ 12 years ○ Ebglyss™: moderate to severe for ages ≥ 12 years ○ Eucisa™: mild to moderate for ages ≥ 3 months ○ Dupixent®: moderate to severe for ages ≥ 6 months ○ Nemluvio®: moderate to severe for ages ≥ 12 years ○ Opzelura™: mild to moderate for ages ≥ 2 years ○ pimecrolimus: mild to moderate for ages ≥ 2 years ○ Protopic® 0.03%: moderate to severe for ages ≥ 2 years ○ Protopic® 0.1%: moderate to severe for ages ≥ 16 years ○ Vtama® for ages ≥ 2 years ○ Zoryve® cream 0.15%: mild to moderate for ages ≥ 6 years ○ Zoryve cream, 0.05%: mild to moderate atopic dermatitis in ages 2 to 5 AND the one that applies: *Clinical Criteria for pimecrolimus, or tacrolimus • Prior documented trial & failure of 8 weeks of one (1) topical corticosteroid of medium to high potency (e.g., mometasone, fluocinolone) **Clinical Criteria for Eucrisa™, Dupixent® and Adbry® (reviewed electronically, AutoPA look back of 180 days for no more than a 30 day) • Prior documented trial & failure of 30-day trial (or contraindication) of • One (1) topical corticosteroid of medium to high potency (e.g., mometasone, fluocinolone) OR • One (1) topical calcineurin inhibitors (tacrolimus or pimecrolimus) *** Clinical Criteria for Ebglyss™, Nemluvio® Opzelura™ (topical Janus kinase inhibitor), Vtama and Zoryve cream 0.05% and 0.15% • Prior documented trial & failure of 8 weeks of each ○ One (1) topical corticosteroid of medium to high potency (e.g., mometasone, fluocinolone) AND ○ One (1) topical calcineurin inhibitors (tacrolimus or pimecrolimus) AND ○ A trial and failure of Dupixent® • Other indications: Opzelura cream can not be approved for the indication of nonsegmental vitiligo in adult and pediatric members ≥ 12 years old

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Preferred Agents	Non-Preferred Agents	SA Criteria														
		Clinical Criteria for Nemluvio® for Prurigo nodularis • Age 18 or older • A diagnosis of prurigo nodularis • An 8-week trial and failure of Dupixent **Clinical Criteria for Vtama® for Plaque Psoriasis • Age 18 or older • A diagnosis of plaque psoriasis ***** Clinical Criteria for Anzupgo (delgocitinib 2% topical cream) • The member is 18 years of age or older, AND • The member has a diagnosis of moderate to severe chronic hand atopic dermatitis, AND • The member has had inadequate response to or has been unable to use topical steroids <u>QUANTITY LIMITS for Atopic Dermatitis</u> <table><tr><th>Drug</th><th>Rolling qty limit</th></tr><tr><td>Adbry®</td><td>4 syringes/14 days (initial dose) 4 syringes/28 days (maintenance)</td></tr><tr><td>Dupixent</td><td>2 syringes (or pens)/14 days (initial dose), 2 syringes (or pens)/28 days (maintenance)</td></tr><tr><td>pimecrolimus</td><td>30gm per 30 days</td></tr><tr><td>Eucrisa® (crisaborole)</td><td>300gm per year</td></tr><tr><td>Opzelura (ruxolitinib)</td><td>240 gm (4 x 60gm) per 30 days</td></tr><tr><td>Protopic® (tacrolimus)</td><td>30gm per 30 days</td></tr></table> <u>Dupixent other Diagnosis: Fax Form</u> ♦ Diagnosis of moderate to severe asthma Member must have moderate to severe asthma diagnosed as ONE of the following types: o Asthma with eosinophilic phenotype with eosinophil count ≥ 150 cells/mcL; OR	Drug	Rolling qty limit	Adbry®	4 syringes/14 days (initial dose) 4 syringes/28 days (maintenance)	Dupixent	2 syringes (or pens)/14 days (initial dose), 2 syringes (or pens)/28 days (maintenance)	pimecrolimus	30gm per 30 days	Eucrisa® (crisaborole)	300gm per year	Opzelura (ruxolitinib)	240 gm (4 x 60gm) per 30 days	Protopic® (tacrolimus)	30gm per 30 days
Drug	Rolling qty limit															
Adbry®	4 syringes/14 days (initial dose) 4 syringes/28 days (maintenance)															
Dupixent	2 syringes (or pens)/14 days (initial dose), 2 syringes (or pens)/28 days (maintenance)															
pimecrolimus	30gm per 30 days															
Eucrisa® (crisaborole)	300gm per year															
Opzelura (ruxolitinib)	240 gm (4 x 60gm) per 30 days															
Protopic® (tacrolimus)	30gm per 30 days															

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	Preferred Agents	Non-Preferred Agents	SA Criteria
			- o Oral corticosteroid dependent asthma with at least 1 month of daily oral corticosteroid use within the last 3 months; AND - o Member must be 6 years of age or older ♦ Diagnosis of eosinophilic esophagitis (EoE); - o Member ≥1 years old; AND - o Member weighs ≥ 15 kg; AND - o Prescribed by or consultation with an allergist or gastroenterologist; AND - o Member did not respond clinically to treatment with a topical glucocorticosteroid or proton pump inhibitor ♦ Diagnosis of chronic rhinosinusitis with nasal polyposis (CRSwNP); - o Member ≥ 12 years old - o Member has inadequate response after 3 consistent months use of preferred PDL intranasal steroids or oral corticosteroids; AND - o Member is concurrently treated with intranasal corticosteroids ♦ Diagnosis of Prurigo Nodularis for the treatment of adult members with prurigo nodularis (PN) - o Age 18 or older - o Diagnosis of PN - o Prescribed by or in consultation with dermatologist, allergist, or immunologist ♦ Diagnosis of inadequately controlled chronic obstructive pulmonary disease (COPD) and an eosinophilic phenotype: - o Age 18 or older - o Prescribed by or in consultation with a pulmonologist - o Member has a diagnosis of COPD that is inadequately controlled and a minimum blood eosinophil count of 300 cells/mcL at screening, measured within the past 12 months - o Member is receiving maximal inhaled therapy consisting of a long-acting muscarinic antagonist (LAMA), long-acting beta agonist (LABA), and inhaled corticosteroid (ICS) (or therapy of LAMA plus LABA if ICS is contraindicated) ♦ Diagnosis of Chronic Spontaneous Urticaria: - o Age 12 or older - o Diagnosis of Chronic Spontaneous Urticaria - o Member had a 30 day trial and failure, or intolerance to H1 antihistamine treatment

