



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

1/1/24

Virginia Medicaid's Pharmacy Benefits Management System

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

General Information:

- Virginia Medicaid's 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.

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



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
Analgesics		
*♦ Opioids – Long Acting (LAO)		
Preferred (Sch III-VI)	Non-Preferred	<p>*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)</p> <p>LENGTH OF AUTHORIZATIONS</p> <ul style="list-style-type: none"> 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).
<p>Butrans® (buprenorphine) Transdermal Patch</p>	<p><i>Belbuca</i> (buprenorphine buccal film)</p> <p>buprenorphine (gen Butrans®)</p> <p><i>buprenorphine</i> (gen <i>Belbuca</i>)</p> <p><i>ConZip®</i> (tramadol ER)</p> <p><i>Ryzolt™</i> (tramadol ER)</p> <p><i>tramadol ER</i></p> <p><i>Ultram ER®</i> (tramadol ER)</p>	
Preferred (Sch II)	Non-Preferred	<p>Daily dose limits have been established for all LAO. Quantity limits can be found at: Daily Dose Limits for Short & Long Acting Opioids</p>
<p>fentanyl 12, 25, 50, 75 & 100 mcg patches</p> <p>morphine sulfate ER tab</p>	<p><i>Arymo™ER</i></p> <p><i>Embeda</i></p> <p><i>Exalgo®</i></p> <p><i>fentanyl 37.5 mcg, 62.5 mcg, and 87.5 mcg patches</i></p> <p><i>hydromorphone ER</i></p> <p><i>Hysingla ER™</i></p> <p><i>Kadian® ER</i></p> <p><i>Morphabond™ ER</i></p> <p><i>morphine ER cap (generic Avinza®)</i></p> <p><i>morphine ER cap (generic Kadian®)</i></p> <p><i>MS Contin®</i></p> <p><i>Nucynta® ER</i></p> <p><i>Oramorph® SR®</i></p> <p><i>oxycodone-long acting</i></p> <p><i>OxyContin®</i></p> <p><i>oxymorphone ER</i></p> <p><i>Xartemis™ XR</i></p> <p><i>Xtampza ER®</i></p> <p><i>Zohydro ER™</i></p>	



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

1/1/24

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*Methadone Drugs		
	<i>Dolophine[®]</i> <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		
*Transmucosal Immediate Release Fentanyl		LENGTH OF AUTHORIZATIONS:
	<i>Actiq[®]</i> <i>Fentora[®]</i> <i>fentanyl citrate</i> <i>Lazanda[®]</i> <i>Subsys[®]</i>	<ul style="list-style-type: none"> 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).
Short-Acting Opioids		
codeine/APAP hydrocodone/APAP hydrocodone/ibuprofen hydromorphone morphine IR oxycodone IR oxycodone/APAP tramadol HCl 50mg tramadol HCl/APAP	<i>Abstral[®]</i> <i>Apadaz[™]</i> <i>codeine tab/soln</i> <i>butalbital comp with codeine</i> <i>butalbital/caffeine/APAP</i> <i>w/codeine</i> <i>butorphanol tartrate nasal</i> <i>dihydrocodeine/APAP/caffeine</i> <i>dihydrocodeine/ASA/caffeine</i> <i>hydromorphone liq/supp</i> <i>mepidine tab</i> <i>morphine supp</i> <i>Nucynta[®]</i> <i>Oxaydo[®]</i> <i>oxycodone/APAP(gen</i> <i>PrimLev[™])</i> <i>oxycodone conc</i> <i>oxycodone oral syringe</i> <i>oxycodone/ASA</i> <i>oxycodone/ibuprofen</i> <i>oxymorphone HCl</i>	*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)



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

1/1/24

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	Panlor [®] pentazocine/naloxone PrimLev [™] RoxyBond [™] Seglentis [®] tramadol 100 Mg tramadol soln (AG for Qdolo [®]) Ultracet [®] Ultram [®] Zamicet [®] soln																											
♦ Opioid Dependency CLOSED CLASS		*All Buprenorphine Containing Drugs (non-preferred) require submission of Clinical SA. Refer (Oral Buprenorphine SA Form)																										
*buprenorphine SL *buprenorphine/naloxone tab SL *Suboxone [®] film *Sublocade [™] SQ Kloxxado [™] Spray naloxone syringe & vial naloxone nasal spray naloxone nasal spray OTC Naloxone Carpuject naltrexone tab Narcan [®] Nasal Spray Vivitrol [®] Zimhi [™]	Brixadi[™] *buprenorphine/naloxone film SL *Probuphine [®] implant *Zubsolv [™] **Lucemyra [®]	**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 border="1" data-bbox="1003 987 1623 1398"> <tbody> <tr><td>buprenorphine SL tab 2mg</td><td>3/day</td></tr> <tr><td>buprenorphine SL tab 8mg</td><td>2/day</td></tr> <tr><td>buprenorphine/naloxone SL tab 2-0.5mg</td><td>3/day</td></tr> <tr><td>buprenorphine/naloxone SL tab 8-2mg</td><td>3/day</td></tr> <tr><td>buprenorphine/naloxone SL film 2-0.5mg</td><td>3/day</td></tr> <tr><td>buprenorphine/naloxone SL film 4-1mg</td><td>1/day</td></tr> <tr><td>buprenorphine/naloxone SL film 8-2mg</td><td>3/day</td></tr> <tr><td>Suboxone[®] SL film 2-0.5mg</td><td>3/day</td></tr> <tr><td>Suboxone[®] SL film 4-1mg</td><td>1/day</td></tr> <tr><td>Suboxone[®] SL film 8-2mg</td><td>3/day</td></tr> <tr><td>Suboxone[®] SL film 12-3mg</td><td>2/day</td></tr> <tr><td>Zubsolv[™] SL tab 0.7-0.18 mg</td><td>2/day</td></tr> <tr><td>Zubsolv[™] SL tab 1.4-0.36mg</td><td>2/day</td></tr> </tbody> </table>	buprenorphine SL tab 2mg	3/day	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
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1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria								
		<table border="1"> <tr> <td>Zubsolv™ SL tab 2.9–0.71mg</td> <td>2/day</td> </tr> <tr> <td>Zubsolv™ SL tab 5.7–1.4mg</td> <td>2/day</td> </tr> <tr> <td>Zubsolv™ SL tab 8.6–2.1mg</td> <td>2/day</td> </tr> <tr> <td>Zubsolv™ SL tab 11.4–2.9mg</td> <td>2/day</td> </tr> </table>	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
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 (OTC & Rx) Infant's ibuprofen drops meloxicam tab naproxen tab naproxen sodium (OTC) naproxen EC (Rx) sulindac	<i>Anaprox® IR & DS®</i> <i>Advil®</i> <i>Aleve®</i> <i>Arthrotec®</i> <i>Cataflam®</i> <i>*Celebrex® & *celecoxib</i> <i>Daypro®</i> <i>diclofenac potassium</i> <i>diclofenac potassium gen Zipsor™</i> <i>diclofenac sodium ER</i> <i>diclofenac sodium/misoprostol</i> <i>diflunisal</i> <i>Duexis®</i> <i>etodolac IR & SR</i> <i>Feldene®</i> <i>fenoprofen</i> <i>flurbiprofen</i> <i>ibuprofen tab chew OTC</i> <i>ibuprofen-famotidine (gen Duexis®)</i> <i>Indocin® supp</i> <i>indomethacin IR, SR & rectal</i> <i>ketoprofen IR & ER</i> <i>ketorolac</i> <i>ketorolac tromethamine (generic)</i> <i>Sprix® nasal spray)</i> <i>meclofenamate</i> <i>mefenamic</i>	Routine PDL edits plus *Step edit required for Celebrex® and celecoxib <ul style="list-style-type: none"> • 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® for which preferred drugs are not indicated. 								



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

1/1/24

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	<p><i>meloxicam susp</i> <i>meloxicam (generic Vivlodex™)</i> <i>Mobic® susp</i> <i>Motrin®</i> <i>nabumetone</i> <i>Nalfon®</i> <i>Naprelan®</i> <i>Naprosyn®</i> <i>naproxen CR (generic Naprelan®)</i> <i>naproxen-esomeprazole mag (generic Vimovo®)DR</i> <i>naproxen sodium (RX)</i> <i>naproxen susp</i> <i>oxaprozin</i> <i>piroxicam</i> <i>Ponstel®</i> <i>Prevacid Naprapac®</i> <i>Sprix® nasal spray</i> <i>Tivorbex™</i> <i>tolmetin sodium</i> <i>Vimovo®</i> <i>Vivlodex™</i> <i>Voltaren®XR</i> <i>Zipsor®</i> <i>Zorvolex™</i></p>	
<p>Topical NSAIDs diclofenac sodium 1% gel diclofenac sodium 1% gel (OTC)</p>	<p>**diclofenac sodium 3 % gel *diclofenac 2% topical sol gen Pennsaid® *Flector® patch (QL) Licart™ patch (diclofenac epolamine 0.013gm (QL) *Pennsaid® top soln, soln pkt & pump</p>	<p>LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus Clinical Criteria for <u>Non-Preferred Topical NSAIDs</u>; *Flector®, <u>diclofenac 2% topical sol, Pennsaid®, Vopac MDS, & Xrylix™ Kit:</u></p> <ul style="list-style-type: none"> 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). For example, a member who failed ibuprofen or naproxen will still need to try oral diclofenac for approval of Flector®.



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

1/1/24

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		*Vopac MDS *Voltaren® 1% gel *Voltaren® 1% gel (OTC) *Xrylix™ Kit	<ul style="list-style-type: none"> diclofenac 2% topical sol, Pennsaid®, Vopac MDS, and Xrylix™ Kit can only be approved for the FDA approved indication of osteoarthritis of the knee. <p>Quantity limit for Flector® and Licart™ patch = 30 patches per RX</p>
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)	<p>LENGTH OF AUTHORIZATIONS: 1 year</p> <p>Routine PDL edits plus</p> <p>**Tobi Podhaler®</p> <ul style="list-style-type: none"> Requires a clinical reason as to why one of the preferred tobramycin inhalation nebulizer solutions cannot be used <p>***Clinical Criteria for Arikayce®</p> <p>Duration of Approval: 12 months</p> <p>Initial Approval Criteria</p> <ul style="list-style-type: none"> Patient is ≥ 18 years of age; AND Diagnosis of Mycobacterium avium complex (MAC) lung disease as determined by the following: <ul style="list-style-type: none"> 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 Patient 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 Patient 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 <p>*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®.</p>



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
		<p>Quantity Limitations: Arikayce®= 590 mg/8.4 MI (28 vials)/28 days Each carton contains a 28-day supply of medication (28 vials) Bethkis® = 224MI (56 amps)/28 days Cayston® = 84 MI (56 amps)/28 days Kitabis™ Pak = 280MI (56 amps)/28 days Tobi Podhaler® = 224 capsule/28 day Tobi® inhalation neb = 280MI (56 amps)/28 days tobramycin nebs = 280MI (56 amps)/28 days</p>
Antifungals, Oral		
<p>fluconazole tab/susp griseofulvin susp nystatin tab/susp terbinafine tab</p>	<p>Ancobon® Brexafemme® clotrimazole (mucous mem) Cresemba® Diflucan® tab/susp flucytosine Gris-Peg® griseofulvin tab griseofulvin ultramicrosize itraconazole itraconazole soln (generic for Sporanox® soln) ketoconazole Lamisil® tab/granules Noxafil® Noxafil® Powdermix posaconazole tab (generic for Noxafil) *Sporanox® cap/soln Tolsura™ Vfend® tab/susp *Vivjoa®</p>	<p>LENGTH OF AUTHORIZATIONS: Duration of the prescription (up to 12 months)</p> <p>Routine PDL edits plus</p> <p>Clinical Criteria for all <u>Non-Preferred oral Antifungals</u>. Requires the submission of a Clinical SA. Refer to Antifungal Oral SA Form</p> <p>*Clinical Criteria for Vivjoa: Duration of approval: Date of service</p> <ul style="list-style-type: none"> • Patient 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 • Member has tried and failed or has a contraindication or intolerance to maintenance antifungal therapy with oral fluconazole. <p>Quantity limit: 18 tablets per treatment course</p>