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Preferred Agents		Non-Preferred Agents	SA Criteria
			♦ Diagnosis of Bullous Pemphigoid: - o Age 18 or older - o Diagnosis of Bullous Pemphigoid
Psoriasis, Topical			
calcipotriene cr/oint/soln	<i>betamethasone/ calcipotriene</i> <i>calcitriol</i> <i>calcipotriene/betamethasone</i> <i>(generic for Taclonex®)</i> <i>calcipotriene foam</i> <i>*Enstilar® Foam (AGE)</i> <i>Sorilux™</i> <i>Taclonex® & Taclonex® Scalp</i> <i>Zoryve® 0.3% cream</i> <i>**Zoryve foam 0.3%</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits plus *♦ <u>Clinical Criteria for Enstilar® Foam</u> • Length of Authorization: 4 weeks • Diagnosis of plaque psoriasis; AND • Minimum age of 12 years **♦ <u>Clinical Criteria for Zoryve foam 0.3%</u> • Member is 9 years of age or older and has a diagnosis of seborrheic dermatitis • Member has a history of failure, contraindication, or intolerance to BOTH of the following therapies: - o 30 days of therapy with ONE (1) topical corticosteroid (i.e., clobetasol, fluocinonide or mometasone cream/ointment/solution) in the past 180 days AND - o 30 days of therapy with ONE (1) topical antifungal (ciclopirox shampoo/gel, ketoconazole cream/shampoo, selenium sulfide 2.25% shampoo) in the past 180 days	
Rosacea Agents, Topical			

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Preferred Agents		Non-Preferred Agents	SA Criteria
metronidazole cr/gel		azelaic acid (generic for Finacea®) brimonidine gel pump (generic Mirvaso®) Epsolay® Finacea® foam ivermectin (generic Soolantra) Metrocream® Metrogel® metronidazole lot, gel pump Mirvaso® Rhofade® Rosadan™ Kit Soolantra®	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
Steroids			
Steroids, Topical Low Potency			LENGTH OF AUTHORIZATIONS: 1 year
hydrocortisone cr/gel/lot/oint hydrocortisone cr/oint OTC hydrocortisone/aloe cr OTC		alclometasone cr/oint aqua glycolic HC Capex® shampoo Derma-smoothe-FS Desowen® lot fluocinolone 0.01% oil Texacort®	Routine PDL edits
Steroids, Topical Medium Potency			
fluticasone propionate cr/oint mometasone furoate cr/oint/soln		betamethasone valerate foam Beser Lotion and Kit clocortolone cr Cordran® tape Elocon® cr/oint/soln fluocinolone acetonide cr/oint/soln flurandrenolide cr/oint/tape fluticasone propionate lot hydrocortisone valerate cr/oint	

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Preferred Agents		Non-Preferred Agents	SA Criteria
		hydrocortisone butyrate (generic for Locoid Lotion) hydrocortisone butyrate cr/oint/soln/ emollient Momexin [®] prednicarbate oint Synalar [®] Synalar TS [®] Ticanase kit [®]	
Steroids, Topical High Potency			LENGTH OF AUTHORIZATIONS: 1 year
betamethasone valerate cr/lot/oint triamcinolone acetonide cr/lot/oint	amcinonide cr/lot/oint betamet diprop & prop gly cr/lot/oint betamet diprop cr/foam/gel/lot/oint DermacinRx [®] SilaPak [™] DermacinRx [®] Silazone DermacinRx [®] Therazole Pak desoximetasone cr/gel/oint/spray desoximetasone (generic Topicort [®] spray) diflorasone diacetate cr/oint Diprolene [®] lot/oint DiproleneAF [®] cr Ellzia [™] Pak Kit fluocinonide cr/ emollient/ gel/oint/soln halcinonide cr/soln Halog [®] cr/oint/soln Kenalog [®] aerosol Silazone [®] II Kit Topicort [®] cr/gel/oint/spray Trianex [®] oint triamcinolone spray	Routine PDL edits plus	

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Preferred Agents		Non-Preferred Agents	SA Criteria
		triamcinolone/dimethicone <i>Whytederm[®]Tdpak</i>	
Steroids, Topical Very High Potency			
clobetasol emollient clobetasol propionate cr/gel/oint/soln halobetasol propionate cr		<i>Apexicon[™]E</i> <i>clobetasol lot/shampoo</i> <i>clobetasol propionate foam/spray</i> <i>Clobex[®] lot/shampoo/spray</i> <i>Clobetex[®] Kit</i> <i>Clodan[®] kit</i> <i>Halonate[®]</i> <i>halobetasol prop foam</i> <i>halobetasol prop oint (generic for Lexette[®])</i> <i>Impeklo[™]</i> <i>Olux[®]</i> <i>Olux[®]-E</i> <i>Ultravate[®] cr/lotion/oint</i> <i>Ultravate[®] PAC & Ultravate[®] X</i>	
Endocrine and Metabolic Agents			
Androgenic Agents (Testosterone – Topical)			
testosterone pump (generic for AndroGel [®] Pump) testosterone packet (authorized generic for Vogelxo [™])	<i>Axiron[®] soln</i> <i>Fortesta[®]</i> <i>Natesto Nasal Gel[®]</i> <i>Testim[®]</i> <i>testosterone (generic for Axiron[®])</i> <i>testosterone gel/pump (generic for Vogelxo[™])</i> <i>testosterone (generic for Fortesta[®])</i> <i>Vogelxo[™] gel/packet/pump</i> <i>Xyosted[™]</i>	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits	

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Preferred Agents		Non-Preferred Agents	SA Criteria
Antihyperuricemics			
allopurinol 100mg and 300mg colchicine tabs Probenecid[®] probenecid & colchicine	<i>allopurinol 200MG</i> <i>colchicine caps</i> <i>febuxostat (generic Uloric[®])</i> <i>Gloperba[®]</i> <i>Mitigare[®]</i> <i>Zyloprim[®]</i>	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits	
Contraceptives*(long-acting IUDs & injectable)			
Depo-Provera[®] 104mg Kyleena[™] Liletta[®] medroxyprogesterone 150mg Mirena[®] Nexplanon[®] Paragard[®] Skyla[®]	<i>Depo-Provera[®] 150mg</i>	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits	
♦Diabetes Hypoglycemics: Injectable and Oral Incretin Mimetics CLOSED CLASS			
Trulicity[™] Victoza[®]	<i>exenatide</i> <i>liraglutide</i> <i>Mounjaro[®]</i> <i>Ozempic[®]</i> <i>Rybelsus Tab[®]</i> <i>Soliqua[®] 100/33</i> <i>Tanzeum[™]</i> <i>Xultophy[®] 100/3.6</i>	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits Preferred Agents Clinical Criteria: • Diagnosis of type II diabetes Non-Preferred Incretin Mimetics SA Form	
Diabetes Hypoglycemics: Injectable Insulins			
Insulin Mix		LENGTH OF AUTHORIZATIONS: 1 year	
Humulin[®] 70/30 pen/vial(OTC)	<i>Humalog[®] Mix 50/50 Kwikpen</i> <i>Humalog[®] Mix 75/25 Kwikpen</i> <i>Humalog Mix 75/25 vial</i> <i>Novolin[®] 70/30 vial (OTC)</i>	Routine PDL edits	