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1/1/24

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		voriconazole tab & powder for susp	
Cephalosporins, Oral			
Second Generation Cephalosporins			LENGTH OF AUTHORIZATIONS: Date of service only; no refills.
cefaclor capsule	cefaclor ER		Routine PDL edits plus Clinical Criteria for <u>Non-Preferred Cephalosporins</u> <ul style="list-style-type: none"> • Infection caused by an organism resistant to preferred drugs, OR • A therapeutic failure to no less than a three-day trial of one preferred cephalosporin; OR • The member is completing a course of therapy with a non-preferred drug initiated in the hospital.
cefprozil tab/susp	cefaclor susp		
cefuroxime tab	Ceftin® tab/susp		
Third Generation Cephalosporins			
cefdinir cap/susp	Cedax® cap/susp ceftibuten cefditoren pivoxil cefixime suspension cefpodoxime proxetil cap/susp Spectracef® Suprax® chewable tab/cap/susp		
Macrolides, Oral			
Macrolides & Ketolides			LENGTH OF AUTHORIZATIONS: Date of service only; no refills
azithromycin pack/susp/tab	Biaxin® tab		Routine PDL edits plus Clinical Criteria for <u>Non-Preferred Macrolides and Ketolides</u> <ul style="list-style-type: none"> • Infection caused by an organism resistant to preferred drugs, OR • A therapeutic failure to no less than a three-day trial of one preferred drug within the same class; OR • The member is completing a course of therapy with a non-preferred drug which was initiated in the hospital.
clarithromycin tab/susp	clarithromycin ER		
E.E.S.® 200 susp	Eryped® 200 susp		
erythromycin base cap DR	Eryped® 400 susp		
erythromycin ethylsuccinate	Ery-tab®		
200mg susp	E.E.S.® 400 tab		
erythromycin stearate	Erythrocin® Stearate		
	erythromycin base tab		
	erythromycin ethylsuccinate		
	400mg tab(Generic E.E.S.® 400)		
	PCE®		
	Zithromax® pac/tab/susp		
	ZMAX® susp		
Otic			



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1/1/24

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Ciprodex[®] ciprofloxacin/dexamethasone (generic Ciprodex) ofloxacin neomycin/polymyxin/hc soln/sus		<i>Cetraxal[®]</i> <i>Cipro HC[®]</i> <i>ciprofloxacin and fluocinolone acetonide (generic Otovel)</i> <i>Otovel</i>	<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 Routine PDL edits plus: <u>Clinical Criteria for Non-Preferred Quinolones</u> <ul style="list-style-type: none"> • Infection caused by an organism resistant to preferred drugs; OR • A therapeutic failure to no less than a three-day trial of one preferred quinolone; OR • The member is completing a course of therapy with a non-preferred drug initiated in the hospital.
ciprofloxacin susp/tab	<i>Cipro[®] IR, XR & susp</i> <i>ciprofloxacin ER</i> <i>Noroxin[®]</i> <i>ofloxacin</i>		
Third Generation Quinolones			
levofloxacin tab	<i>Baxdela[™] tab</i> <i>Levaquin[®] tab/susp</i> <i>levofloxacin susp</i> <i>moxifloxacin</i>		
Topical Antibiotics			
mupirocin ointment	<i>*Altabax[™] (QL)</i> <i>Bactroban[®] cr/ointment</i> <i>Centany[®]</i> <i>Centany AT[®] Kit</i>	<u>LENGTH OF AUTHORIZATIONS:</u> Date of service only; no refills Routine PDL edits <i>*Quantity Limit = 15 grams per 34 days</i>	
Vaginal Antibiotics			
Cleocin[®] Ovules Clindesse[®] cr metronidazole gel Nuessa[™]	<i>Cleocin[®] cr</i> <i>clindamycin cr</i> <i>Metrogel[®]</i> <i>Vandazole[™] gel</i> <i>Xaciatto[™] gel</i>	<u>LENGTH OF AUTHORIZATIONS:</u> Date of Service Routine PDL edits	



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1/1/24

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Antivirals		
*Hepatitis C Agents		CLOSED CLASS
Interferon		LENGTH OF AUTHORIZATIONS:
	<i>Pegasys® syringe/vial</i>	*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		
sofosbuvir /velpatasvir (generic Epclusa®)	<i>Epclusa® Epclusa® Pellet packs Sovaldi® Vosevi™</i>	* Hepatitis C Therapies are limited to 3 one-month fills (total 84 day supply) with out SA
*NS5A, NS3/4A Inhibitor Combinations		<ul style="list-style-type: none"> • Mavyret™ • sofosbuvir /velpatasvir (generic Epclusa®)
Mavyret™ Mavyret™ Pellet packs	<i>Technivie™ Viekira Pak™ Viekira XR™ Zepatier®</i>	
*NS5B & Protease Inhibitor Combinations		
	<i>Harvoni® ledipasvir/sofosbuvir (generic Harvoni®)</i>	
Herpes Oral		
acyclovir cap/tab/susp famciclovir valacyclovir	<i>Famvir® Sitavig® buccal tab Valtrex® Zovirax® tab/susp</i>	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
Herpes Topical		
acyclovir oint docosanol	<i>Abreva OTC® acyclovir cr (generic Zovirax cr) Denavir® penciclovir Xerese® cr Zovirax® oint Zovirax® cr</i>	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits



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1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
HIV/AIDS CLOSED CLASS		
abacavir tab/soln abacavir-lamivudine tab abacavir-lamivudine-zidov tab Aprelude Aptivus [®] soln atazanavir sulfate cap Biktarvy [®] tab Cabenuva Cimduo [®] tab Complera [®] tab 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 Fuzeon [®] Vial Genvoya [®] tab Intelence [®] tab Invirase [®] tab Isentress [®] chew tab/ tab/powder pack Juluca [®] tab lamivudine tab/soln lamivudine-zidovudine tab Lexiva [®] tab lopinavir-ritonavir soln/tab	Aptivus [®] cap Atripla [®] tab Combivir [®] tab darunavir didanosine cap DR efavirenz cap efavirenz-lamiv-tenof tab Epivir [®] tab Epzicom [®] tab Kaletra [®] tab/soln Lexiva [®] susp Norvir [®] tab Retrovir [®] syrup Reyataz [®] cap stavudine cap Sustiva [®] cap Trizivir [®] tab Trogarzo [®] (intraven) Viracept [®] tab Viread [®] tab Ziagen [®] tab	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits Quantity limits see list here HIV Quantity Limits



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
<p> Maraviroc nevirapine IR tab/ ER tab/susp Norvir[®] soln/powder pack Odefsey[®] tab Pifeltro[™] tab Prezcobix[®] tab Prezista[®] tab/susp Retrovir[®] cap/vial Reyataz[®] powder pack Rilpivirine ER vial ritonavir tab/soln Rukobia[™] ER tab Selzentry[®] tab/soln Stribild[®] tab Sunlenca[®] tab/vial Sustiva[®] tab Symfi[®] and Symfi Lo[®] tab Symtuza[®] tab Temixys[™] tab tenofovir disoproxil fum tab Tivicay[®] tab/tab for susp Triumeq[®] tab Triumeq[®] PD Truvada[®] tab Tybost[®] tab Viramune[®] tab/susp Viread[®] powder Vocabria Ziagen[®] soln zidovudine tab/cap/syrup </p>		
Influenza		
<p>oseltamivir susp/ cap</p>	<p> <i>Flumadine[®] tab</i> <i>rimantadine</i> <i>Relenza Disk[®]</i> <i>Tamiflu[®] susp/cap</i> </p>	<p> <u>LENGTH OF AUTHORIZATIONS:</u> Date of service only Routine PDL edits </p>



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
<i>Xofluza™</i>		
Blood Modifiers		
Bile Salts		
ursodiol cap, tab	<i>*Bilvay™</i> <i>Chenodal®</i> <i>Cholbam®</i> <i>**Livmarli™</i> <i>Ocaliva®</i> <i>Reltone®</i> <i>Urso® & Urso® Forte tab</i>	<p>LENGTH OF AUTHORIZATIONS: 1 year</p> <p>Routine PDL edits</p> <p>*Livmarli™ (PDL criteria do not apply)</p> <ul style="list-style-type: none"> Must have a confirmed diagnosis of cholestatic pruritus in patients 3 months of age and older with Alagille syndrome (ALGS) <p>** Bilvay™ (PDL criteria do not apply)</p> <ul style="list-style-type: none"> Must have a confirmed diagnosis of cholestatic pruritus in: <ul style="list-style-type: none"> Patients 12 months of age and older with Alagille syndrome (ALGS) Patients 3 months of age and older with progressive familial intrahepatic cholestasis (PFIC)
Hemophilia Treatment (Closed Class)		
Factor VIII Products		
Advate® Adynovate® Afstyla® Alphanate® Altuviiio™ Eloctate® Esperoct® Hemofil-M® Humate-P® Jivi®		<p>LENGTH OF AUTHORIZATIONS: N/A</p> <p>All products are preferred without EDITs</p>



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

1/1/24

<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
Koate-DVI[®] Kogenate FS[®] Kovaltry[®] Novoeight[®] Nuwiq[®] Obizur[®] Recombinate[®] Xyntha[®] Kit & Solofuse Syr		
Factor IX Products		
AlphaNine SD[®] Alprolix[®] BeneFIX[®] Idelvion[®] Ixinity[®] Profilnine SD[®] Rebinyn[®] 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[®]		
Von Willebrand		



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

1/1/24

Preferred Agents		Non-Preferred Agents	SA Criteria
Vonvendi® Wilate®			
Phosphate Binders			
calcium acetate 667mg cap calcium acetate 668 mg sevelamer carbonate tab	<i>Auryxia™</i> <i>calcium acetate 667mg tab</i> <i>Calphron</i> <i>Eliphos</i> <i>Fosrenol® chewable tab</i> <i>lanthanum carbonate chewable tab</i> <i>Phoslyra® powder, tab</i> <i>Renagel®</i> <i>Renvela®</i> <i>sevelamer carbonate powder packet</i> <i>sevelamer HCL (gen Renagel)</i> <i>Velphoro® chewable tab</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits	
Sickle Cell (CLOSED CLASS)			
Droxia Endari Oxbryta	<i>Adakveo IV</i> <i>Siklos</i>	*Non-preferred agents require the submission of a Clinical SA. Preferred agents do not require submission of any SA. Refer to Sickle Cell SA Form	
Bone Resorption Suppression and Related Agents			
Bisphosphonates			
alendronate tab ibandronate	<i>Actonel®</i> <i>alendronate soln</i> <i>Atelvia DR®</i> <i>Boniva®</i> <i>Binosto™</i> <i>etidronate</i> <i>Fosamax® tab & Fosamax® plusD</i> <i>risedronate DR</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits	



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

1/1/24

Preferred Agents		Non-Preferred Agents	SA Criteria
Calcitonins			
calcitonin-salmon nasal	Miacalcin [®] spray calcitonin-salmon (gen Miacalcin [®])	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits	
Others			
raloxifene	Evista [®] *Forteo [®] teriparatide *Tymlos [™]	LENGTH OF AUTHORIZATIONS: Initial approval will be for 1 year Routine PDL edits for Evista[®] *♦ Clinical SA must be completed for (Forteo[®] OR Tymlos[™] SA Form)	
Cardiac			
Anticoagulants		CLOSED CLASS	
Low Molecular Weight Heparin includes FactorXA Inhibitor		LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus	
enoxaparin (generic Lovenox [®])	Arixtra [®] fondaparinux Fragmin [®] syringe & vial Lovenox [®]		
Oral Anticoagulants		Clinical Criteria for Savaysa[™]	
Eliquis [™] Eliquis [™] Dose Pack Jantoven [®] Pradaxa [®] warfarin Xarelto [®] & Starter Pack, susp	*Savaysa [™] Pradaxa [®] Pellet Pack	<ul style="list-style-type: none"> • Diagnosis of: Non-valvular Atrial Fibrillation, OR • Deep vein thrombosis, OR • Pulmonary embolism; AND • Documentation that CrCl is NOT ≥ 95 mL/min calculated by Cockcroft-Gault equation 	
Antihypertensive Agents			
ACE Inhibitors		LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits	
benazepril enalapril lisinopril ramipril quinapril	Accupril [®] Altace [®] captopril Epaned [™] soln enalapril soln		



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	<i>fosinopril</i> <i>Lotensin®</i> <i>Mavik®</i> <i>moexipril</i> <i>Monopril®</i> <i>perindopril</i> <i>Qbrelis™</i> <i>trandolapril</i> <i>Univasc®</i> <i>Vasotec®</i> <i>Zestril®</i>	
ACE Inhibitors + Calcium Channel Blocker Combinations		
amlodipine/benazepril	<i>Lotrel®</i> <i>Tarka®</i> <i>trandolapril-verapamil ER</i>	
ACE Inhibitors + Diuretic Combinations		
benazepril/HCTZ lisinopril/HCTZ enalapril/HCTZ	<i>Accuretic®</i> <i>captopril/HCTZ</i> <i>fosinopril/HCTZ</i> <i>Lotensin HCT®</i> <i>moexipril/HCTZ</i> <i>quinapril /HCTZ</i> <i>quinapril-HCTZ (gen Accuretic)</i> <i>Vaseretic®</i> <i>Zestoretic®</i>	
Angiotensin Receptor Blockers		
*Entresto™ (QL) irbesartan losartan olmesartan	<i>Atacand®</i> <i>Avapro®</i> <i>Benicar®</i> <i>candesartan</i>	<p><u>LENGTH OF AUTHORIZATIONS:</u> 1 year</p> <p>Routine PDL edits plus</p> <p><i>Quantity Limit = 2 per day for Entresto™</i></p>