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Preferred Agents		Non-Preferred Agents	SA Criteria
insulin aspart/insulin aspart protamine vial insulin lispro protamine mix kwikpen (Authorized Generic)		Novolog [®] Mix 70/30 pen/vial	*Clinical Criteria for Toujeo Solostar: The member has tried and failed or is intolerant to Lantus
Insulin N			
Humulin[®] N pen/vial (OTC)		Humulin [®] N pen Novolin [®] N vial (OTC)	
Insulin R			
Humulin[®] R pen/vial		Novolin [®] R vial (OTC)	
Long-Acting Insulins			
Lantus[®] Solostar[®] & vial *Toujeo Solostar[®] pen		Basaglar [®] KwikPen [®] insulin degludec insulin glargine max solo u300 insulin glargine solostar u300 insulin glargine-yfgn Rezvoglar [®] Kwikpen Semglee [™] Toujeo [®] Max Solostar [®] 300 Units/mL Tresiba [®] FlexTouch [®] Pen 100 U/mL, 200 U/mL	
Rapid-Acting Insulins			
Humulin 500 U/M pen/vial insulin lispro vial insulin lispro Jr. Kwikpen insulin lispro Pen insulin aspart cartridge pen/vial insulin aspart/insulin aspart protamine insulin pen		Admelog [®] Solostar Pen/vial Afrezza [®] cartridge (inhalation) Apidra [®] cartridge/Solostar/vial Fiasp [®] /FlexTouch [®] Pen/PenFill [®] Humalog Kwikpen 200 unit/mL Humalog vial/pen Humalog Cartridge Humalog Kwikpen 100 unit/mL	

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Preferred Agents		Non-Preferred Agents	SA Criteria
		Humalog Jr. Kwikpen Kirsty vial/pen Lyumjev TM Merilog TM Novolog [®] cart/vial/Flexpen	
Diabetes Oral Hypoglycemics			
Oral Hypoglycemics Alpha-Glucosidase Inhibitors			LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus
acarbose		miglitol (generic Glyset [®])	♦ Age edit for all Oral Hypoglycemics is 18 years of age or older, except Metformin which is 10 years of age.
Oral Hypoglycemics Biguanides			
metformin 500, 850, 1000mg metformin ER (generic for Glucophage [®] XR)		Riomet [®] susp metformin 625 mg metformin 750 mg metformin ER (generic Fortamet [®]) metformin ER (generic Glumetza [®]) metformin (generic Riomet [®])	
Oral Hypoglycemics Biguanide Combination Drugs			
glyburide/metformin		glipizide/metformin Glucovance [®]	
Oral Hypoglycemics DPP-IV Inhibitors & Combination			
CLOSED			
Janumet [®] Januvia [®] Jentadueto TM Jentadueto XR TM Tradjenta TM		alogliptin (generic Nesina TM) alogliptin/metformin (generic Kazano TM) alogliptin/pioglitazone (generic Oseni TM) Brynovin TM Glyxambi [®] Janumet XR [®] linagliptin/metformin	

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Preferred Agents		Non-Preferred Agents	SA Criteria
		Kazano TM Nesina TM Oseni TM Qtern [®] saxagliptin hcl(generic Onglyza TM) saxagliptin/metformin ER (generic Kombiglyze XR TM) Steglujan TM Trijardy TM XR Zituvio TM Zituvimet TM Zituvimet TM XR	
Oral Hypoglycemics Meglitinides			
nateglinide repaglinide		repaglinide/metformin Starlix	
Oral Hypoglycemics Second Generation Sulfonylureas			
glimepiride 1,2,4 mg glipizide glipizide ER glyburide glyburide micronized		glimepiride 3mg Glucotrol [®] Glucotrol XL [®]	
Oral Hypoglycemics Sodium Glucose Co-Transporter 2 Inhibitor CLOSED CLASS			
Farxiga TM Jardiance [®] Synjardy [®] Synjardy [®] XR Xigduo TM XR		dapagliflozin dapagliflozin-metformin ER Inpefa TM Invokana TM Invokamet TM Invokamet TM XR Segluromet TM Steglatro TM	

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Preferred Agents	Non-Preferred Agents	SA Criteria
Oral Hypoglycemics Thiazolidinediones		
pioglitazone	Actoplus Met® IR & XR Actos® Avandaryl® Avandamet® Duetact® pioglitazone/metformin pioglitazone/glimepiride	
Erythropoiesis Stimulating Proteins		
Epogen® Retacrit™ (PFIZER)	Aranesp® vial/syringe Mircera® Procrit® Reblozyl® Retacrit™ (VIFOR) Vafseo®	LENGTH OF AUTHORIZATIONS: for duration of the prescription up to 6 months Routine PDL edits
Glucagon Agents CLOSED CLASS		
Baqsimi nasal Proglycem suspension oral Zegalogue Autoinjector & syringe SQ	diazoxide suspension oral Glucagon emergency kit (Lilly) inj Glucagon inj Glucagon emergency kit (Fresenius and Amphastar) inj Gvoke pen, syringe, vial SQ	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
Glucocorticoids, Oral		
budesonide EC dexamethasone soln/tab hydrocortisone methylprednisolone dose pk methylprednisolone 4 mg tab prednisolone sod phosph soln prednisolone soln	Alkindi® Sprinkle Cortef® cortisone acetate dexamethasone elixir/intensol Entocort® EC Eohilia™ Flo-Pred®	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits

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Preferred Agents		Non-Preferred Agents	SA Criteria
prednisone soln/tab/dose pk		<i>Hemady[®]</i> <i>Khindivi</i> <i>Medrol[®] dose pk/tab</i> <i>methylprednisolone 8,16 & 32mg tab</i> <i>Millipred DP[®] tab Does Pk</i> <i>Millipred[®] tab</i> <i>Orapred[®] ODT</i> <i>prednisolone sod phosphate ODT/ (10 mg/5ml&20 mg/5ml soln)</i> <i>prednisone intensol</i> <i>prednisone DR tab</i> <i>TaperDex[®]</i> <i>Veripred[®]</i>	
*Growth Hormone		CLOSED CLASS	
Genotropin [®] Cartridge, Miniquick Norditropin FlexPro [®]		<i>Humatrope[®] cartridge/vial</i> <i>Ngenla[™]</i> <i>Nutropin AQ[®] NuSpin[™]</i> <i>Omnitrope[®] cartridge/vial</i> <i>Serostim[®] vial</i> <i>Skytrofa[™]</i> <i>Sogroya[®]</i> <i>Zomacton[®] vial</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year ♦ALL Growth Hormone drugs (preferred and non-preferred) require the submission of a Clinical SA. Refer to (Growth Hormone SA Fax Form)
*Hereditary Angioedema (HAE) Agents			
Berinert [®] Cinryze [™] icatibant (generic Firazyr [®]) Kalbitor [®] Sajazir [™]		<i>Andembry[®]</i> <i>Dawnzera[™]</i> <i>Ekterly[®]</i> <i>Firazyr[®]</i> <i>Haegarda[®]</i> <i>Orladeyo[™]</i> <i>Ruconest[®]</i> <i>Takhzyro[™]</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits plus *♦ALL Hereditary Angioedema drugs (preferred and non-preferred) require the submission of a Clinical SA. Refer to Hereditary Angioedema (HAE) SA Form **Black box warning KALBITOR[®] Anaphylaxis has been reported after administration of KALBITOR [®] . Because of the risk of anaphylaxis, KALBITOR should only be administered by a healthcare professional with appropriate medical support to manage anaphylaxis and

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Preferred Agents		Non-Preferred Agents	SA Criteria
			hereditary angioedema. Healthcare professionals should be aware of the similarity of symptoms between hypersensitivity reactions and hereditary angioedema and members should be monitored closely.
*Pancreatic Enzymes			
Creon® Zenpep®	Pancreaze® Pertzye® Ultresa® Viokace®		LENGTH OF AUTHORIZATION: 1 year Routine PDL edits plus *Clinical Criteria for Pancreatic Enzymes - All drugs preferred and non preferred require a diagnosis of pancreatic insufficiency due to cystic fibrosis or chronic pancreatitis or pancreatectomy. - If a member has a diagnosis of Cystic Fibrosis they do not have to try and fail of a preferred. - If member has a feeding tube, then two different pancreatic enzymes can be approved for use together.
Progestational Agents CLOSED CLASS			
medroxyprogesterone acetate (tab only) norethindrone acetate progesterone cap/inj	Aygestin® Crinone (Vaginal) Depo-Provera 400 MG/ML Prometrium® Provera®		LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
Vaginal Estrogens			
Premarin® Vaginal cr Vagifem® Vaginal tab	Estrace® Vaginal cr estradiol cream (gen Estrace®) estradiol vaginal tab Estring® Vaginal ring Femring® Vaginal ring Imvexxy® Intrarosa™ Yuvaferm®		LENGTH OF AUTHORIZATIONS: 6 months Routine PDL edits
Weight Management Agents CLOSED CLASS			
orlistat Xenical	Imcivree SQ liraglutide(gen Saxenda®)		Length of authorization: As indicated on fax form

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<i>Preferred Agents</i>		<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
phendimetrazine IR and ER phentermine benzphetamine diethylpropion IR and ER		<i>Lomaira™</i> <i>phentermine/topiramate ER</i> <i>Saxenda®</i> <i>Wegovy®</i> <i>Zepbound™</i>	Routine PDL edits plus Weight Management Agents SA Form GLP-1 Receptor Agonist For Cardiovascular Risk Reduction SA Form GLP-1 Receptor Agonist For Obstructive Sleep Apnea SA Form GLP-1 Receptor Agonists for Metabolic Dysfunction-Associated Steatohepatitis SA Form ♦ The selected GLP-1 Receptor Agonist is brand Saxenda
Gastrointestinal			
G I Antibiotics			
metronidazole 250mg, 500mg tab vancomycin cap vancomycin for reconstitution oral soln kit (Firvanq™ soln)		<i>*Aemcolo™</i> <i>Alinia®</i> <i>Dificid®</i> <i>fidaxomicin</i> <i>Firvanq™ soln</i> <i>Flagyl® cap/ER</i> <i>Likmez™</i> <i>metronidazole 125mg</i> <i>metronidazole cap</i> <i>neomycin</i> <i>nitazoxanide (generic Alinia®)</i> <i>paromomycin</i> <i>Solosec®</i> <i>Tindamax®</i> <i>tinidazole</i> <i>vancomycin sol</i> <i>Vancocin®</i> <i>**Vowst™</i>	Length of authorization: 1 year Routine PDL edits *♦ Clinical Criteria for Aemcolo - Diagnosis of travelers' diarrhea with moderate diarrhea that is distressing or interferes with planned activities AND - Documentation of a history of failure, contraindication, or intolerance to one or more of the following: Azithromycin (generic Zithromax), Ciprofloxacin (generic Cipro), Levofloxacin (generic Levaquin), Ofloxacin (generic Floxin) **♦ Clinical Criteria for Vowst™: - Member ≥ 18 years of age; AND - Member has a confirmed diagnosis of recurrent <i>Clostridioides difficile</i> infection (CDI) with a total of ≥3 episodes of CDI within 12 months; AND - Antibiotic treatment for recurrent CDI must be completed 2 to 4 days prior to initiation of Vowst therapy; AND

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Preferred Agents		Non-Preferred Agents	SA Criteria
			- Member will take 10 oz of magnesium citrate (or 250 mL polyethylene glycol electrolyte solution for members with impaired kidney function) the evening prior to initiation of Vowst therapy; AND - Member must not have absolute neutrophil count (ANC) < 500 cells/mm³, toxic megacolon, or small bowel ileus - Quantity limit = 12 capsules per 3 days - Renewal Criteria: Not applicable Length of authorizations: 30 Days
Antiemetic/Antivertigo Agents			
Cannabinoids (delta-9THC derivatives)			LENGTH OF AUTHORIZATIONS: 6 months
*dronabinol cap	*Marinol [®]		*Dronabinol plus all non-preferred Antiemetic/Antivertigo agents require submission of a Clinical SA. Refer to Antiemetic/Antivertigo SA form
5HT₃ Receptor Blockers			LENGTH OF AUTHORIZATIONS: 3 months, unless otherwise noted
ondansetron ODT/soln/tab	Akynzeo [®] granisetron tab, vial ondansetron vial, amp, syringe palonosetron vial, syringe Posfrea [™] Sancuso [®] patch Sustol [®]		Routine PDL edits plus: For ondansetron 16mg ODT: The member has tried and failed or is intolerant to ondansetron 8mg ODT.
NK-1 Receptor Antagonist			LENGTH OF AUTHORIZATIONS: Length of chemotherapy regimen or a maximum of 6 months
	Aponvie [™] aprepitant capsule/pack Cinvanti [™] (Intraven) Emend [®] Bi Pak/ cap Emend [®] Tri-fold pack/susp Emend [®] vial Focinvez [™] fosaprepitant vial		