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

1/1/24

<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
valsartan	<i>Cozaar[®]</i> <i>Diovan[®]</i> <i>Edarbi[®]</i> <i>eprosartan mesylate</i> <i>Micardis[®]</i> <i>Teveten[®]</i> <i>valsartan solution</i>	
Angiotensin Receptor Blockers + Calcium Channel Blocker Combinations		
amlodipine/olmesartan amlodipine/valsartan	<i>Azor[®]</i> <i>amlodipine/olmesartan/HCTZ</i> <i>amlodipine/valsartan/HCTZ</i> <i>Exforge[®] & Exforge[®] HCT</i> <i>Tribenzor[®]</i>	
Angiotensin Receptor Blockers + Diuretic Combinations		
irbesartan/HCTZ losartan/HCTZ olmesartan/HCTZ valsartan/HCTZ	<i>Atacand HCT[®]</i> <i>Avalide[®]</i> <i>Benicar HCT[®]</i> <i>candesartan/HCTZ</i> <i>Diovan HCT[®]</i> <i>Edarbyclor[®]</i> <i>Hyzaar[®]</i> <i>Micardis HCT[®]</i> <i>telmisartan/HCTZ</i> <i>Teveten HCT[®]</i>	
Antihypertensives, Sympatholytics		
Catapres[®]-TTS clonidine (transdermal) clonidine tab guanfacine methyldopa	<i>Clorpres[®]</i> <i>methyldopa/HCTZ</i> <i>Tenex[®]</i>	



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
Beta Blockers		*Clinical Criteria for Hemangeol™
atenolol bisoprolol carvedilol labetalol metoprolol tartrate metoprolol succinate propranolol tab & ER/soln Sorine® sotalol AF sotalol HCL	<i>acebutolol</i> <i>Betapace® IR & AF®</i> <i>betaxolol</i> <i>Carvedilol ER</i> <i>Coreg® IR & CR®</i> <i>Corgard®</i> <i>*Hemangeol™</i> <i>Inderal® XL</i> <i>Innopran® XL</i> <i>Kaspargo™ Sprinkle</i> <i>LevatoI®</i> <i>Lopressor®</i> <i>nebivolol</i> <i>nadolol</i> <i>pindolol</i> <i>propranolol LA</i> <i>Sectral®</i> <i>Sotylize™</i> <i>Tenormin®</i> <i>timolol maleate</i> <i>Toprol XL®</i> <i>Trandate®</i> <i>Zebeta®</i>	<ul style="list-style-type: none"> • Diagnosis of proliferating infantile hemangioma requiring systemic therapy
Beta Blockers + Diuretic Combinations		
atenolol/chlorthalidone bisoprolol/HCTZ	<i>Corzide®</i> <i>Dutoprol®</i> <i>Lopressor HCT®</i> <i>metoprolol/HCTZ</i> <i>nadolol/bendroflumethiazide</i> <i>propranolol/HCTZ</i> <i>Tenoretic®</i> <i>Ziac®</i>	



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
Calcium Channel Blockers –Dihydropyridine		
Afeditab CR[®] amlodipine Nifedical XL[®] nifedipine nifedipine ER	<i>Adalat CC[®]</i> <i>Conjupri[®]</i> <i>Consensi[®] (amlodipine/celecoxib)</i> <i>felodipine ER</i> <i>isradipine</i> <i>Katerzia[™]oral suspension</i> <i>levamlodipine (generic</i> <i>Conjupri[®])</i> <i>nisoldipine</i> <i>nicardipine</i> <i>Norliqva[®]</i> <i>Norvasc[®]</i> <i>Procardia XL[®]</i> <i>Sular[®]</i>	
Calcium Channel Blockers- Non-Dihydropyridine		
Cartia XT[®] diltiazem IR, ER q12hr & 24hr Taztia XT[®] verapamil tab IR & ER	<i>Calan[®] IR & SR</i> <i>Cardizem[®] IR, CD & LA</i> <i>Isoptin SR[®]</i> <i>diltiazem LA</i> <i>Matzim LA</i> <i>Tiazac[®]</i> <i>verapamil 360 cap</i> <i>verapamil ER cap</i> <i>Verelan[®] & Verelan PM[®]</i>	
Direct Renin Inhibitors (includes combination)		
	<i>aliskiren 150 & 300mg (generic for Tekturna)</i> <i>Tekamlo[®]</i> <i>Tekturna[®]</i> <i>Tekturna HCT[®]</i>	



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	<i>Twynsta[®]</i> <i>telmisartan/amlodipine</i>	
Lipotropics		
Bile Acid Sequestrants		LENGTH OF AUTHORIZATIONS: 1 year
cholestyramine powder reg & light colestipol tab Prevalite[®]	<i>Colestid[®] granule/packet/tab</i> <i>colesevelam tab and Pkt (generic)</i> <i>Welchol</i> <i>colestipol HCl granules</i> <i>Questran[®] powder/powder Light</i> <i>Welchol[®] pack, tab</i>	Routine PDL edits plus
Cholesterol Absorption Inhibitor (CAI) and /or ACL Inhibitor (adenosine triphosphate citrate lyase)		Routine PDL Plus ♦ Clinical criteria Nexletol[™] or Nexlizet[™]
ezetimibe	<i>Nexletol[®] (bempedoic acid)</i> <i>Nexlizet[®] (bempedoic acid/ezetimibe)</i> <i>Zetia[®]</i>	Initial Approval Criteria <ul style="list-style-type: none"> • Patient is ≥ 18 years of age; AND • Patient has diagnosis of heterozygous familial hypercholesterolemia (HeFH) or established atherosclerotic cardiovascular disease (ASCVD); AND • Patient has failed to achieve a target LDL-C despite physician attestation that the patient is adherent to maximally tolerated doses of statins prior to the lipid panel demonstrating suboptimal reduction; AND • Patient can be classified into ONE of the following risk factor groups: <ul style="list-style-type: none"> ○ 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



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
		<ul style="list-style-type: none"> Therapy will not be used with concurrent doses of simvastatin > 20 mg or pravastatin > 40mg; <p>Renewal Criteria</p> <ul style="list-style-type: none"> Laboratory analyses demonstrate a reduction in LDL-C when compared to the baseline values (prior to initiating bempedoic acid or bempedoic acid/ezetimibe); AND Patient has shown continued adherence to maximally tolerated statin dosage
Fibric Acid Derivatives		
fenofibrate (generic <i>Tricor</i> [®] 48mg 145mg) gemfibrozil	<i>Antara</i> [®] <i>fenofibrate</i> (generics for <i>Antara</i> [®] , <i>Fenoglide</i> [®] & <i>Lipofen</i> [®]) <i>fenofibrate</i> (generics for <i>Triglide</i> [®]) <i>fenofibric acid</i> <i>Fenoglide</i> [®] <i>Fibricor</i> [®] <i>Lipofen</i> [®] <i>Lofibra</i> [®] <i>Lopid</i> [®] <i>Tricor</i> [®] <i>Triglide</i> [®] <i>Trilipix</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>Ezallor Sprinkle</i> (rosuvastatin) <i>Lipitor</i> [®] <i>Livalo</i> [®] <i>rosuvastatin-ezetimibe</i>	



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

1/1/24

Preferred Agents		Non-Preferred Agents	SA Criteria
		<i>simvastatin soln (generic Flolipid)</i> <i>simvastatin/ezetimibe</i> <i>Vytorin[®]</i> <i>Zocor[®]</i> <i>ZypitamagTM</i>	
HMG CoA Reductase Inhibitors and Combinations (Statins)			
lovastatin pravastatin		<i>Atoprev[®]</i> <i>fluvastatin</i> <i>Lescol[®] and Lescol XL[®]</i> <i>Mevacor[®]</i> <i>Pravachol[®]</i>	
Microsomal Triglyceride Transfer Protein Inhibitor			
		* <i>JuxtapidTM</i>	* ♦ Clinical Criteria for JuxtapidTM. Refer to JuxtapidTM SA Fax Form
Niacin Derivatives			
niacin ER		<i>Niaspan[®]</i> <i>Niacor[®]</i>	
Omega 3 Fatty Acid Agent			
*** <i>omega-3 acid ethyl esters (ST)</i> Omega-3 OTC Vascepa[®]		<i>icosapent ethyl (generic Vascepa[®])</i> *** <i>Lovaza[®] (ST)</i>	*** ♦ Clinical Criteria for Lovaza[®] and omega-3 acid ethyl esters <ul style="list-style-type: none"> • Step edit requires trial and failure of any other lipotropic; OR • Documented high triglycerides of ≥ 500 mg/dL.
*Proprotein Convertase Subtilisin Kexin Type 9 (PCSK9) Inhibitors			
		<i>Leqvio[®]</i> <i>Praluent[®]</i> <i>Repatha[®]</i>	LENGTH OF AUTHORIZATIONS: Three months for initial approval; six months for renewal * ♦ ALL PCSK9 Inhibitors require the submission of a Clinical SA. Refer to Lipotropics, Other SA Form
Platelet Inhibitors			
Brilinta [®] clopidogrel dipyridamole prasugrel (generic Effient [®])		* <i>Aggrenox[®]</i> * <i>ASA/dipyridamole</i> ** <i>ASA/omeprazole (generic Yosprala[®])</i> ** <i>Durlaza ERTM</i> <i>Effient[®]</i>	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus Clinical Criteria for Select <u>Non-Preferred Platelet Inhibitors</u> * <u>Aggrenox[®] & ASA/dipyridamole</u>



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

1/1/24

Preferred Agents		Non-Preferred Agents	SA Criteria
		<i>Persantine[®]</i> <i>Plavix[®]</i> ** <i>Yosprala[®] Tab</i> *** <i>ZontivityTM</i>	<ul style="list-style-type: none"> Aspirin and dipyridamole are covered as separate drugs without SA; clinical reason as to why the individual drugs cannot be used separately. <p>**<u>Durlaza ERTM</u> & *<u>Yosprala[®] Tab</u></p> <ul style="list-style-type: none"> Aspirin is covered without SA; clinical reason as to why aspirin cannot be used. <p>*** <u>ZontivityTM</u></p> <ul style="list-style-type: none"> Diagnosis of MI (myocardial infarction) or PAD (peripheral arterial disease); AND Members must not have a history of stroke, TIA, ICH, GI bleed and peptic ulcer; AND Must have concomitant therapy with clopidogrel, unless member has a contraindication to clopidogrel in which case member must have concomitant therapy with aspirin; AND Member is 18 years of age or older; AND Prescribed by or in consultation with a cardiologist.
*Pulmonary Arterial Hypertension Agents			
Inhaled Prostacyclin Analogues			LENGTH OF AUTHORIZATIONS: 1 year
Ventavis[®]		<i>Tyvaso[®]</i> <i>Tyvaso DPITM (treprostinil)</i>	
Oral Endothelin Receptor Antagonist			Routine PDL edits plus
ambrisentan (generic Letairis) 5 & 10mg <u>Tracleer[®] tab</u>		<u>bosentan (generic Tracleer[®])</u> <i>Letairis[®] 5 & 10 mg</i> <i>Opsumit[®]</i> <i>Tracleer[®] susp</i>	
*Phosphodiesterase 5 Inhibitors (PDE-5)			*♦ <u>Clinical Criteria for all preferred and non-preferred PDE-5</u>
Alyq (tadalafil) sildenafil tab/susp tadalafil(generic Adcirca[®])		<i>Adcirca[®]</i> <u>Liqrev[®]</u> <i>Revatio[®] tab/inj/susp</i> <i>Tadliq[®] Suspension</i>	<ul style="list-style-type: none"> Diagnosis of pulmonary hypertension in members >18 years is required; 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[®]
Prostacyclin Vasodilator and Receptor Agonist			
		<i>OrenitramTM</i> <u>OrenitramTM titration kit</u> <i>Uptravi[®]</i>	
Soluble Guanylate Cyclase Stimulators			