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Preferred Agents		Non-Preferred Agents	SA Criteria
Other			LENGTH OF AUTHORIZATIONS: 1 year, unless otherwise noted
*Diclegis® meclizine (OTC, Rx) metoclopramide soln, tab, vial prochlorperazine tab **promethazine soln, syrp, tab, 12.5mg&25mg supp (AGE)	Antivert® Barhemsys® *Bonjesta™ Compazine® supp/tab Compro® dimenhydrinate tab, vial *doxylamine succinate/ vit B6 metoclopramide ODT metoclopramide syringe **Phenergan® (AGE) prochlorperazine supp, vial **promethazine vial, ampul (AGE) **Promethegan 50 mg supp(AGE) Reglan® scopolamine (gen Transderm-Scop®) Tigan® Transderm-Scop® trimethobenzamide	**♦Promethazine products approved for members ≥ 2 years *♦Bonjesta®/Diclegis®/doxylamine succinate/vit B6: Approve through EDD. member must be pregnant and at least 18 years of age.	
*GI Motility, Chronic			
**Linzess™ lubiprostone and AG Movantik®	alosetron Amitiza® Ibsrela® Lotronex® Motegrity™ prucalopride Symproic® Viberzi™	LENGTH OF AUTHORIZATIONS: 6 months Routine PDL edits plus *♦All GI Motility drugs (preferred and non-preferred) require the submission of a Clinical SA. Refer to Chronic GI Motility SA form **Pediatric use of Linzess 72 mcg only: 1. Diagnosis of functional constipation (FC) 2. Member is 6 to 17 years of age	

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Preferred Agents		Non-Preferred Agents	SA Criteria
			3. Prescriber attestation that other causes of constipation have been ruled out 4. Member has had treatment failure on at least TWO of the following: Osmotic Laxatives (i.e., lactulose, polyethylene glycol, sorbitol), OR Bulk Forming Laxatives (i.e., psyllium, fiber), OR Stimulant Laxatives (i.e., bisacodyl, senna)
H. Pylori Treatment			
Pylera®	<i>bismuth-metronidazole-tetracycline (gen Pylera®)</i> <i>Helidac®</i> <i>Omeclamox®-Pak</i> <i>lansoprazole/amoxicillin/clarithromycin</i> <i>Prevpac®</i> <i>Talicia®</i> <i>Voquezna®</i>		LENGTH OF AUTHORIZATIONS: 14 days Routine PDL edits
Laxatives & Cathartics CLOSED CLASS			
Bowel Prep Agents			LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
Gavilyte-C solution Gavilyte-G solution PEG 3350/electrolytes solution for reconstitution PEG-3350/flavor packs solution for reconstitution sodium, potassium, mag sulfates (Authorized Generic Suprep)	<i>Clenpiq</i> <i>Golytely</i> <i>Nulytely With Flavor Packs</i> <i>PEG 3350/electrolytes powder pack (gen Moviprep)</i> <i>Suflave Powder</i> <i>Suprep</i> <i>Sutab</i>		
Constipation			
lactulose solution	*lactulose packet *Kristalose Packet		* Kristalose/lactulose packet: Must have tried, failed or be intolerant to generic lactulose solution
Proton Pump Inhibitors			
omeprazole RX pantoprazole Protonix® susp	<i>dexlansoprazole dr (gen Dexilant®)</i> <i>Dexilant®</i>		LENGTH OF AUTHORIZATIONS: 12 weeks; unless member meets an exception; then 1 year

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Preferred Agents		Non-Preferred Agents	SA Criteria
		esomeprazole magnesium cap/susp/tab esomeprazole strontium lansoprazole cap Konvomep® Nexium® omeprazole OTC omeprazole magnesium OTC omeprazole/sodium bicarbonate pantoprazole susp (generic) Protonix® susp Prevacid® RX, OTC & SoluTab Prilosec® Rx & Susp Protonix® rabeprazole DR tab	Routine PDL edits plus *♦All non-preferred Proton Pump Inhibitors require submission of a Clinical SA. Refer to Proton Pump Inhibitor SA form Preferred agents require a SA for use over 90 days
Ulcerative Colitis Oral and Rectal Preparations (5-ASA DERIVATIVES)			
Ulcerative Colitis – Oral			LENGTH OF AUTHORIZATIONS: 1 year
balsalazide disodium mesalamine ER (AG of Apriso) Pentasa® sulfasalazine DR & IR	Asacol® HD Azulfidine® IR & DR budesonide ER (gen Uceris™) Delzicol™ Dipentum Lialda® mesalamine (gen Apriso®) mesalamine (gen Asacol® HD) mesalamine (gen Delzicol) mesalamine (gen Lialda®) mesalamine ER (gen Pentasa®)	Routine PDL edits	
Ulcerative Colitis – Rectal			
mesalamine rectal supp mesalamine enema	budesonide (generic Uceris foam) Canasa® rectal supp mesalamine kit Rowasa® enema/kit		

Preferred Agents		Non-Preferred Agents	SA Criteria
		SFRowasa®	
Genitourinary			
Alpha-Blockers and Androgen Hormone Inhibitors For Benign Prostatic Hypertrophy (BPH)			
Alpha-Blockers for BPH		LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits 	

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Preferred Agents		Non-Preferred Agents	SA Criteria
		Toviaz™ VESicare® and LS	
Immunological Agents			
*♦Multiple Sclerosis		CLOSED CLASS	
Avonex® Avonex® Adm Pack Copaxone 20 mg syringe® dimethyl fumarate and Starter Pack fingolimod **Kesimpta® (ST) teriflunomide (generic Aubagio®)		Aubagio® Bafiertam™ Betaseron® Briumvi™ cladribine Copaxone® 40 mg syringe® Gilenya® glatiramer 20 mg/ml syringe Glatopa™ Mavenclad® Mayzent® Ocrevus® Ocrevus Zunovo™ Plegridy® Ponvory™ Rebif® SQ Rebif® Rebidose Pen® Tascenso ODT™ Tecfidera® Tecfidera® Starter Pack Tyruko Tysabri® Vumerity™ Zeposia®	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus *Adherence to the documented PI age and diagnosis required, reviewed electronically, AutoPA **Kesimpta® requires step trial of dimethyl fumarate (generic Tecfidera®) or a preferred Injectable agent, to be determined by an AutoPA. Multiple Sclerosis SA Form Briumvi SA Form Ocrevus/Ocrevus Zunovo SA Form Tysabri and biosimilars SA Form Ampyra® and dalfampridine ER require the submission of a Clinical SA. Refer to MS - Ampyra® SA form
Drugs to improve walking in members with (MS)			
dalfampridine ER (generic Ampyra®)		Ampyra®	

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Preferred Agents	Non-Preferred Agents	SA Criteria
♦Cytokine and CAM Antagonists And Related Agents		
adalimumab-adbm (unbranded version of Cyltezo made by Boehringer Ingelheim) Enbrel® Pen, Sureclick, Hadlima™ Pyzchiva® Syringe, Vial Starjemza™	<i>Abrilada™</i> <i>Actemra® SQ & ACTPEN</i> <i>adalimumab-aacf</i> <i>adalimumab-aaty</i> <i>adalimumab-adbm(Quallent)</i> <i>adalimumab-adaz</i> <i>adalimumab-fkjp Pen</i> <i>adalimumab-ryvk</i> <i>Amjevita™</i> <i>Amjevita™ Autoinjector</i> <i>Arcalyst®</i> <i>Avsola™</i> <i>Avtozma®</i> <i>Bimzelx®</i> <i>Cibinqo™</i> <i>Cimzia® & Cimzia® Syringe Kit</i> <i>Cosentyx™</i> <i>Cosentyx™ UnoReady pen</i> <i>Cyltezo®</i> <i>Enspryng™</i> <i>Entyvio®</i> <i>Hulio®</i> <i>Humira® Pen, Syringe</i> <i>Hyrimoz®</i> <i>Ilaris®</i> <i>Ilumya™</i> <i>Inflectra®</i> <i>infliximab (gen Remicade®)</i> <i>Kevzara® inj, pen</i> <i>Kineret®</i> <i>Olumiant®</i> <i>OmvoH™</i> <i>Orencia®</i> <i>Otezla®</i>	CLOSED CLASS LENGTH OF AUTHORIZATION: 1 year Routine PDL edits plus For preferred agents members must meet FDA approved age and indication. (reviewed electronically, AutoPA look back of 180 days for diagnosis) For preferred ustekinumab products members must also have tried and failed ONE preferred TNF inhibitor *All non-preferred Cytokine and CAM Antagonists require submission of a Clinical SA. For brand name Humira and Stelara please see additional instructions in the SA form. Refer to Cytokine and CAM Antagonists and Related Agents SA Form ** See Cytokine and CAM Antagonists Appendix A for Clinical Criteria** For a list of Cytokine and CAM Antagonists and criteria for approval see Appendix A

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Preferred Agents		Non-Preferred Agents	SA Criteria
		<i>Otezla XRTM</i> <i>Otufi[®]</i> <i>Remicade[®]</i> <i>Renflexis[®]</i> <i>Rinvoq[®]</i> <i>Rinvoq[®] LQ</i> <i>Selarsdi[®]</i> <i>SimlandiTM</i> <i>Simponi[®]</i> <i>Simponi[®] Aria</i> <i>SkyriziTM syringe, Pen</i> <i>Spevigo[®]</i> <i>Stelara[®] vial/syringe</i> <i>Stegeyma[®]</i> <i>SotyktuTM</i> <i>Taltz[®]</i> <i>TofidenceTM</i> <i>TremfyaTM</i> <i>Tyenne[®]</i> <i>Uplizna[®]</i> <i>ustekinumab-aaaz</i> <i>ustekinumab-aekn</i> <i>ustekinumab-ttwe</i> <i>Velsipity[®]</i> <i>XeljanzTM, XeljanzTM XR and sol</i> <i>YesintekTM</i> <i>Yuflyma[®]</i> <i>YusimryTM</i> <i>ZymfentraTM</i>	
♦Methotrexate CLOSED CLASS			<u>LENGTH OF AUTHORIZATIONS: 1 year</u>
methotrexate tab/PF vial/MDV	<i>Jylamvo[®]</i> <i>RasuvoTM</i> <i>Trexall[®]</i>		<u>Routine PDL edits</u>

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Preferred Agents		Non-Preferred Agents	SA Criteria
		Xatmep TM	
Ophthalmic			
Antibiotics			
bacitracin/polymyxin b sulfate oint ciprofloxacin drops erythromycin gentamicin drops/oint moxifloxacin drops (generic Vigamox) ofloxacin drops polymyxin/trimethoprim tobramycin	<i>AzaSiteTM drops</i> <i>bacitracin</i> <i>besifloxacin susp</i> <i>Besivance[®] drops</i> <i>Ciloxan[®] drops/oint</i> <i>Garamycin[®] drops/oint</i> <i>gatifloxacin 0.5% soln</i> <i>levofloxacin drops</i> <i>moxifloxacin drops (generic Moxeza[®])</i> <i>Natacyn[®]</i> <i>neomycin/polymix/gramicidin</i> <i>neomycin/bacitracin/polymyxin oint</i> <i>Neosporin[®]</i> <i>Ocuflox[®]</i> <i>Polytrim[®]</i> <i>sulfacetamide oint/ soln</i> <i>Tobrex[®] drops/oint</i> <i>Vigamox[®]</i> <i>Zymaxid[®]</i>	<u>LENGTH OF AUTHORIZATIONS:</u> Routine PDL edits	Date of service only; no refills
Antibiotic/Steroid Combinations			
neomycin/polymyxin/dexame thasone oint/susp sulfacetamide/prednisolone Tobradex[®] oint tobramycin/dexamethasone susp	<i>Blephamide[®] S.O.P.</i> <i>loteprednol/tobramycin ophthalmic (generic Zylet)</i> <i>Maxitrol[®] oint/susp</i> <i>neomycin/bacitracin/poly/HC</i> <i>neomycin/polymyxin/HC</i> <i>Pred-G[®] oint</i>	<u>LENGTH OF AUTHORIZATION:</u> Routine PDL edits	Date of service only; no refills