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

1/1/24

Preferred Agents		Non-Preferred Agents	SA Criteria
		<i>Adempas</i> [®]	
Central Nervous System			
Alzheimer's Agents			
Cholinesterase Inhibitors		<i>Adlarity</i> [®] <i>Aricept</i> [®] ODT, tab <i>Exelon</i> [®] cap & <i>Exelon</i> [®] (transderm) <i>galantamine IR, ER tab/soln</i> <i>Namzaric</i> [®] (donepezil/memantine) <i>Razadyne</i> [®] IR, ER <i>rivastigmine cap</i>	LENGTH OF AUTHORIZATIONS: Length of prescription (up to 3 months) Routine PDL edits
donepezil OTD & tab			
rivastigmine (transderm)			
NMDA Receptor Antagonist		<i>memantine Dose Pack, soln</i> <i>memantine ER</i> <i>Namenda</i> [®] Dose Pack/XR tab <i>Namenda</i> [®] tab	
memantine tab			
Anticonvulsants		CLOSED CLASS	
Barbiturates		<i>Mysoline</i> [®]	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL
phenobarbital elixir/tab			
primidone			
Benzodiazepines		<i>clonazepam ODT</i> <i>clorazepate</i> <i>Klonopin Tab</i> <i>Onfi</i> [®] susp/tab <i>Sympazan</i> [™] (clobazam) <i>Tranxene</i> [®]	* ♦ Clinical Criteria for Nayzilam [®] (reviewed electronically, AutoPA) <ul style="list-style-type: none"> Member is 12 years of age or older; AND Diagnosis of acute treatment of intermittent, stereotypic episodes of frequent seizure activity (i.e., seizure clusters, acute repetitive seizures) that are distinct from a patient's usual seizure pattern.
clobazam tab/susp			
clonazepam tab			
Diastat [®] rectal			
Diastat [®] AcuDial [™] rectal			
diazepam rectal & Device			
rectal			
Nayzilam [®]			
Valtoco [®] Nasal			
Cannabidiol			

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

1/1/24

Preferred Agents		Non-Preferred Agents	SA Criteria
*Epidioex [®] (cannabidiol)			*♦ Clinical Criteria for Epidiolex[®] (reviewed electronically, AutoPA) Duration of Approval: 1 year <ul style="list-style-type: none"> • Patient must be ≥ 1 years of age; AND • Patient has been diagnosed with: <ul style="list-style-type: none"> ○ Epilepsy and recurrent seizures including Lennox-Gastaut syndrome, ○ Dravet syndrome, or tuberous sclerosis complex
Carbamazepine Derivatives			
carbamazepine chewable tab/susp/tab Carbatrol[®] Epitol oxcarbazepine tab Tegretol[®]XR Trileptal[®] susp	<i>Aptiom[®]</i> <i>carbamazepine ER</i> <i>carbamazepine XR</i> <i>Equetro[®] cap</i> <i>OxtellarTM XR</i> <i>oxcarbazepine susp</i> <i>Tegretol[®] susp/tab</i> <i>Trileptal[®] tab</i>		
Hydantoins			
Dilantin 30mg phenytoin cap/chew tab/ susp phenytoin ext cap	<i>Dilantin[®] cap & Infatab, susp</i> <i>Peganone[®]</i> <i>Phenytek[®]</i>		
Succinimides			
ethosuximide cap/syrup	<i>Celontin[®]</i> <i>Zarontin[®] cap/syrup</i>		
Valproic Acid and Derivatives			
divalproex tab/sprinkle divalproex ER valproic acid cap, soln	<i>Depakote[®] ER & sprinkle</i>		
Other Anticonvulsants			
Gabitril[®] lacosamide soln/tab (gen Vimpat [®]) Lamictal[®] ODT dose pk lamotrigine ODT lamotrigine tab	<i>Banzel[®] susp/tab</i> <i>Briviact[®] tab/soln</i> <i>Diacomit[®] cap/powder pack</i> <i>Elepsia[®]XR tab</i> <i>EprontiaTM</i> <i>felbamate susp/tab</i>	*♦ Clinical Criteria for Fintepla[®] <ul style="list-style-type: none"> • Patient is at least two years of age or older; AND • Patient must have a diagnosis of Dravet syndrome or Lennox-Gastaut syndrome 	



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
<p>lamotrigine chew tab lamotrigine XR levetiracetam soln/tab levetiracetam ER rowepra (generic levetiracetam) subvenite tab (generic lamotrigine) tiagabine topiramate tab/sprinkle cap zonisamide cap</p>	<p>Felbatol[®] susp/tab * Fintepla[®] Fycompa[®] susp/tab Keppra[®] soln/tab Keppra[®] XR lacosamide solution unit dose Lamictal[®] XR & XR dose pk Lamictal[®] tab/dose pk Lamictal[®] ODT lamotrigine tab dose pk lamotrigine ODT dose pk rufinamide susp/tab (generic Banzel[®]) Qudexy[™] XR Sabril[®] powder pack/tab Spritam[®] subvenite dose pk Topamax[®] tab/sprinkle topiramate ER cap Trokendi[™] XR vigabatrin tab/ powder pack (generic Sabril[®]) Vimpat[®] soln/tab Xcopri[®] Zonisade[™] Solution **Ztalmly[®]</p>	<p>** ♦ Clinical Criteria for Ztalmly:</p> <ul style="list-style-type: none"> • Initial: <ul style="list-style-type: none"> ○ Patient is two years of age or older; AND ○ 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: <ul style="list-style-type: none"> ○ 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: 1 year
<p>bupropion IR, SR & XL desvenlafaxine ER tab (generic for Pristiq[®]) mirtazapine ODT/tab trazodone tab venlafaxine IR tab & ER cap vilazodone tab</p>	<p>Aplenzin[®] Auvelity[™] Brintellix[®] bupropion XL (gen Forfivo[®] XL) desvenlafaxine ER tab (gen Khedzla[™]) Effexor[®] XR</p>	<p>Routine PDL edits</p>



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	<p><i>Emsam[®] transdermal</i> <i>Fetzima[®]</i> <i>Forfivo[®] XL</i> <i>KhedezlaTM</i> <i>Marplan[®]</i> <i>Nardil[®]</i> <i>nefazodone</i> <i>Parnate[®]</i> <i>phenelzine</i> <i>Pristiq[®]</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[®] IR, SR & XL</i></p>	
SSRI		
<p>citalopram soln/tab escitalopram tab fluoxetine cap/soln fluvoxamine tab paroxetine tab sertraline conc, soln, tab</p>	<p><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>Luvox[®] CR</i> <i>paroxetine CR</i> <i>paroxetine susp</i> <i>Paxil[®] tab/susp & Paxil[®] CR</i> <i>Pexeva[®]</i> <i>Prozac[®] cap/weekly</i> <i>Sarafem[®]</i> <i>sertraline cap</i> <i>Zoloft[®] conc/tab</i></p>	
Antimigraine Agents		



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
<p>sumatriptan succinate tab cartridge/vial/pen Imitrex® nasal rizatriptan tab/MLT</p>	<p>almotriptan Alsuma® Amerge® Axert® Cambia® eletriptan (generic Relpax®) 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®) Tosymra Treximet® Zembrace™ SymTouch™ zolmitriptan nasal Zomig® tab/nasal spray/ZMT</p>	<p>LENGTH OF AUTHORIZATIONS: 1 year</p> <p>Routine PDL edi</p>
<p>Antimigraine Agents, Others CLOSED CLASS Calcitonin Gene-related Peptide Antagonist (CGRP)</p>		<p>*♦ All Nonpreferred CGRPs require the submission of a Clinical SA. Refer to fax form Antimigraine Agents, Others SA Form</p>
<p>Preventive Treatment</p>		<p>Preferred Agents *step edit required</p>
<p>Aimovig™ Ajovy® Autoinjector & Syringe Ajovy Autoinjector 3-Pk Emgality™ Syringe Emgality™ Pen Nurtec™ ODT</p>	<p>Emgality™ 100MG Syringe Qulipta™</p>	<p>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 = 1 mL per 30 days Nurtec® ODT = 18 tabs per 30 days Qulipta™ = 1 tab per day</p>



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
Acute treatment of migraine		Preferred Agents (No SA with trial of 2 generic triptans)
Nurtec™ ODT Ubrelvy®	Reyvow® Trudhesa™ Nasal Spray Zavzpret™	Quantity limit for acute treatment Nurtec® ODT = 18 tabs per 30 days Reyvow® = 50mg or 100mg - 8 tabs per strength per 30 days Ubrelvy® = 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 quetiapine fumarate ER quetiapine tab risperidone ODT/soln/tab Vraylar™ ziprasidone cap	Abilify® tab/IM inj ***Abilify Mycite® (with sensor) **aripiprazole ODT, soln asenapine (generic for Saphris®) Caplyta Capsule™ Clozaril® clozapine ODT Fanapt® tab & titration pk Fazaclor® **Geodon® tab, IM Invega® Latuda® Lybalvi® **Nuplazid™ tab, cap (QL) **olanzapine/fluoxetine paliperidone ER quetiapine fumarate ER (authorized generic only) Rexulti® tab Risperdal® ODT/soln/tab Saphris® Secuado® Patch Seroquel® IR Seroquel® XR Versacloz™ ziprasidone IM (gen Geodon®) Zyprexa® tab/IM/Zydis	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™ <ul style="list-style-type: none"> Indicated for the treatment of hallucinations and delusions associated with Parkinson's disease psychosis. <i>Quantity Limit Nuplazid™ = 2 per day</i> *** ♦ Clinical Criteria for Abilify Mycite® Initial Approval Criteria: For Three months SA Patient must: <ul style="list-style-type: none"> Be ≥ 18 years of age; AND Have tolerability to oral aripiprazole with suboptimal effects (as assessed by prescriber) that may be due to adherence problems; AND Have a smart phone compatible with the device; AND Give consent to a healthcare provider and caregiver (if applicable) to monitor the portal; AND There is a documented intervention by prescriber if nonadherence is detected Renewal Criteria: Every 3 Months Reevaluate <ul style="list-style-type: none"> Patient must: Continue to meet initial criteria; AND



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

1/1/24

	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
			<ul style="list-style-type: none"> • Have prescriber attestation that patient benefited from therapy; AND • Have prescriber attestation that there is a continued need for device (e.g., continued suboptimal effects and/or compliance); AND • Have a healthcare provider and caregiver (if applicable) agree to continue to monitor device; AND • Not have worsened target symptoms; AND • Not have had any treatment-limited adverse effects (e.g., hypersensitivity, suicidality, neuroleptic malignant syndrome, tardive dyskinesia, metabolic changes, pathological gambling and other compulsive behaviors, orthostatic hypotension, falls, seizures, cognitive and motor impairment, dysphagia, disruption in body temperature regulation, and leukopenia, neutropenia, and agranulocytosis); AND • Have a healthcare provider state reason why the patient cannot use long-acting injectable atypical antipsychotic if there is continued nonadherence.
	Atypical, Long-Acting Injectable		LENGTH OF AUTHORIZATIONS: 1 year
	Abilify Asimtufii[®] Abilify Maintena [®] Aristada [®] Aristada [®] Initio Invega Hafyera [™] , Sustenna [®] & Trinza [®] Perseris [™] Risperdal Consta [®] Uzedly[™]	Zyprexa [®] Relprevv [™]	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	Adasuve [®] Haldol decanoate (injection) pimozide Moban [®] molindone	Routine PDL edits



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

1/1/24

<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
perphenazine trifluoperazine thiothixene thioridazine		
*Movement Disorders CLOSED CLASS		
Austedo [®] tab Austedo XR [®] tab Austedo XR[®] titration pack Ingrezza [®] cap Ingrezza [®] Initiation Pack tetrabenazine tab (generic) Xenazine [®] Xenazine [®] tab		<p><u>LENGTH OF AUTHORIZATIONS:</u> 1 year</p> <p><u>*Clinical Criteria for Movement Disorders</u></p> <ul style="list-style-type: none"> • Diagnoses of Tardive Dyskinesia or Huntington's disease • Prescribed by or in consult with a neurologist or psychiatrist • Quantity limit <ul style="list-style-type: none"> ○ 4 tabs/day Austedo[®] ○ 42 tablets per 365 Austedo XR[®] titration pack ○ 1 cap/day Ingrezza[®] ○ 4 tabs/day Xenazine[®]



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
Neuropathic Pain		
<p>capsaicin OTC topical duloxetine 20, 30 & 60 mg (generic for Cymbalta®) gabapentin cap/tab/soln lidocaine 5% patch pregabalin cap</p>	<p><i>Cymbalta®</i> <i>Dermacinrx® PHN Pak™ Kit</i> <i>Drizalma™ Sprinkle</i> <i>duloxetine 40 mg (generic)</i> <i>Irenka™</i> <i>Gralise™ (gabapentin, ER)</i> <i>Horizant™ (gabapentin enacarbil ER)</i> <i>Irenka™</i> <i>Lidoderm® patch</i> <i>LidoPure 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>Zilacaine Patch</i> <i>Ztlido™ 1.8% (lidocaine topical system)</i></p>	<p>LENGTH OF AUTHORIZATIONS: 1 year</p> <p>Routine PDL</p> <p>Plus</p> <p>Criteria for Lyrica solution:</p> <ul style="list-style-type: none"> - If diagnosis is epilepsy/seizures, patient must have a problem swallowing tabs/capsules AND clinical reason why at least TWO preferred seizure medications cannot be used. - For other diagnoses, patient 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		
<p>pramipexole ropinirole HCl</p>	<p><i>Mirapex® IR & ER</i> <i>Neupro®</i> <i>pramipexole ER</i> <i>Requip® XR</i> <i>ropinirole HCl ER</i></p>	<p>LENGTH OF AUTHORIZATIONS: 1 year</p> <p>Routine PDL edits</p>
Sedatives / Hypnotics		
<p>temazepam 15 & 30 mg</p>	<p><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></p>	<p>LENGTH OF AUTHORIZATIONS: Length of the prescription (up to 3 months)</p> <p>Routine PDL edits</p>