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Preferred Agents		Non-Preferred Agents	SA Criteria
		Tobradex® ST Zylet®	
Antihistamines/Mast Cell Stabilizers			
Antihistamines		<u>LENGTH OF AUTHORIZATIONS:</u> 1 year	
Alaway OTC® ketotifen fumerate olopatadine (generic Patanol & Pataday) (OTC)	alcaftadine (gen Lastacast®) azelastine bepotastine (gen Bepreve®) Bepreve® Elestat® epinastine 0.05% eye drops Lastacast® olopatadine (generic Patanol & Pataday) (Rx) olopatadine 0.7% Optivar® Patanol® Rx and OTC Pataday® Rx and OTC Zaditor® OTC Zerviate™	Routine PDL edits	
Mast Cell Stabilizers			
cromolyn sodium			
Anti-inflammatory Agents			
Corticosteroids			
Durezol® prednisolone acetate	Alrex™ Dexamethasone difluprednate (gen Durezol®) Flarex® fluorometholone FML®, FML Forte® loteprednol etabonate Maxidex®		

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Preferred Agents	Non-Preferred Agents	SA Criteria
	Omnipred[®] Pred Forte[®] & Pred Mild[®] prednisolone sod phosphate Vexol[®]	
Immunomodulators CLOSED CLASS		
Restasis[®] Xiidra[®]	Cequa[™] cyclosporine *Eysuvis Miebo[™] Restasis Multidose[®] Tryptyr[®] Tyrvaya[™] Nasal Spray **Verkazia[®]	*♦DUR DURATION LIMIT Eysuvis 1 bottle in 3 months, message returned to pharmacist stating "Limitation Exceeded: Quantity Limit of 1 Bottle per 3 Months. Member needs Intraocular Pressure (IOP) Test. If IOP test completed, Physician to call 1-800-932-6648 for a Clinical SA" **♦<u>Verkazia criteria (PDL criteria do not apply)</u> • Member is ≥4 years of age; AND • Diagnosis of moderate to severe vernal keratoconjunctivitis; AND • Trial and failure, contraindication, or intolerance to one of the following: - o Topical ophthalmic "dual-action" mast cell stabilizer and antihistamine (e.g., olopatadine, azelastine) OR - o Topical ophthalmic mast cell stabilizers (e.g., cromolyn); AND • Prescribed by ophthalmologist or optometrist in consultation with an ophthalmologist • Quantity Limit: 120 single-dose vials per 30 days • Length of approval: 1 year
NSAIDs		LENGTH OF AUTHORIZATIONS: Date of service only; no refills
diclofenac sodium flurbiprofen sodium ketorolac 0.5%	Acular[®] 0.5% & LS[®] 0.4% Acuvail[®] bromfenac sodium 0.07% bromfenac sodium 0.075% bromfenac 0.09% BromSite[™] *Ilevro[™] 0.3% (QL)	Routine PDL edits *Ilevro[™] is limited to 1 bottle plus 1 refill

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Preferred Agents		Non-Preferred Agents	SA Criteria
		<i>Inveltys™ (loteprednol etabonate)</i> <i>ketorolac LS 0.4%</i> <i>Nevanac®</i> <i>Ocufen®</i> <i>Prolensa™</i>	
Glaucoma Agents			
Alpha 2 Adrenergic Agents			LENGTH OF AUTHORIZATIONS: 1 year
Alphagan P® 0.1 & 0.15%	brimonidine 0.2%	<i>apraclonidine 0.5% drops</i> <i>brimonidine tartrate 0.1 & 0.15%</i> <i>Iopidine® 0.5% & 1%</i>	Routine PDL edits
Beta Blockers			
carteolol 1%	Combigan®	<i>Betagan® 0.5%</i> <i>Betimol®</i> <i>betaxolol 0.5%</i> <i>brimonidine-timolol (gen Combigan®)</i> <i>Istalol® 0.5%</i> <i>timolol (generic for Betimol®)</i> <i>timolol (generic for Timoptic Ocudose)</i>	
levobunolol 0.5%	metipranolol 0.3%		
timolol maleate			
Carbonic Anhydrase Inhibitors			
Azopt®	dorzolamide	<i>brinzolamide</i> <i>Cosopt® 0.5%-2%</i> <i>Cosopt® PF</i> <i>Simbrinza™</i> <i>Trusopt® 2%</i> <i>dorzolamide/timolol PF</i>	*Indicated for the treatment of presbyopia in adults
dorzolamide/timolol			
Cholinergic Muscarinic Receptor Agonist			
		<i>pilocarpine</i> <i>*Vuity™</i>	

Preferred Agents		Non-Preferred Agents	SA Criteria
Rho Kinase Inhibitor			
Rhopressa®			
Rocklatan			
Prostaglandin Analogs			
latanoprost		<i>bimatoprost</i>	
Travatan Z®		<i>Iyuzeh™</i>	
		<i>Lumigan® 0.03% & 0.01%</i>	
		<i>Rescula®</i>	
		<i>tafluprost</i>	
		travoprost 0.004%	
		<i>Vyzulta™</i>	
		<i>Xalatan® 0.005%</i>	
		<i>Zioptan™</i>	
Respiratory			
*Anti-Allergens, Oral			
Grass Pollen			LENGTH OF AUTHORIZATIONS: 1 year
		<i>*Grastek®</i>	*♦ All non-preferred Anti-Allergen drugs require the submission of a Clinical SA. Refer to Anti-Allergens, Oral SA Form or Palforzia SA Form
		<i>*Odactra®</i>	
		<i>*Oralair® SL</i>	
		<i>*Ragwitek™</i>	
Peanut			
		<i>*Palforzia™</i>	
Antihistamines: First and Second Generation			
First Generation Antihistamines			LENGTH OF AUTHORIZATIONS: 1 year
Generic only class		<i>All Brands require a SA</i>	Routine PDL edits
Second Generation Antihistamines and Combinations			
cetirizine liquid 1mg/1mL (RX/OTC)		<i>cetirizine chew tab (OTC)</i>	
cetirizine tabs OTC		<i>cetirizine liquid 5mg/5mL (OTC)</i>	
		<i>cetirizine D tab (OTC)</i>	