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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
Sedatives / Hypnotics (Non-Benzodiazepine)		
eszopiclone zaleplon zolpidem	<i>Ambien® IR & CR</i> <i>Belsomra®</i> <i>Dayvigo™</i> <i>doxepin (generic for Silenor®)</i> <i>Elduar™</i> <i>*Hetlioz®</i> <i>**Hetlioz™ LQ</i> <i>Intermezzo®</i> <i>Lunesta®</i> <i>Quviviq™</i> <i>Rozerem®</i> <i>Silenor®</i> <i>Sonata®</i> <i>tasimelteon</i> <i>zolpidem CR</i> <i>zolpidem (generic Intermezzo®)</i> <i>zolpidem tartrate capsule</i>	<p>LENGTH OF AUTHORIZATIONS: 6 months. For Renewal - must document therapeutic benefit and confirm compliance</p> <p>Routine PDL edits plus</p> <p>* Clinical Criteria for Hetlioz® and tasimelteon</p> <ul style="list-style-type: none"> 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. <p>** Clinical Criteria for Hetlioz™ LQ oral suspension</p> <ul style="list-style-type: none"> For the treatment Nighttime sleep disturbances in SMS in pediatric patients AND Member must be 3 years to 15 years of age.
Skeletal Muscle Relaxants		
baclofen chlorzoxazone cyclobenzaprine HCL dantrolene sodium methocarbamol tizanidine tab	<i>Amrix®</i> <i>baclofen soln</i> <i>baclofen susp (gen Fleqsuvy™)</i> <i>*carisoprodol</i> <i>*carisoprodol/ASA</i> <i>*carisoprodol/ASA/codeine</i> <i>cyclobenzaprine ER</i> <i>Dantrium®</i> <i>Fexmid®</i> <i>Fleqsuvy™</i> <i>Lorzone®</i> <i>Lyvispah™</i> <i>metaxalone</i> <i>orphenadrine citrate</i> <i>orphenadrine/ASA/caffeine</i>	<p>LENGTH OF AUTHORIZATIONS:</p> <ul style="list-style-type: none"> 1 year for chronic conditions Duration of prescription (up to 3 months) for acute conditions <p>Routine PDL edits plus</p> <p>* Clinical Criteria for Carisoprodol Drugs</p> <ul style="list-style-type: none"> Patient 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



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	Parafon Forte® DSC Robaxin® Skelaxin® *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 Zyban®	LENGTH OF AUTHORIZATIONS: 6 months Routine PDL edits
*Stimulants/ADHD Medications (AGE) CLOSED CLASS		
Amphetamine Drugs		
Adderall®XR amphetamine salts combo (generic for Adderall IR) dextroamphetamine (generic for Dexedrine) Vyvanse® cap/chewable tab (lisdexamfetamine)	Adderall® IR (amphetamine salts combo) Adzenys XR ODT™ Adzenys ER™ susp Adzenys® ER amphetamine salts combo XR amphetamine sulfate (generic Evekeo®) amphetamine susp (generic Adzenys ER™ susp) Desoxyn® Dexedrine® dextroamphetamine SR & soln Dyanavel® XR susp Evekeo® Evekeo™ ODT lisdexamfetamine(generic Vyvanse®) methamphetamine	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits *♦ 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. Refer to Stimulant SA form (Stimulant/ADHD Medications SA Form) This does not include nonstimulant agents such as atomoxetine (generic for Strattera®), clonidine ER, guanfacine ER or others



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	<i>Mydayis ERTM</i> <i>Procentra[®] soln</i> <i>XelstrylTM</i> <i>ZenzediTM</i>	
Methylphenidate Drugs		
All methylphenidate IR generic methylphenidate soln Concerta[®] Daytrana[®] Transdermal dexmethylphenidate IR dexmethylphenidate XR	<i>AdhansiaTMXR</i> <i>AptensioTM XR</i> <i>AzstarysTM</i> <i>Cotempla XR-ODTTM</i> <i>Focalin[®] IR & Focalin[®] XR</i> <i>Jornay PM</i> <i>Metadate CD[®] & Metadate ER[®]</i> <i>Methylin ER[®], soln IR</i> <i>methylphenidate chew</i> <i>methylphenidate ER, 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, LA[®] & SR[®]</i> <i>Relexxii[®]</i> <i>QuilliChew ERTM</i> <i>QuillivantTM XR susp</i>	
Miscellaneous Drugs		
atomoxetine (generic for Strattera[®]) clonidine ER guanfacine ER	***armodafinil ***modafinil ***NuvigilTM ***Provigil[®] <i>Intuniv[®]</i> <i>Qelbree[®]</i> ***SunosiTM	***♦ <u>NuvigilTM/Provigil[®]/armodafinil/modafinil:</u> Refer to <u>Narcolepsy Medications SA Form</u>



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	Strattera® ***Wakix®	
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 clindamycin Phos 1% pledget, swab clindamycin/benzoyl peroxide (Duac®) erythromycin solution Panoxyl 4 Acne Cr Wash (OTC) Panoxyl 10 cleansing bar, foaming wash (OTC)	Acanya™ w/pump Acne Clearing System® (OTC) Aczone® Gel and Gel Pump Akliel® Amzeeq™ Arazlo™ Avar Cleanser, Medicated Pad Avar-E Avar-E LS Avar LS Cleanser, Medicated Pad Azelex® Benzaclin® & Benzaclin® Pump BP 10-1 Benzefoam™ regular & Ultra™ Benzepro benzoyl peroxide wash/cr/gel/ lotion/foam/towelette (RX) benzoyl peroxide 6%, 9% cleanser (OTC) BPO Kit Cleocin T® Clindacin™ Pac Kit Clindagel® clindamycin phosphate (generic for Clindagel®) clindamycin/benzoyl peroxide (generic for Acanya® Pump)	Routine PDL edits plus * ♦ Clinical Criteria for Dermatologic Acne Agents <ul style="list-style-type: none"> • 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.



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

1/1/24

	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		<p><i>clindamycin/benzoyl peroxide (generics for Benzaclin®)</i> <i>clindamycin phosphate foam, el, lotion, med swab</i> <i>clindamycin/tretinoin (generic Veltin®)</i> <i>Delos™ Lotion</i> <i>Duac® gel</i> <i>erythromycin gel/med. swab</i> <i>Evoclin™</i> <i>Inova™</i> <i>Lavoclen™ Cleanser & Kit</i> <i>Neuac™ topical/kit</i> <i>Onexton™ gel & w/Pump</i> <i>Ovace® Wash</i> <i>Ovace® Plus</i> <i>shampoo/cr/lotion/foam</i> <i>Pacnex® HP & LP</i> <i>Panoxyl® 3% cr (OTC)</i> <i>Promiseb® Complete</i> <i>Rosula Cleanser</i> <i>Se BPO® Wash Kit & cleanser</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></p>	



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	<i>Veltin®</i> <i>Winlevi®</i>	
Retinoids/Combinations, Topical		
adapalene gel OTC Retin® A 0.025%, 0.05, 0.1 % cr & 0.01, 0.025% gel	<i>Acnefree® Severe Kit (OTC)</i> <i>adapalene 0.1% cr/gel/lot</i> <i>adapalene 0.3% gel/gel w/pump</i> <i>adapalene-benzoyl peroxide (gen</i> <i>Epiduo® and Epiduo® Forte)</i> <i>Altreno™</i> <i>Atralin® 0.05% gel</i> <i>Avage® 0.1% cr</i> <i>Avita® 0.025% cr/gel</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>Renova® 0.02% cr/cr pump</i> <i>Retin®-A Micro 0.04%, 0.1% gel</i> <i>Retin®-A Micro 0.08%, 0.04%,</i> <i>0.1% pump</i> <i>Tazorac® cr/gel</i> <i>tazarotene 0.1% cr, foam</i> tretinoin 0.025, 0.1% cr & 0.01, 0.025, 0.05% gel <i>tretinoin microsphere 0.04% &</i> <i>0.1% gel</i> <i>Twynéo®</i> <i>Ziana® gel</i>	<p>* ♦ Age Edit for Fabior™ Foam</p> <ul style="list-style-type: none"> Members must be between the ages of 12 and 18 years of age
Antifungal Topical		
antifungal 1% cr, powder antifungal 2% cr ciclopirox soln ciclodan 8% soln	<i>Alevazol® OTC</i> <i>Azolen® Tincture OTC</i> <i>Bensal HP®</i> <i>Ciclodan® Kit</i>	<p>LENGTH OF AUTHORIZATIONS: 6 months</p> <p>Routine PDL edits plus</p>



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
<p>clotrimazole cr (OTC & RX) clotrimazole soln (OTC & RX) clotrimazole-betamethasone cr ketoconazole shampoo ketoconazole cr miconazole cr/spray (OTC) nystatin cr/oint/ powder terbinafine cr (OTC) tolnaftate cr/powder(OTC)</p>	<p><i>ciclopirox cr/shampoo/gel</i> <i>ciclopirox kit</i> <i>ciclopirox suspension</i> <i>clotrimazole-betamethasone lot</i> <i>*CNL 8™ Kit</i> <i>Desenex® Aero Powder (OTC)</i> <i>econazole</i> <i>Ertaczo®</i> <i>Exelderm® cr & soln</i> <i>Extina®</i> <i>Fungi-Nail® (OTC)</i> <i>Fungoid® Kit (OTC)</i> <i>Fungoid® (OTC)</i> <i>*Jublia®</i> <i>ketoconazole foam</i> <i>*Kerydin®</i> <i>Lamisil AT® cr/gel (OTC)</i> <i>Lamisil® Spray (OTC)</i> <i>Loprox® Kit/ Shampoo/susp</i> <i>Lotrimin AF® cr (OTC)</i> <i>Lotrimin Ultra® (OTC)</i> <i>Lotrisone® cr</i> <i>luliconazole (generic for Luzu)</i> <i>**Luzu®</i> <i>miconazole nitrate (OTC)</i> <i>miconazole Oint/ powder (OTC)</i> <i>Mentax®</i> <i>Naftin® cr/gel</i> <i>Naftifine CR</i> <i>Nyata Kit®</i> <i>Nizoral A-D® Shampoo (OTC)</i> <i>nystatin-triamcinolone cr/oint</i> <i>oxiconazole cr (generic Oxistat®)</i> <i>Oxistat® cr</i> <i>Oxistat® Lotion</i> <i>Pediaderm AF®</i></p>	<p>Select non-preferred topical Antifungals (CNL-8™, Jublia®, Kerydin™, Luzu®, Penlac®) require the submission of a Clinical SA. Refer to Antifungal Topical SA Form</p>



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	PediPak® *Penlac® tavorole (generic Kerydin®) Tinactin® Aero powder/ spray(OTC) tolnaftate aero powde/spray (OTC) tolnaftate soln (OTC) Vusion®	
Immunomodulators Atopic Dermatitis CLOSED CLASS		
Adbry™ **Dupixent® *Elidel® **Eucrisa™ *tacrolimus	*Opzelura™ pimecrolimus (generic Elidel) Protopic®	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus ♦ Clinical Criteria for Atopic Dermatitis-all drugs <ul style="list-style-type: none"> • Member must have an FDA approved diagnosis: Atopic dermatitis AND <ul style="list-style-type: none"> • Age and severity for drug as follows below: <ul style="list-style-type: none"> ○ Adbry™: moderate to severe for ages ≥ 18 years ○ Elidel®: tacrolimus mild to moderate for ages > 2 years ○ Eucisa™: mild to moderate for ages > 3months ○ Dupixent; moderate to severe for ages > 6 months ○ Opzelura™: mild to moderate for ages equal to or > 12 years ○ Protopic® 0.03%: moderate to severe for ages > 2 years. ○ Protopic® 0.1%: moderate to severe for ages > 16 years. AND the one that applies: <u>*Clinical Criteria for Elidel® or tacrolimus</u> <ul style="list-style-type: none"> • Prior documented trial & failure of 8 weeks of one (1) topical corticosteroid of medium to high potency (e.g., mometasone, fluocinolone) <u>**Clinical Criteria for Eucrisa™, Dupixent® and Adbry®</u> (reviewed electronically, AutoPA look back of 180 days for no more than a 30 day) <ul style="list-style-type: none"> • Prior documented trial & failure of 30-day trial (or contraindication) of <ul style="list-style-type: none"> ○ One (1) topical corticosteroid of medium to high potency (e.g., mometasone, fluocinolone) OR ○ One (1) topical calcineurin inhibitors (tacrolimus or pimecrolimus)



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

1/1/24

	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>														
			<p>*** Clinical Criteria for Opzelura™ (topical Janus kinase inhibitor)</p> <ul style="list-style-type: none"> • Prior documented trial & failure of 8 weeks of each <ul style="list-style-type: none"> ○ 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® AND ○ A trial and failure of Eucrisa™ • Other indications: Opzelura cream can not be approved for the indication of nonsegmental vitiligo in adult and pediatric patients ≥ 12 years old <p>QUANTITY LIMITS for Atopic Dermatitis</p> <table border="1" data-bbox="1003 743 1934 971"> <thead> <tr> <th>Drug</th> <th>Rolling qty limit</th> </tr> </thead> <tbody> <tr> <td>Adbry®</td> <td>4 syringes/28 days (initial dose) 2 syringes/28 days</td> </tr> <tr> <td>Dupixent</td> <td>2 300 mg inj), Maintance 300 mg (Q2W)</td> </tr> <tr> <td>Elidel® (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> </tbody> </table> <p>Dupixent other Diagnosis: Fax Form</p> <ul style="list-style-type: none"> ◆ Diagnosis of moderate to severe asthma Patient must have moderate to severe asthma diagnosed as ONE of the following types: <ul style="list-style-type: none"> ○ Asthma with eosinophilic phenotype with eosinophil count ≥ 150 cells/mcL; OR ○ Oral corticosteroid dependent asthma with at least 1 month of daily oral corticosteroid use within the last 3 months; AND ○ Patient must be 6 years of age or older ◆ Diagnosis of eosinophilic esophagitis (EoE); <ul style="list-style-type: none"> ○ Patient ≥12 years old; AND ○ Patient weighs ≥ 40 kg; AND ○ Prescribed by or consultation with an allergist or gastroenterologist; AND 	Drug	Rolling qty limit	Adbry®	4 syringes/28 days (initial dose) 2 syringes/28 days	Dupixent	2 300 mg inj), Maintance 300 mg (Q2W)	Elidel® (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/28 days (initial dose) 2 syringes/28 days																
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Virginia Medicaid's Preferred Drug List (PDL)/ Common Core Formulary

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
		<p>o Patient did not respond clinically to treatment with a topical glucocorticosteroid or proton pump inhibitor</p> <p>♦ Diagnosis of chronic rhinosinusitis with nasal polyposis (CRSwNP);</p> <ul style="list-style-type: none"> o Patient ≥ 18 years old o Patient has inadequate response after 3 consistent months use of preferred PDL intranasal steroids or oral corticosteroids; AND o Patient is concurrently treated with intranasal corticosteroids <p>♦ Diagnosis of Prurigo Nodularis for the treatment of adult patients with prurigo nodularis (PN)</p> <ul style="list-style-type: none"> o Age 18 or older o Diagnosis of (PN)
Psoriasis, Topical		
<p>calcipotriene cr/oint/soln</p>	<p><i>betamethasone/ calcipotriene</i> <i>Calcitrene[®]</i> <i>calcitriol</i> <i>calcipotriene/betamethasone</i> <i>(generic for Taclonex[®])</i> <i>Dovonex[®]</i> <i>Duobrii[™]</i> <i>*Enstilar[®] Foam (AGE)</i> <i>Micanol[®]</i> <i>Sorilux[™]</i> <i>Taclonex[®] & Taclonex[®] Scalp</i> <i>Vectical[™]</i> <i>Vtama[®]</i> <i>Zoryve[®]</i></p>	<p><u>LENGTH OF AUTHORIZATIONS:</u> 1 year</p> <p>Routine PDL edits plus</p> <p>*♦ <u>Clinical Criteria for Enstilar[®] Foam</u></p> <ul style="list-style-type: none"> • Length of Authorization: 4 weeks • Diagnosis of plaque psoriasis; AND • Minimum age of 18 years
Rosacea Agents, Topical		



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

1/1/24

Preferred Agents		Non-Preferred Agents		SA Criteria	
metronidazole cr/gel		azelaic acid (generic for Finacea®) brimonidine gel pump (generic Mirvaso®) Epsolay® Finacea® foam/gel ivermectin (generic Soolantra) Metrocream® Metrogel® metronidazole lot Mirvaso® Noritate® Rhofade® Rosadan™ Kit Soolantra® Zilxi™		LENGTH OF AUTHORIZATIONS: Routine PDL edits	1 year
Steroids					
Steroids, Topical Low Potency				LENGTH OF AUTHORIZATIONS: Routine PDL edits	1 year
hydrocortisone cr/gel/lot/oint hydrocortisone cr/oint OTC hydrocortisone/aloe cr OTC		anusol-hc 2.5% cr alclometasone cr/oint aqua glycolic HC Capex® shampoo Derma-smoothe-FS desonate gel/cr/lot/oint Desowen® lot fluocinolone 0.01% oil Micort™-HC Pediaderm® HC & Pediaderm® TA Scalpicin OTC Texacort®			
Steroids, Topical Medium Potency					
fluticasone propionate cr/oint		betamethasone valerate foam Beser Lotion and Kit			



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
mometasone furoate cr/oint/soln	<i>clocortolone cr</i> <i>Cloderm®</i> <i>Cordran® tape</i> <i>Cutivate® cr/lot</i> <i>Dermatop® cr/oint</i> <i>Elocon® cr/oint/soln</i> <i>fluocinolone acetoneide cr/oint/soln</i> <i>flurandrenolide cr/oint/tape</i> <i>fluticasone propionate lot</i> <i>hydrocortisone valerate cr/oint</i> <i>hydrocortisone butyrate (generic for Locoid Lotion)</i> <i>hydrocortisone butyrate cr/oint/soln/ emollient</i> <i>Locoid Lipocream</i> <i>Locoid Lotion</i> <i>Luxiq®</i> <i>Momexin®</i> <i>Pandel®</i> <i>prednicarbate cr/oint</i> <i>Synalar®</i> <i>Synalar TS®</i> <i>Ticanase kit®</i>	
Steroids, Topical High Potency		LENGTH OF AUTHORIZATIONS: 1 year
betamethasone valerate cr/lot/oint triamcinolone acetoneide cr/lot/oint	<i>amcinonide cr/lot/oint</i> <i>betamet diprop & prop gly cr/lot/oint</i> <i>betamet diprop cr/foam/gel/lot/oint</i> <i>DermacinRx® SilaPak™</i> <i>DermacinRx® Silazone</i> <i>DermacinRx® Therazole Pak</i> <i>desoximetasone cr/gel/oint/spray</i>	Routine PDL edits plus * ♦ Clinical Criteria for Sernivo™ <ul style="list-style-type: none"> Length of Authorization: 4 weeks (treatment beyond 4 weeks is not recommended). Member must have diagnosis of mild to moderate plaque psoriasis: AND At least 18 years of age



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	<i>desoximetasone (generic Topicort® spray) diflorasone diacetate cr/oint Diprolene® lot/oint DiproleneAF® cr Ellzia™ Pak Kit fluocinonide cr/ emollient/ gel/oint/soln halcinonide cr Halog® cr/oint/soln Kenalog® aerosol Loprox® Suspension Kit *Sernivo™ Silazone® II Kit Topicort® cr/gel/oint/spray Trianex® oint triamcinolone spray triamcinolone/dimethicone Vanos® cr Whytederm® Tdpak</i>	
Steroids, Topical Very High Potency		
clobetasol emollient clobetasol propionate cr/gel/oint/soln halobetasol propionate cr	<i>Apexicon™ E Bryhali™ (halobetasol propionate) clobetasol lot/shampoo clobetasol propionate foam/spray Clobex® lot/shampoo/spray Clobetex® Kit Clodan® kit Halonate® halobetasol propionate oint (generic for Lexette®) Impeklo™ Olux®</i>	



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

1/1/24

	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		<i>Olux[®]-E</i> <i>Temovate[®] oint</i> <i>Ultravate[®] cr/lotion/oint</i> <i>Ultravate[®] PAC & Ultravate[®] X</i>	
Endocrine and Metabolic Agents			
Androgenic Agents (Testosterone – Topical)			
	AndroGel[®] Pump Androderm[®] Patch testosterone pump (generic for AndroGel[®] Pump)	<i>AndroGel[®] packet, patch</i> <i>Axiron[®] soln</i> <i>Fortesta[®]</i> <i>Natesto Nasal Gel[®]</i> <i>Testim[®]</i> <i>testosterone (generic for Axiron[®])</i> <i>testosterone gel/packet/pump (generic for VogelxoTM)</i> <i>testosterone (generic for Fortesta[®])</i> <i>VogelxoTM gel/packet/pump</i> <i>XyostedTM</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits Plus
Antihyperuricemics			
	allopurinol 100mg and 300mg colchicine tabs Probenecid[®] probenecid & colchicine	<i>allopurinol 200MG</i> <i>colchicine caps</i> <i>Colcrys[®]</i> <i>febuxostat (generic Uloric[®])</i> <i>Gloperba[®]</i> <i>Mitigare[®]</i> <i>Uloric[®]</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits plus
Contraceptives*(long-acting IUDs & injectable)			



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

1/1/24

Preferred Agents		Non-Preferred Agents		SA Criteria
Depo-Provera® 104mg Kyleena™ Liletta® medroxyprogesterone 150mg Mirena® Nexplanon® Paragard® Skylla®		<i>Depo-Provera® 150mg</i>		LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
Diabetes Hypoglycemics: Injectable Amylin Analogs				
		<i>*SymLin®</i> <i>*SymLin® Pens</i>		LENGTH OF AUTHORIZATIONS: 1 year *Clinical Criteria for Injectable Amylin Analogs <ul style="list-style-type: none"> • Member must have a history of at least a 90day trial of insulin. • SymLin® is only indicated as adjunct therapy with insulin. • Member meeting ALL of the following criteria may be approved: <ul style="list-style-type: none"> ○ Diagnosis of Type 1 or 2 diabetes; AND ○ On insulin therapy, AND ○ Failure to achieve adequate glycemic control (HbA1c ≤ 6.5%)
♦Diabetes Hypoglycemics: Injectable and Oral Incretin Mimetics CLOSED CLASS				
Byetta® Trulicity™ Victoza®		<i>Adlyxin™</i> <i>Bydureon™ Bcise SQ</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 Non-Preferred Incretin Mimetics SA Form
Diabetes Hypoglycemics: Injectable Insulins				
Insulin Mix				LENGTH OF AUTHORIZATIONS: 1 year
Humalog® Mix 50/50 vial Humalog® Mix 75/25 vial Humulin® 70/30 pen/vial(OTC)		<i>Humalog® Mix 50/50 Kwikpen</i> <i>Humalog® Mix 75/25 Kwikpen</i>		Routine PDL edits



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

1/1/24

Preferred Agents		Non-Preferred Agents	SA Criteria
insulin aspart/insulin aspart protamine vial		insulin lispro protamine mix kwikpen Novolin [®] 70/30 vial (OTC) Novolog [®] Mix 70/30 pen/vial	
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			
insulin glargine pen and vial Lantus [®] Solostar [®] & vial Levemir [®] pen/vial		Basaglar [®] KwikPen [®] insulin degludec insulin glargine-yfgn Rezvoglar [®] Kwikpen Semglee [™] Toujeo [®] Solostar [®] 300 Units/mL Tresiba [®] FlexTouch [®] Pen 100 U/ml, 200 U/ml	
Rapid-Acting Insulins			
Humulin 500 U/M pen/vial Humalog [®] vial/pen Humalog [®] Cartridge Humalog Kwikpen 100 unit/ml Humalog Jr. Kwikpen Novolog [®] cart/vial/Flexpen 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 Lyumjev [™]	



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
Diabetes Oral Hypoglycemics		
Oral Hypoglycemics Alpha-Glucosidase Inhibitors		LENGTH OF AUTHORIZATIONS: 1 year
acarbose	Glyset® miglitol (generic Glyset®)	Routine PDL edits plus
♦ 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 metformin ER (generic for Glucophage® XR)	Fortamet® Glucophage® IR & XR Glutmetza® Riomet® susp metformin 625 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® Janumet XR® Januvia® Jentadueto™ Jentadueto XR™ Kombiglyze XR™ Onglyza™ Tradjenta™	alogliptin (generic Nesina™) alogliptin/metformin (generic Kazano™) alogliptin/pioglitazone (generic Oseni™) Glyxambi® Kazano™ Nesina™ Oseni™ Qtern® saxagliptin hcl(generic Onglyza™)	



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

1/1/24

Preferred Agents		Non-Preferred Agents	SA Criteria
		<i>saxagliptin/metformin ER</i> <i>(generic Kombiglyze XR™)</i> <i>Steglujan™</i> <i>Trijardy™ XR</i>	
Oral Hypoglycemics Meglitinides			
nateglinide repaglinide		<i>Prandin® & PrandiMet™</i> <i>repaglinide/metformin</i> <i>Starlix</i>	
Oral Hypoglycemics Second Generation Sulfonylureas			
glimepiride glipizide glipizide ER glyburide glyburide micronized		<i>Amaryl®</i> <i>Diabeta®</i> <i>Glucotrol®</i> <i>Glucotrol XL®</i>	
*Oral Hypoglycemics Sodium Glucose Co-Transporter 2 Inhibitor CLOSED CLASS			
Farxiga™ Invokana™ Invokamet™ Jardiance® Synjardy® Xigduo™ XR		<i>Brenzavvy®</i> <i>bexagliflozin</i> <i>Inpefa™</i> <i>Invokamet™ XR</i> <i>Segluromet™</i> <i>Steglatro™</i> <i>Synjardy® XR</i>	
Oral Hypoglycemics Thiazolidinediones			
pioglitazone		<i>Avandia®</i> <i>Actoplus Met® IR & XR</i> <i>Actos®</i> <i>Avandaryl®</i> <i>Avandamet®</i> <i>Duetact®</i> <i>pioglitazone/metformin</i> <i>pioglitazone/glimepiride</i>	



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
Erythropoiesis Stimulating Proteins		
Epogen® Retacrit™ (PFIZER)	<i>Aranesp® vial/syringe</i> <i>Procrit®</i> <i>Reblozyl®</i> <i>Retacrit™ (VIFOR)</i> <i>Mircera®</i>	<u>LENGTH OF AUTHORIZATIONS:</u> for duration of the prescription up to 6 months Routine PDL edits
Glucagon Agents (CLOSED CLASS)		
Baqsimi nasal Glucagon inj Glucagon emergency kit (Fresenius and Amphastar) inj Gvoke pen, syringe, vial SQ Proglycem suspension oral	<i>Diazoxide suspension oral</i> <i>Glucagon emergency kit (Lilly) inj</i> <i>Zegalogue Autoinjector & syringe SQ</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL
Glucocorticoids, Oral		
budesonide EC dexamethasone soln/tab hydrocortisone methylprednisolone dose pk methylprednisolone 4 mg tab prednisolone sod phosph soln prednisolone soln prednisone soln/tab/dose pk	<i>Alkindi® Sprinkle</i> <i>Cortef®</i> <i>cortisone acetate</i> <i>dexamethasone elixir/intensol</i> <i>Dexpak®</i> <i>*Emflaza™ (AGE)</i> <i>Entocort® EC</i> <i>Flo-Pred®</i> <i>Hemady®</i> <i>Medrol® dose pk/tab</i> <i>methylprednisolone 8,16 & 32mg tab</i> <i>Millipred DP® tab Does Pk</i> <i>Millipred® soln/tab</i> <i>Orapred® ODT</i> <i>Ortikos™</i> <i>prednisolone sod phosphate ODT/ soln</i> <i>prednisone intensol</i> <i>Rayos® DR tab</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits plus *♦Clinical Criteria for Emflaza™ <ul style="list-style-type: none"> • Trial and failure of all drugs does not apply to Emflaza™ • Diagnosis for treatment of Duchenne muscular dystrophy (DMD) • Minimum Age Limit = 2 years of age



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	<i>TaperDex</i> [®] <i>Tarpeyo</i> [™] <i>Veripred</i> [®]	
*Growth Hormone CLOSED CLASS		
Genotropin [®] Cartridge, Miniquick Norditropin FlexPro [®] Nutropin AQ [®] NuSpin [™]	<i>Humatrope</i> [®] cartridge/vial <i>Omnitrope</i> [®] cartridge/vial <i>Saizen</i> [®] cartridge/vial <i>Serostim</i> [®] vial <i>Skytrofa</i> [™] <i>Sogroya</i> [®] <i>Zomacton</i> [®] vial	<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>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 hereditary angioedema. Healthcare professionals should be aware of the similarity of symptoms between hypersensitivity reactions and hereditary angioedema and patients should be monitored closely.
*Pancreatic Enzymes		
Creon [®] Zenpep [®]	<i>Pancrease</i> [®] <i>Pertzye</i> [®] <i>Ultresa</i> [®] <i>Viokace</i> [®]	<u>LENGTH OF AUTHORIZATION:</u> 1 year Routine PDL edits plus <u>*Clinical Criteria for Pancreatic Enzymes</u> <ul style="list-style-type: none"> All drugs preferred and non preferred require a diagnosis of pancreatic insufficiency due to cystic fibrosis or chronic pancreatitis or pancreatectomy. <ul style="list-style-type: none"> If a member has a diagnosis of Cystic Fibrosis they do not have to try and fail of a preferred.



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
		<ul style="list-style-type: none"> 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	<i>Aygestin</i> [®] <i>Crinone (Vaginal)</i> <i>Depo-Provera 400 MG/ML</i> <i>Prometrium</i> [®] <i>Provera</i> [®]	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits
Progestins Used For Cachexia		
megestrol acetate	<i>Megace</i> [®] <i>Megace</i> [®] ES <i>megestrol suspension ES</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits
Vaginal Estrogens		
Premarin [®] Vaginal cr Vagifem [®] Vaginal tab	<i>Estrace</i> [®] Vaginal cr <i>estradiol cream (gen Estrace</i> [®]) <i>estradiol vaginal tab</i> <i>Estring</i> [®] Vaginal ring <i>Femring</i> [®] Vaginal ring <i>Imvexxy</i> [®] <i>Intrarosa</i> TM <i>Yuvafem</i> [®]	<u>LENGTH OF AUTHORIZATIONS:</u> 6 months Routine PDL edits
Weight Management Agents (CLOSED CLASS)		
Contrave orlistat Saxenda SQ Xenical Wegovy SQ	<i>Imcivree SQ</i> <i>Zepbound</i> TM	Length of authorization: As indicated on fax form Routine PDL edits plus Weight Management Agents SA Form
Gastrointestinal		
G I Antibiotics		



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
<p>Firvanq™ soln metronidazole tab vancomycin cap</p>	<p>*<i>Aemcolo™</i> <i>Alinia®</i> <i>Dificid®</i> <i>Flagyl® cap/ER</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>Xifaxan®</i> <i>vancomycin compounded oral soln kit</i> <i>vancomycin sol</i> <i>Vancocin®</i> ***<i>Vowst™</i></p>	<p>Length of authorization: 1 year</p> <p>Routine PDL edits</p> <p>*♦Clinical Criteria for Aemcolo</p> <ul style="list-style-type: none"> • 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) <p>**♦Clinical Criteria for Xifaxan</p> <p>Xifaxan®: 200 mg tabs: –</p> <ul style="list-style-type: none"> • Treatment of travelers' diarrhea caused by noninvasive strains of E. coli in patients • ≥ 12 years of age – • Quantity limit = 9 tabs per claim • Length of authorization 1 month <p>Xifaxan®: 550 mg tabs: –</p> <ul style="list-style-type: none"> • Reduction in risk of overt hepatic encephalopathy recurrence in patients <ul style="list-style-type: none"> ○ ≥ 18 years of age. ○ 2 tablets/day <p>OR –</p> <ul style="list-style-type: none"> • Treatment of irritable bowel syndrome with diarrhea (IBS-D) in patients <ul style="list-style-type: none"> ○ ≥ 18 years of age – ○ 3 times a day for 14 days. ○ 42 tablets /14 days, can be retreated up to two times with the same regimen. ○ Max 126 tablets per 365 days <p>Length of authorization = 3 months</p> <p>***♦ Clinical Criteria for Vowst™:</p> <ul style="list-style-type: none"> • Patient ≥ 18 years of age; AND • Patient has a confirmed diagnosis of recurrent <i>Clostridioides difficile</i> infection (CDI) with a total of ≥3 episodes of CDI within 12 months; AND



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
		<ul style="list-style-type: none"> • Antibiotic treatment for recurrent CDI must be completed 2 to 4 days prior to initiation of Vowst therapy; AND • Patient will take 10 oz of magnesium citrate (or 250 mL polyethylene glycol electrolyte solution for patients with impaired kidney function) the evening prior to initiation of Vowst therapy; AND • Patient 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 <p>Length of authorizations: 30 Days</p>
Antiemetic/Antivertigo Agents		
Cannabinoids (delta-9THC derivatives)		
*dronabinol cap	*Cesamet™ *Marinol® *Syndros®	<p>LENGTH OF AUTHORIZATIONS: 6 months</p> <p>*Dronabinol plus all non-preferred Antiemetic/Antivertigo agents require submission of a Clinical SA. Refer to Antiemetic/Antivertigo SA form</p>
5HT3 Receptor Blockers		
ondansetron ODT/soln/tab	Aloxi® Anzemet® Akynzeo® granisetron tab, vial Kytril® tab ondansetron vial, amp, syringe palonosetron vial, syringe Sancuso® patch Sustol® Zofran® ODT/soln/tab Zuplenz® film	<p>LENGTH OF AUTHORIZATIONS: 3 months, unless otherwise noted</p> <p>Routine PDL</p>
NK-1 Receptor Antagonist		
	Aponvie™ aprepitant capsule/pack Cinvanti™ (Intraven)	<p>LENGTH OF AUTHORIZATIONS: Length of chemotherapy regimen or a maximum of 6 months</p>



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	Emend® Bi Pak/ cap Emend® Tri-fold pack/susp Emend® vial fosaprepitant vial Varubi™ IV, Tab	
Other		LENGTH OF AUTHORIZATIONS: 1 year, unless otherwise noted
*Diclegis® meclizine (OTC, Rx) metoclopramide soln, syr, tab, vial **Phenadoz® supp prochlorperazine tab **promethazine soln, syr, tab (AGE)	Antivert® Barhemsys® *Bonjesta™ Compazine® supp/tab Compro® dimenhydrinate tab, vial doxylamine succinate/ vit B6 Metozolv® ODT metoclopramide ODT **Phenergan® (AGE) prochlorperazine supp, vial **promethazine 50mg supp, vial, ampul (AGE) Reglan® scopolamine (gen Transderm-Scop®) Tigan® Transderm-Scop® trimethobenzamide	**♦Promethazine products approved for members ≥ 2 years *♦Bonjesta®/Diclegis®: Approve through EDD. patient must be pregnant and at least 18 years of age.
*GI Motility, Chronic		
Amitiza® **Linzess™ lubiprostone Movantik®	alosetron Lotronex® lubiprostone (authorized generic) Motegrity™ Relistor® Symproic® Trulance™	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



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	<i>Viberzi™</i>	**Pediatric use of Linzess 72 mcg only: 1. Diagnosis of functional constipation (FC) 2. Member is 6 to 17 years of age 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>	LENGTH OF AUTHORIZATIONS: 14 days Routine PDL edits
Histamine-2 Receptor Antagonists (H-2 RA)		
famotidine (OTC & RX) *famotidine oral susp	<i>cimetidine tab/syrup (OTC/RX)</i> <i>nizatidine cap/susp</i> <i>Pepcid® susp/tab (OTC/RX)</i>	LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits plus *♦AutoPA for famotidine oral susp <ul style="list-style-type: none"> • Is under 12 years of age • Continuation of therapy, has a claim in past 90 days • Is using a Tube feeding
Proton Pump Inhibitors		
omeprazole RX pantoprazole Protonix® susp	<i>Aciphex® DR tab/sprinkle</i> <i>dexlansoprazole dr (gen Dexilant®)</i> <i>Dexilant®</i> <i>esomeprazole magnesium cap/susp</i>	LENGTH OF AUTHORIZATIONS: 12 weeks; unless member meets an exception; then 1 year Routine PDL edits plus



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

1/1/24

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



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
Genitourinary		
Alpha-Blockers and Androgen Hormone Inhibitors For Benign Prostatic Hypertrophy (BPH)		
Alpha-Blockers for BPH		LENGTH OF AUTHORIZATIONS: 1 year
alfuzosin tamsulosin HCL	<i>Flomax[®]</i> <i>Rapaflo[®]</i> <i>silodosin (generic Rapaflo)</i> <i>Uroxatral[®]</i>	Routine PDL edits plus
Androgen Hormone Inhibitors for BPH		
dutasteride finasteride	<i>Avodart[®]</i> <i>dutasteride/tamsulosin</i> <i>Jalyn[®]</i> <i>Proscar[®]</i>	
Androgen Hormone Inhibitors and Phosphodiesterase (PDE) 5 Inhibitor for BPH		
	<i>Entadfi[™]</i>	
Phosphodiesterase (PDE) 5 Inhibitor for BPH		*Step edit for Cialis[®] & tadalafil 2.5 and 5mg - trial and failure of Alpha Blockers and Androgen Inhibitors for BPH. Prescriber must attest that the member is not on the state's sex offenders list. Consult or evaluation by Urologist.
	<i>*Cialis[®] (ST)</i> <i>tadalafil 5 mg (ST)</i>	
Urinary Antispasmodics (Bladder Relaxant)		
oxybutynin tab/syrup oxybutynin ER solifenacin Toviaz[™]	<i>darifenacin ER (generic Enablex[®])</i> <i>Detrol[®] & Detrol[®] LA</i> <i>Ditropan[®] & *Ditropan[®] XL</i> <i>fesoterodine fumarate er</i> <i>flavoxate</i> <i>Gelnique[™] gel/gel Pump</i> <i>Gemtesa[®]</i> <i>Myrbetriq[™] tab, Granules</i> <i>Oxytrol[®] transdermal includes OTC</i> <i>Sanctura XR</i> <i>trospium IR & ER</i>	



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

1/1/24

	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		<p><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>Hadlima™</i> <i>Hulio®</i> <i>Hyrimoz®</i> <i>Idacio®</i> <i>Ilaris®</i> <i>Ilumya™</i> <i>Inflectra®</i> <i>Kevzara® inj, pen</i> <i>Kineret®</i> <i>Ohumiant®</i> <i>Orencia®</i> <i>Otezla®</i> <i>Otrexup®</i> <i>Rasuvo™</i> <i>RediTrex®</i> <i>Remicade®</i> <i>Renflexis®</i> <i>Rinvoq™</i> <i>Skyrizi™ syringe, Pen</i> <i>Siliq®</i> <i>Simponi®</i> <i>Simponi® Aria</i> <i>Spevigo®</i> <i>Stelara® vial/syringe</i> <i>Sotyktu™</i> <i>Taltz®</i> <i>Tremfya™</i> <i>Trexall®</i></p>	<p>For a list of Cytokine and CAM Antagonists and criteria for approval see Appendix A</p>



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

1/1/24

	<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
		Uplizna [®] Xatmep [™] Xeljanz [™] , Xeljanz [™] XR and solution Yuflyma [®] Yusimry [™]	
Ophthalmic			
Antibiotics			
	bacitracin/polymyxin b sulfate oint ciprofloxacin drops erythromycin gentamicin drops/oint moxifloxacin drops (generic Vigamox) ofloxacin drops polymyxin/trimethoprim tobramycin	AzaSite [™] drops bacitracin Besivance [®] drops Ciloxan [®] drops/oint Garamycin [®] drops/oint gatifloxacin 0.5% soln Ilotycin [®] levofloxacin drops moxifloxacin drops (generic Moxeza [®]) Natacyn [®] neomycin/polymix/gramicidin neomycin/bacitracin/polymyxin oint Neosporin [®] Ocuflax [®] Polytrim [®] sulfacetamide oint/ soln Tobrex [®] drops/oint Vigamox [®] Zymaxid [®]	<u>LENGTH OF AUTHORIZATIONS:</u> Date of service only; no refills Routine PDL edits
Antibiotic/Steroid Combinations			
	neomycin/polymyxin/dexamethasone oint/susp sulfacetamide/prednisolone	Blephamide [®] S.O.P. Maxitrol [®] oint/susp neomycin/bacitracin/poly/HC	<u>LENGTH OF AUTHORIZATION:</u> Date of service only; no refills Routine PDL edits



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

1/1/24

Preferred Agents		Non-Preferred Agents	SA Criteria
Tobradex[®] oint/susp tobramycin/dexamethasone susp		<i>neomycin/polymyxin/HC</i> <i>Pred-G[®] oint</i> <i>Tobradex[®] ST</i> <i>Zylet[®]</i>	
Antihistamines/Mast Cell Stabilizers			
Antihistamines			LENGTH OF AUTHORIZATIONS: 1 year
Alaway OTC[®] ketotifen fumerate olopatadine (generic Patanol & Pataday) Zaditor[®] OTC		<i>bepotastine (gen Bepreve[®])</i> <i>Bepreve[®]</i> <i>Elestat[®]</i> <i>epinastine 0.05% eye drops</i> <i>Lastacast[®]</i> <i>Optivar[®]</i> <i>Patanol[®] Rx and OTC</i> <i>Pataday[®] Rx and OTC</i> <i>Zerviate[™]</i>	Routine PDL edits
Mast Cell Stabilizers			
cromolyn sodium		<i>Alocril[®]</i> <i>Alomide[®]</i>	
Anti-inflammatory Agents			
Corticosteroids			
Durezol[®] fluorometholone prednisolone acetate		<i>Alrex[™]</i> <i>Dexamethasone</i> <i>difluprednate (gen Durezol[®])</i> <i>Flarex[®]</i> <i>FML[®], FML Forte[®]</i> <i>loteprednol etabonate (generic Lotemax[™] gel)</i> <i>Lotemax[™] drops/gel/oint</i> <i>Maxidex[®]</i> <i>Omnipred[®]</i> <i>Pred Forte[®] & Pred Mild[®]</i>	



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	<p>prednisolone sod phosphate Vexol[®]</p>	
Immunomodulators	CLOSED CLASS	
<p>Restasis[®] Restasis Multidose[®] Xiidra[®]</p>	<p><i>Cequa[™]</i> <i>cyclosporine</i> <i>*Eysuvis</i> <i>Miebo[™]</i> <i>Tyrvaya[™] Nasal Spray</i> <i>**Verkazia[®]</i></p>	<p>*♦DUR DURATION LIMIT Eysuvis 1 bottle in 3 months, message returned to pharmacist stating "Limitation Exceeded: Quantity Limit of 1 Bottle per 3 Months. Patient needs Intraocular Pressure (IOP) Test. If IOP test completed, Physician to call 1-800-932-6648 for a Clinical SA"</p> <p>**♦Verkazia criteria (PDL criteria do not apply)</p> <ul style="list-style-type: none"> • Patient 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: <ul style="list-style-type: none"> 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
<p>diclofenac sodium flurbiprofen sodium ketorolac 0.4%& 0.5%</p>	<p><i>Acular[®] 0.5% & LS[®] 0.4%</i> <i>Acuvail[®]</i> <i>bromfenac 0.09%</i> <i>BromSite[™]</i> <i>*Ilevro[™]0.3% (QL)</i></p>	<p>Routine PDL edits</p> <p>*Ilevro[™] is limited to 1 bottle plus 1 refill</p>



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	<i>Inveltys™ (loteprednol etabonate)</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® levobunolol 0.5% metipranolol 0.3% timolol maleate	<i>Betagan® 0.5%</i> <i>betaxolol 0.5%</i> <i>Betoptic-S® 0.25%</i> <i>brimonidine-timolol (gen Combigan®)</i> <i>Istalol® 0.5%</i> <i>timolol (generic for Timoptic OcuDose)</i> <i>Timoptic® drops 0.25% & 0.5%</i> <i>Timoptic® XE 0.25% & 0.5%</i> <i>soln-gel</i>	
Carbonic Anhydrase Inhibitors		
Azopt® dorzolamide dorzolamide/timolol	<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>	
Cholinergic Muscarinic Receptor Agonist		
	*Vuity™	*Indicated for the treatment of presbyopia in adults



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

1/1/24

Preferred Agents		Non-Preferred Agents	SA Criteria
Rho Kinase Inhibitor			
Rhopressa®			
Rocklatan			
Prostaglandin Analogs			
latanoprost	<i>bimatoprost</i>		
Travatan Z®	<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>Xelpros® (latanoprost)</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>		
levocetirizine tab/OTC	<i>cetirizine D tab (OTC)</i>		
	<i>Clarinex®</i>		



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

1/1/24

Preferred Agents		Non-Preferred Agents	SA Criteria
loratadine tab/syrup OTC		<i>Clarinex-D[®]</i> <i>desloratadine ODT</i> <i>fexofenadine</i> <i>fexofenadine/PSE ER</i> <i>fexofenadine suspension</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
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>	Routine PDL edits plus
Short Acting Metered Dose Inhalers or Devices			
Proair[®] HFA ProAir[®] RespiClick Ventolin[®] HFA		<i>albuterol HFA (Proair[®])</i> <i>albuterol HFA (Ventolin[®])</i> <i>albuterol HFA (Proventil[®])</i> <i>levalbuterol tartrate HFA</i> <i>ProAir[®] Digihaler[™]</i> <i>Proventil[®] HFA</i> <i>Xopenex[®] HFA</i>	
Short Acting Nebulizers			
albuterol sulfate (premixed)		<i>levalbuterol soln</i> <i>Xopenex[®]</i>	



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

1/1/24

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® Lonhala™ Magnair™ Tudorza™ *Yupelri™ (revefenacin)	<u>LENGTH OF AUTHORIZATION:</u> 1 year Routine PDL edits plus *Clinical Criteria Yupelri™ (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® (AGE) Stiolto Respimat™ (AGE)		<u>LENGTH OF AUTHORIZATION:</u> 1 year Routine PDL edits



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

1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
Phosphodiesterase 4 (PDE4) Inhibitors CLOSED CLASS		
<p>*roflumilast</p>	<p><i>*Daliresp[®]</i></p>	<p>LENGTH OF AUTHORIZATION: 1 year</p> <p>*Clinical Criteria for Daliresp[®] and roflumilast</p> <ul style="list-style-type: none"> • 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
<p>Advair[®] Diskus Advair[®] HFA Dulera[®] Symbicort[®] Trelegy[®] Ellipta</p>	<p><i>AirDuo[®] Digihaler[®]</i> <i>Airduo[™] Respiclick[®]</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></p>	<p>Routine PDL edits</p>
Inhaled Corticosteroids: Metered Dose Inhalers CLOSED CLASS		
<p>Arnuity[™] Ellipta[®] Asmanex Twister[®] fluticasone HFA (gen Flovent[®] HFA)</p>	<p><i>Alvesco[®]</i> <i>Aerospan[™]</i> <i>ArmonAir[®] Digihaler[™]</i> <i>Asmanex HFA[®]</i></p>	



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

1/1/24

Preferred Agents		Non-Preferred Agents		SA Criteria
fluticasone Diskus (gen Flovent® Diskus) Flovent® Diskus Flovent® HFA Pulmicort Flexhaler®		<i>QVAR® & QVAR® Redihaler</i>		LENGTH OF AUTHORIZATIONS: 1 year Routine PDL edits
Inhaled Corticosteroids: Nebulizer Solution CLOSED CLASS				
budesonide respules		<i>Pulmicort® Respules</i>		
Nasal Steroids				
Dymista® fluticasone Rx		<i>azelastine/fluticasone nasal spray (generic for Dymista®)</i> <i>Beconase AQ®</i> <i>budesonide (generic for Rhinocort® Aqua)</i> <i>budesonide (generic Rhinocort® Allergy OTC)</i> <i>Children's Qnasl™</i> <i>Clarispray OTC</i> <i>Flonase®</i> <i>Flonase Sensimist (OTC)</i> <i>flunisolide</i> <i>fluticasone OTC</i> <i>ipratropium bromide spray</i> <i>mometasone (generic Nasonex®)</i> <i>Nasonex OTC</i> <i>Omnaris®</i> <i>Qnasl™</i> <i>Rhinocort Aqua®</i> <i>Rhinocort® Allergy OTC</i> <i>Ryaltris®</i> <i>Ticanase®</i> <i>triamcinolone OTC</i> <i>triamcinolone acetonide</i> <i>Veramyst®</i>		



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1/1/24

Preferred Agents	Non-Preferred Agents	SA Criteria
	<i>XhanceTM</i> <i>ZetonnaTM</i>	
*♦Cough and Cold Drug		
Ala-Hist DM benzonatate cap codeine/ promethazine guaifenesin/codeine phosphate hydrocodone/homatrop syrup Iophen-C NR phenylephrine HCl/promethazine HCl promethazine DM syrup Tusnel[®] Pediatric Drops	<i>lohist-DM syrup</i> <i>All other Legend cough and cold drugs are non-preferred</i> <i>Tessalon[®] perle</i>	<u>LENGTH OF AUTHORIZATION:</u> Date of Service Only Routine PDL edits * Children under the age of 6 years are not eligible for cough and cold drugs. FDA advisory: cough and cold medicines containing codeine or hydrocodone are limited to adults 18 years and older because the risks of these medicines outweigh their benefits in children younger than 18
Epinephrine, Self-Injected		
epinephrine 0.15 mg & 0.3 mg (authorized generic EpiPen[®] & EpiPen[®] Jr) Epipen[®] Epipen[®] Jr	<i>Auvi-Q[®]</i> <i>epinephrine 0.15 mg & 0.3mg (generic EpiPen[®] and EpiPen[®] Jr)</i> <i>epinephrine 0.15mg & 0.3mg (generic Adrenaclick)</i> <i>SymjepiTM (epinephrine)</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits
Intranasal Antihistamines		
azelastine 0.1%	<i>Astepro[®] 0.15%</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits



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1/1/24

<i>Preferred Agents</i>	<i>Non-Preferred Agents</i>	<i>SA Criteria</i>
Leukotriene Receptor Antagonists		
montelukast tabs/chewable tabs	<i>Accolate®</i> <i>Singulair® tabs/chew tabs/granules</i> <i>montelukast granules</i> <i>zafirlukast</i> <i>Zyflo™</i> <i>Zyflo CR™</i> <i>zileuton ER</i>	<u>LENGTH OF AUTHORIZATIONS:</u> 1 year Routine PDL edits