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levocetirizine tab loratadine tab/syrup OTC		<i>Clarinet[®]</i> <i>Clarinet-D[®]</i> <i>desloratadine ODT, sol</i> <i>fexofenadine</i> <i>fexofenadine/PSE ER</i> <i>fexofenadine suspension</i> <i>levocetirizine sol</i> <i>loratadine ODT</i> <i>loratadine D 12 & 24 hr</i>	
Beta-Adrenergic Agents			
Long Acting Beta Adrenergic s (LABA) MDIs or Nebulizers		LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits	
arformoterol (authorized generic Brovana[®]) Serevent Diskus[®]		<i>arformoterol (generic Brovana[®])</i> <i>Arcapta DS</i> <i>Brovana[®]</i> <i>formoterol fumarate (generic Perforomist[®])</i> <i>Perforomist[®]</i> <i>Striverdi[®] Respimat</i>	
Short Acting Metered Dose Inhalers or Devices			
<i>albuterol HFA (Proair) (Authorized Generic)</i> <i>albuterol HFA (Proventil) and (Authorized Generic)</i> Ventolin[®] HFA		<i>albuterol HFA (Proair[®])</i> <i>albuterol HFA (Ventolin[®])</i> <i>levalbuterol tartrate HFA</i> <i>ProAir[®] RespiClick</i> <i>Xopenex[®] HFA</i>	
Short Acting Nebulizers			
albuterol sulfate (premixed)		<i>levalbuterol soln</i> <i>Xopenex[®]</i>	

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Preferred Agents		Non-Preferred Agents	SA Criteria
COPD: Anticholinergics, Bronchodilators and Phosphodiesterase 4 (PDE4) Inhibitors CLOSED CLASS			
Short Acting Anticholinergics/Short Acting Bronchodilators CLOSED CLASS			
Atrovent HFA® ipratropium bromide soln			
Short Acting Anticholinergics/Short Acting Bronchodilators CLOSED CLASS			
Combivent® Respimat ipratropium/albuterol nebs		Bevespi Aerosphere™ Duaklir Pressair	<u>LENGTH OF AUTHORIZATION:</u> 1 year Routine PDL edits
Long Acting Anticholinergics CLOSED CLASS			
Spiriva® Spiriva® Respimat		Incruse™ Ellipta® tiotropium (generic for Spiriva®) Tudorza™ *Yupelri™ (revefenacin)	<u>LENGTH OF AUTHORIZATION:</u> 1 year Routine PDL edits plus <u>*Clinical Criteria Yupelri™</u> (reviewed electronically, AutoPA) • Diagnosis of COPD AND • Trial/failure of Spiriva® Handihaler OR Spiriva® Respimat
Long Acting Anticholinergics/Long Acting Bronchodilators CLOSED CLASS			
Anoro™ Ellipta® Stiolto Respimat™		umeclidinium-vilanterol	<u>LENGTH OF AUTHORIZATION:</u> 1 year Routine PDL edits
Phosphodiesterase 3 (PDE3) inhibitor and Phosphodiesterase 4 (PDE4) inhibitor CLOSED CLASS			
		Ohtuvayre™	

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Preferred Agents		Non-Preferred Agents	SA Criteria
Phosphodiesterase 4 (PDE4) Inhibitors CLOSED CLASS			
*roflumilast		*Daliresp[®]	LENGTH OF AUTHORIZATION: 1 year *Clinical Criteria for Daliresp[®] and roflumilast • If the member has a diagnosis of severe COPD associated with chronic bronchitis and a history of exacerbations; AND • Trial/failure on at least one first-line or second-line agent (inhaled anticholinergics, long-acting beta agonists or inhaled corticosteroids); AND • Adjunctive therapy (Daliresp[®] must be used in conjunction with first-line or second-line agent).
Corticosteroids: Inhaled and Nasal Steroids			
Inhaled Corticosteroids: Combination Drugs (Glucocorticoid and Long Acting Beta Adrenergic) CLOSED CLASS			LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
Advair[®] Diskus Advair[®] HFA Dulera[®] Symbicort[®] Trelegy[®] Ellipta		<i>Airsupra</i> <i>budesonide-formoterol (generic Symbicort[®])</i> <i>Breo[®] Ellipta[™]</i> <i>Breyna[™] (generic Symbicort[®])</i> <i>Breztri Aerosphere[™]</i> <i>fluticasone/salmeterol (generic Airduo)</i> <i>fluticasone/salmeterol powder (generic Advair)</i> <i>fluticasone/salmeterol HFA (generic Advair HFA)</i> <i>fluticasone-vilanterol (generic Breo[®] Ellipta[™])</i> <i>Wixela[®] (fluticasone/salmeterol)</i>	
Inhaled Corticosteroids: Metered Dose Inhalers CLOSED CLASS			
Arnuity[™] Ellipta[®] Asmanex Twisthaler Asmanex HFA[®]		<i>Aerospan[™]</i> <i>Alvesco[®]</i>	

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QVAR® Redihaler		fluticasone HFA (gen Flovent® HFA) fluticasone Diskus (gen Flovent® Diskus)	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
Inhaled Corticosteroids: Nebulizer Solution CLOSED CLASS			
budesonide respules		Pulmicort® Respules	
Nasal Steroids			
Dymista® fluticasone Rx and OTC		azelastine/fluticasone nasal spray (generic for Dymista®) budesonide (generic for Rhinocort® Aqua) budesonide (generic Rhinocort® Allergy OTC) Children's Qnasl™ Flonase® Flonase Sensimist (OTC) flunisolide ipratropium bromide spray mometasone (generic Nasonex®) Nasonex OTC Omnaris® Qnasl™ Rhinocort Aqua® Rhinocort® Allergy OTC Ryaltris® Ticanase® triamcinolone OTC triamcinolone acetanide Veramyst® Xhance™	

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Preferred Agents	Non-Preferred Agents	SA Criteria
Epinephrine, Self-Administered		
epinephrine 0.15 mg & 0.3 mg (authorized generic EpiPen® & EpiPen® Jr) Epipen® Epipen® Jr	Auvi-Q® epinephrine 0.15 mg & 0.3mg (generic EpiPen® and EpiPen® Jr) epinephrine 0.15mg & 0.3mg (generic Adrenaclick) Neffy® Symjepi™ (epinephrine)	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
*♦ Immunomodulators, Asthma CLOSED CLASS		
Fasenra Pen Fasenra Syringe Xolair Autoinjector Xolair Syringe Xolair Vial	Cinqair Nucala Auto-Injector Nucala Syringe Nucala Vial Tezspire Pen Tezspire Syringe	LENGTH OF AUTHORIZATIONS: see fax forms Routine PDL edits plus *♦ All Agents require the submission of a Clinical SA. Refer to fax forms below: Cinqair SA Form Fasenra SA Form Nucala SA Form Tezspire SA Form Xolair SA Form
Intranasal Antihistamines		
azelastine 0.1%	azelastine 0.15% nasal spray	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
Leukotriene Receptor Antagonists		
montelukast tabs/chewable tabs	Accolate® Singulair® tabs/chew tabs/ granules montelukast granules zafirlukast Zyflo™ zileuton ER	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits