

This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

01/01/12 Version 2012.4

- Prior authorization for a non-preferred agent in any category will be given only if there has been a trial of the preferred brand/generic equivalent or preferred formulation of the active ingredient, at a therapeutic dose, that resulted in a partial response with a documented intolerance.
- Prior authorization of a non-preferred isomer, pro-drug, or metabolite will be considered with a trial of a preferred
 parent drug of the same chemical entity, at a therapeutic dose, that resulted in a partial response with documented
 intolerance or a previous trial and therapy failure, at a therapeutic dose, with a preferred drug of a different chemical
 entity indicated to treat the submitted diagnosis. (The required trial may be overridden when documented evidence is
 provided that the use of these preferred agent(s) would be medically contraindicated.)
- Unless otherwise specified, the listing of a particular brand or generic name includes all legend forms of that drug. OTC drugs are not covered unless specified.
- PA criteria for non-preferred agents apply in addition to general Drug Utilization Review policy that is in effect for the entire pharmacy program, including, but not limited to, appropriate dosing, duplication of therapy, etc.
- The use of pharmaceutical samples will not be considered when evaluating the members' medical condition or prior prescription history for drugs that require prior authorization.

Acronyms

- CL Requires clinical PA. For detailed clinical criteria, please refer to: http://www.dhhr.wv.gov/bms/Pharmacy/Pages/PriorAuthorizationCriteria.aspx
- NR New drug has not been reviewed by P & T Committee
- AP Non-preferred and selected preferred drugs, where indicated, are subject to auto-PA criteria. See PA criteria column.



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA		
ACNE AGENTS (To	ACNE AGENTS (Topical) ^{AP}				
		FECTIVE			
	AKNE-MYCIN (erythromycin) AZELEX (azelaic acid) clindamycin erythromycin sodium sulfacetamide	ACZONE (dapsone) CLEOCIN-T (clindamycin) EVOCLIN (clindamycin) KLARON (sodium sulfacetamide)	Thirty (30) day trials each of one preferred retinoid and two unique chemical entities in two other subclasses, including the generic version of a requested non-preferred product, are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present. (In cases of pregnancy, a trial of retinoids will not be required.)		
	RETI	NOIDS			
	RETIN A liquid & Micro (tretinoin) TAZORAC (tazarotene) tretinoin cream, gel	adapalene AVITA (tretinoin) DIFFERIN (adapalene) RETIN-A cream, gel (tretinoin) TRETIN-X (tretinoin)	PA required after 17 years of age for tretinoin products.		
	KERATOLYTICS (Benzoyl Peroxides)			
	benzoyl peroxide ETHEXDERM (benzoyl peroxide) OSCION (benzoyl peroxide)	BENZAC WASH (benzoyl peroxide) BENZEFOAM (benzoyl peroxide) BENZEFOAM ULTRA (benzoyl peroxide) BREVOXYL (benzoyl peroxide) DESQUAM (benzoyl peroxide) LAVOCLEN (benzoyl peroxide) TRIAZ (benzoyl peroxide)	Acne kits are non-preferred.		
		ION AGENTS			
	benzoyl peroxide/urea erythromycin/benzoyl peroxide sulfacetamide sodium/sulfur wash/cleanser	ACANYA (clindamycin phosphate/benzoyl peroxide) AVAR (sulfur/sulfacetamide) BENZACLIN GEL (benzoyl peroxide/ clindamycin) BENZAMYCIN PAK (benzoyl peroxide/ erythromycin) benzoyl peroxide/clindamycin gel	Thirty (30) day trials each of one preferred retinoid and two unique chemical entities in two other subclasses, including the generic version of a requested non-preferred product, are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is		



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
		CLENIA (sulfacetamide sodium/sulfur) DUAC CS (benzoyl peroxide/ clindamycin) EPIDUO (adapalene/benzoyl peroxide) INOVA 4/1 (benzoyl peroxide/salicylic acid) NUOX (benzoyl peroxide/sulfur) PLEXION (sulfacetamide sodium/sulfur) PRASCION (sulfacetamide sodium/sulfur) ROSADERM (sulfacetamide sodium/sulfur) ROSANIL (sulfacetamide sodium/sulfur) ROSULA (sulfacetamide sodium/sulfur/ urea) sulfacetamide sodium/sulfur/ urea SULFOXYL (benzoyl peroxide/sulfur) SULFATOL (sulfacetamide sodium/sulfur/urea) VELTIN (clindamycin/tretinoin) ZENCIA WASH (sulfacetamide sodium/sulfur) ZIANA (clindamycin/tretinoin)	present. (In cases of pregnancy, a trial of retinoids will not be required.) In addition, thirty day trials of combinations of the corresponding preferred single agents available are required before non-preferred combination agents will be authorized.
ALZHEIMER'S AGE	ENTSAP		
	CHOLINESTER A	ASE INHIBITORS	
	donepezil	ARICEPT (donepezil) ARICEPT 23mg (donepezil) ARICEPT ODT(donepezil) COGNEX (tacrine) donepezil ODT EXELON CAPSULE (rivastigmine) EXELON PATCH (rivastigmine) galantamine galantamine ER RAZADYNE (galantamine) rivastigmine	A thirty (30) day trial of a preferred agent is required before a non-preferred agent in this class will be authorized unless one of the exceptions on the PA form is present. Aricept 23mg tablets will be approved when there is a diagnosis of moderate-to-severe Alzheimer's Disease, a trial of donepezil 10mg daily for at least three (3) months, and donepezil 20mg daily for an additional one (1) month. Aricept and donepezil ODT will be approved only when the oral dosage form is not appropriate for the patient.



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
			Members currently utilizing Exelon patches as of 1/1/2012 may continue.
	NMDA RECEPTO	OR ANTAGONIST	
	NAMENDA (memantine)		
ANALOECICO NAI	DOOTIC CHORT ACTING (Now you	va na ta na INAP	
ANALGESICS, NAI	RCOTIC - SHORT ACTING (Non-pai	· · · · · · · · · · · · · · · · · · ·	0: (0) 1 (1) (1)
	APAP/codeine ASA/codeine codeine dihydrocodeine/ APAP/caffeine hydrocodone/APAP hydrocodone/ibuprofen hydromorphone levorphanol morphine oxycodone oxycodone/APAP oxycodone/APAP pentazocine/APAP pentazocine/naloxone ROXICET (oxycodone/acetaminophen) tramadol tramadol/APAP	ABSTRAL (fentanyl) ACTIQ (fentanyl) butalbital/APAP/caffeine/codeine butalbital/ASA/caffeine/codeine butorphanol COMBUNOX (oxycodone/ibuprofen) DEMEROL (meperidine) DILAUDID (hydromorphone) fentanyl FENTORA (fentanyl) FIORICET W/ CODEINE (butalbital/APAP/caffeine/codeine) FIORINAL W/ CODEINE (butalbital/ASA/caffeine/codeine) LAZANDA (fentanyl) LORCET (hydrocodone/APAP) LORTAB (hydrocodone/APAP) MAGNACET (oxycodone/APAP) meperidine NUCYNTA (tapentadol) OPANA (oxymorphone) ONSOLIS (fentanyl) oxycodone/ibuprofen OXYFAST (oxycodone) OXYIR (oxycodone) PANLOR (dihydrocodeine/ APAP/caffeine) PERCOCET (oxycodone/APAP) PERCODAN (oxycodone/APAP) PERCODAN (oxycodone/APAP) PERCODAN (oxycodone/APAP) TECODAN (pentazocine/APAP) TALACEN (pentazocine/APAP) TALWIN NX (pentazocine/naloxone) TREZIX (dihydrocodeine/ APAP/caffeine) PREZIX (dihydrocodeine/ APAP/caffeine)	Six (6) day trials of at least four (4) chemically distinct preferred agents (based on narcotic ingredient only), including the generic formulation of a requested non-preferred product, are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present. Fentanyl lozenges and Onsolis will only be approved for a diagnosis of cancer and as an adjunct to a long-acting agent. Neither will be approved for monotherapy. Limits: Unless the patient has escalating cancer pain or another diagnosis supporting increased quantities of short-acting opioids, all short acting solid forms of the narcotic analgesics are limited to 120 tablets per 30 days for the purpose of maximizing the use of longer acting medications to prevent unnecessary breakthrough pain in chronic pain therapy.



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
		ULTRACET (tramadol/APAP) ULTRAM (tramadol) VICODIN (hydrocodone/APAP) VICOPROFEN (hydrocodone/ibuprofen) VOPAC (codeine/acetaminophen) XODOL (hydrocodone/acetaminophen) ZAMICET (hydrocodone/APAP) ZYDONE (hydrocodone/acetaminophen) XOLOX (oxycodone/APAP)	
ANALGESICS, NA	RCOTIC - LONG ACTING (Non-pare		
	fentanyl transdermal KADIAN (morphine) 10mg, 20mg, 30mg, 50mg, 60mg, 100mg methadone morphine ER tablets OPANA ER (oxymorphone)	AVINZA (morphine) BUTRANS (buprenorphine) CONZIP ER (tramadol) DOLOPHINE (methadone) DURAGESIC (fentanyl) EXALGO ER (hydromorphone) EMBEDA (morphine/naltrexone) KADIAN (morphine) 80mg, 200mg morphine ER capsules MS CONTIN (morphine) NUCYNTA ER (tapentadol) ORAMORPH SR (morphine) oxycodone ER OXYCONTIN (oxycodone) oxymorphone ER RYZOLT ER (tramadol) tramadol ER ULTRAM ER (tramadol)	Six (6) day trials each of two preferred unique long acting chemical entities are required before a non-preferred agent will be approved unless one of the exceptions on the PDL form is present. The generic form of the requested non-preferred agent, if available, must be tried before the non-preferred agent will be approved. Butrans will be approved if the following criteria are met: 1. Diagnosis of moderate to severe chronic pain requiring continuous around-the-clock analgesia or 2. Patient cannot take oral medications and has a diagnosis of chronic pain and 3. Needs analgesic medication for an extended period of time and 4. Has had a previous trial** of a non-opioid analgesic medication and 5. Previous trial of one opioid medication** and 6. Current total daily opioid dose is ≤ 80 mg morphine equivalents daily or dose of transdermal fentanyl is ≤ 12.5 mcg/hr 7. Patient is not currently being



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
			treated with buprenorphine
			**Requirement is waived for patients who cannot swallow
			Dose optimization is required for achieving equivalent doses of Kadian 80mg and 200mg. AP does not apply.
			Exception: Oxycodone ER will be authorized if a diagnosis of cancer is submitted without a trial of the preferred agents.
ANALGESICS (Topi	ical) ^{AP}		
	capsaicin lidocaine lidocaine/prilocaine xylocaine	EMLA (lidocaine/prilocaine) FLECTOR PATCH (diclofenac) LIDODERM PATCH (lidocaine) LIDAMANTLE (lidocaine) LIDAMANTLE HC (lidocaine/hydrocortisone) LMX 4 (lidocaine) PENNSAID (diclofenac) SYNERA (lidocaine/tetracaine) VOLTAREN GEL (diclofenac) ZOSTRIX (capsaicin)	Ten (10) day trials of each of the preferred topical anesthetics (lidocaine, lidocaine/prilocaine, and xylocaine) are required before a non-preferred topical anesthetic will be approved unless one of the exceptions on the PA form is present. Lidoderm patches will be approved for a diagnosis of post-herpetic neuralgia. Thirty (30) day trials of each of the preferred oral NSAIDS and capsaicin are required before Voltaren Gel will be approved unless one of the exceptions on the PA form is present. Flector patches will be approved only for a diagnosis of acute strain, sprain or injury after a five (5) day trial of one of the preferred oral NSAIDs and for a maximum duration of 14 days unless one of the exceptions on the PA form is



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
			present.
ANDROGENIC AGE	NTS		
	ANDRODERM (testosterone) ANDROGEL (testosterone)	AXIRON (testosterone) FORTESTA (testosterone) TESTIM (testosterone)	The non-preferred agents will be approved only if one of the exceptions on the PA form is present.
ANGIOTENSIN MOI	DULATORS ^{AP}		
	ACE INH	IIBITORS	
	benazepril captopril enalapril fosinopril lisinopril quinapril ramipril	ACCUPRIL (quinapril) ACEON (perindopril) ALTACE (ramipril) CAPOTEN (captopril) LOTENSIN (benazepril) MAVIK (trandolapril) moexipril MONOPRIL (fosinopril) perindopril PRINIVIL (lisinopril) trandolapril UNIVASC (moexipril) VASOTEC (enalapril) ZESTRIL (lisinopril)	Fourteen (14) day trials of each of the preferred agents in the corresponding group, with the exception of the Direct Renin Inhibitors, are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present.
	ACE INHIBITOR CO	MBINATION DRUGS	
	benazepril/amlodipine benazepril/HCTZ captopril/HCTZ enalapril/HCTZ fosinopril/HCTZ lisinopril/HCTZ quinapril/HCTZ	ACCURETIC (quinapril/HCTZ) CAPOZIDE (captopril/HCTZ) LEXXEL (enalapril/felodipine) LOTENSIN HCT (benazepril/HCTZ) LOTREL (benazepril/amlodipine) moexipril/HCTZ PRINZIDE (lisinopril/HCTZ) TARKA (trandolapril/verapamil) trandolapril/verapamil UNIRETIC (moexipril/HCTZ) VASERETIC (enalapril/HCTZ) ZESTORETIC (lisinopril/HCTZ)	
	ANGIOTENSIN II RECEP	TOR BLOCKERS (ARBs)	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	AVAPRO (irbesartan) BENICAR (olmesartan) DIOVAN (valsartan) losartan MICARDIS (telmisartan)	ATACAND (candesartan) COZAAR (losartan) EDARBI (azilsartan) eprosartan TEVETEN (eprosartan)	
	ARB COME	BINATIONS	
	AVALIDE (irbesartan/HCTZ) BENICAR-HCT (olmesartan/HCTZ) DIOVAN-HCT (valsartan/HCTZ) EXFORGE (valsartan/amlodipine) EXFORGE HCT (valsartan/amlodipine/HCTZ) losartan/HCTZ MICARDIS-HCT (telmisartan/HCTZ)	ATACAND-HCT (candesartan/HCTZ) AZOR (olmesartan/amlodipine) HYZAAR (losartan/HCTZ) TEVETEN-HCT (eprosartan/HCTZ) TRIBENZOR (olmesartan/amlodipine/HCTZ) TWYNSTA (telmisartan/amlodipine)	
	DIRECT RENI	N INHIBITORS	
	AMTURNIDE (aliskiren/amlodipine/HCTZ) ^{AP} TEKAMLO (aliskiren/amlodipine) ^{AP} TEKTURNA (aliskiren) ^{AP} TEKTURNA HCT (aliskiren/HCTZ) ^{AP} VALTURNA (aliskiren/valsartan) ^{AP}		A thirty (30) day trial of one preferred ACE, ARB, or combination agents, at the maximum tolerable dose, is required before Tekturna will be approved. Tekturna HCT, Valturna, Tekamlo or Amturnide will be approved if the criteria for Tekturna are met and the patient also needs the other agents in the combination.
ANTICOAGULANT		TABLE	
	ARIXTRA (fondaparinux) FRAGMIN (dalteparin) LOVENOX (enoxaparin)	enoxaparin fondaparinux INNOHEP (tinzaparin)	Trials of each of the preferred agents will be required before a non-preferred agent will be approved unless one of the exceptions on the PA form is present.
	OR	AL	
	PRADAXA (dabigatran) ^{AP} warfarin	XARELTO (rivaroxaban) ^{NR}	Pradaxa will be approved for the diagnosis of non-valvular atrial fibrillation.



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
DRUG CLASS	FREFERRED AGENTS	NON-FREFERRED AGENTS	FA CRITERIA
ANTICONVULSAN			
		VANTS	
	carbamazepine CARBATROL (carbamazepine) DEPAKOTE SPRINKLE (divalproex) divalproex EC divalproex DR EPITOL (carbamazepine) FELBATOL (felbamate) gabapentin GABITRIL (tiagabine) levetiracetam lamotrigine lamotrigine chewable LYRICA (pregabalin) oxcarbazepine tablets topiramate TRILEPTAL SUSPENSION (oxcarbazepine) valproic acid zonisamide	BANZEL(rufinamide) carbamazepine XR DEPAKENE (valproic acid) DEPAKOTE (divalproex) DEPAKOTE ER (divalproex) EQUETRO (carbamazepine) FANATREX SUSPENSION (gabapentin) FANATREX (gabapentin) HORIZANT (gabapentin) KEPPRA (levetiracetam) KEPPRA (levetiracetam) LAMICTAL (lamotrigine) LAMICTAL CHEWABLE (lamotrigine) LAMICTAL XR (lamotrigine) levetiracetam ER NEURONTIN (gabapentin) SABRIL (vigabatrin) STAVZOR (valproic acid) TEGRETOL (carbamazepine) TOPAMAX (topiramate) TRILEPTAL TABLETS (oxcarbazepine) VIMPAT (lacosamide) ZONEGRAN (zonisamide)	A fourteen (14) day trial of one of the preferred agents in the corresponding group is required for treatment naïve patients with a diagnosis of a seizure disorder before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present. A thirty (30) day trial of one of the preferred agents in the corresponding group is required for patients with a diagnosis other than seizure disorders unless one of the exceptions on the PA form is present. Non-preferred anticonvulsants will be approved for patients on established therapies with a diagnosis of seizure disorders with no trials of preferred agents required. In situations where ABrated generic equivalent products are available, "Brand Medically Necessary" must be hand-written by the prescriber on the prescription in order for the brand name product to be reimbursed. Requests for Gralise will be authorized if the following criteria are met: 1. Diagnosis of post herpetic neuralgia 2. Trial of a tricyclic antidepressant



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
			for a least thirty days 3. Trial of gabapentin immediate release formulation (positive response without adequate duration) 4. Request is for once daily dosing with 1800 mg. maximum daily dosage
	BARBITU	IRATES ^{AP}	
	mephobarbital phenobarbital primidone	MEBARAL (mephobarbital) MYSOLINE (primidone)	
	BENZODIA	ZEPINESAP	
	clonazepam DIASTAT (diazepam rectal) diazepam tablets	diazepam rectal gel KLONOPIN (clonazepam)	
	HYDAN	TOINSAP	
	DILANTIN INFATABS (phenytoin) PEGANONE (ethotoin) phenytoin	CEREBYX (fosphenytoin) DILANTIN (phenytoin) PHENYTEK (phenytoin)	
	SUCCIN	NIMIDES	
	CELONTIN (methsuximide) ethosuximide ZARONTIN (ethosuximide)		
ANTIDEPRESSANT	rs, other		
	SNF	RIS ^{AP}	
	CYMBALTA (duloxetine) venlafaxine ER capsules	EFFEXOR (venlafaxine) EFFEXOR XR (venlafaxine) PRISTIQ (desvenlafaxine) venlafaxine VENLAFAXINE ER Tablets (venlafaxine)	A six (6) week trial each of a preferred agent and an SSRI is required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present.
		N NON-SSRI, OTHER ^{AP}	
	bupropion SR bupropion XL mirtazapine SAVELLA (milnacipran) ^{AP*}	APLENZIN (bupropion hbr) bupropion IR DESYREL (trazodone) EMSAM (selegiline)	* Savella will be approved for a diagnosis of fibromyalgia or a previous thirty (30) day trial of a drug that infers fibromyalgia:



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	trazodone	nefazodone OLEPTRO ER (trazodone) REMERON (mirtazapine) WELLBUTRIN (bupropion) WELLBUTRIN SR (bupropion) WELLBUTRIN XL (bupropion) VIIBRYD (vilazodone hcl)	gabapentin, Cymbalta, Lyrica, amitriptyline or nortriptyline.
	SELECT	ED TCAs	
	imipramine hcl	imipramine pamoate TOFRANIL (imipramine hcl) TOFRANIL PM (imipramine pamoate)	A twelve (12) week trial of imipramine hcl is required before a non-preferred TCA will be authorized.
ANTIDEPRESSANT	TS, SSRIs ^{AP}		
	citalopram fluoxetine fluvoxamine LEXAPRO (escitalopram) paroxetine sertraline	CELEXA (citalopram) LUVOX (fluvoxamine) LUVOX CR (fluvoxamine) PAXIL (paroxetine) PAXIL CR (paroxetine) paroxetine ER PEXEVA (paroxetine) PROZAC (fluoxetine) RAPIFLUX (fluoxetine) SARAFEM (fluoxetine) ZOLOFT (sertraline)	Thirty (30) day trials each of two (2) of the preferred agents are required before a non-preferred agent will be approved unless one of the exceptions on the PA form is present. Upon hospital discharge, patients admitted with a primary mental health diagnosis and have been stabilized on a non-preferred SSRI will receive an authorization to continue that drug.
ANTIEMETICS ^{AP}			
	5HT3 RECEPTO	OR BLOCKERS	
	ondansetron ondansetron ODT	ANZEMET (dolasetron) KYTRIL (granisetron) granisetron GRANISOL (granisetron) SANCUSO (granisetron) ZOFRAN (ondansetron) ZOFRAN ODT (ondansetron) ZUPLENZ (ondansetron)	A 3-day trial of a preferred agent is required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present. PA is required for ondansetron when limits are exceeded.
CANNABINOIDS			
		CESAMET (nabilone) dronabinol MARINOL (dronabinol)	Cesamet will be authorized only for the treatment of nausea and vomiting associated with cancer chemotherapy for patients who



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
		ANTAGONIGE	have failed to respond adequately to 3-day trials of conventional treatments such as promethazine or ondansetron and are over 18 years of age. Marinol will be authorized only for the treatment of anorexia associated with weight loss in patients with AIDS or cancer and unresponsive to megestrol; or for the prophylaxis of chemotherapy induced nausea and vomiting unresponsive to 3-day trials of ondansetron or promethazine for patients between the ages of 18 and 65.
		ANTAGONISTS	
	EMEND (aprepitant)		
ANTIFUNGALS (Or	al)		
	clotrimazole fluconazole* ketoconazole CL nystatin terbinafine CL	ANCOBON (flucytosine) DIFLUCAN (fluconazole) flucytosine GRIFULVIN V TABLET (griseofulvin) griseofulvin GRIS-PEG (griseofulvin) itraconazole LAMISIL (terbinafine) MYCELEX (clotrimazole) MYCOSTATIN Tablets (nystatin) NIZORAL (ketoconazole) NOXAFIL (posaconazole) ORAVIG BUCCAL (miconazole) SPORANOX (itraconazole) VFEND (voriconazole) voriconazole	Non-preferred agents will be approved only if one of the exceptions on the PA form is present. *PA is required when limits are exceeded. PA is not required for griseofulvin suspension for children up to 6 years of age for the treatment of tinea capitis.
ANTIFUNGALS (To		INGALS	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THEDADELITIC			
THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	econazole ketoconazole MENTAX (butenafine) NAFTIN (naftifine) nystatin	ciclopirox ERTACZO (sertaconazole) EXELDERM (sulconazole) LOPROX (ciclopirox) MYCOSTATIN (nystatin) NIZORAL (ketoconazole) OXISTAT (oxiconazole) PENLAC (ciclopirox) SPECTAZOLE (econazole) VUSION (miconazole/petrolatum/zinc oxide) XOLEGEL (ketoconazole)	Fourteen (14) day trials of two (2) of the preferred agents are required before one of the non-preferred agents will be authorized unless one of the exceptions on the PA form is present. If a non-preferred shampoo is requested, a fourteen (14) day trial of one preferred product (ketoconazole shampoo) is required. Oxistat cream will be approved for children 12 and under for tinea corporis, tinea cruris, tinea pedis, and tinea (pityriasis) versicolor.
	ANTIFUNGAL/STER	OID COMBINATIONS	,
	clotrimazole/betamethasone nystatin/triamcinolone	KETOCON PLUS (ketoconazole/hydrocortisone) LOTRISONE (clotrimazole/betamethasone) MYCOLOG (nystatin/triamcinolone) AP	
ANTIHISTAMINES,	MINIMALLY SEDATING ^{AP}		
	ANTIHIS*	TAMINES	
	ALAVERT (Ioratadine) cetirizine loratadine TAVIST-ND (Ioratadine)	ALLEGRA (fexofenadine) CLARINEX Tablets (desloratadine) CLARINEX REDITABS (desloratadine) CLARINEX Syrup (desloratadine) CLARITIN (loratadine) fexofenadine (Rx and OTC) levocetirizine XYZAL (levocetirizine) ZYRTEC (Rx and OTC) (cetirizine) ZYRTEC SYRUP (cetirizine)	Thirty (30) day trials of at least two (2) chemically distinct preferred agents (in the age appropriate form), including the generic formulation of a requested non-preferred product, are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present.
	ANTIHISTAMINE/DECONG	SESTANT COMBINATIONS	
	ALAVERT-D (loratadine/pseudoephedrine) cetirizine/pseudoephedrine loratadine/pseudoephedrine SEMPREX-D (acrivastine/ pseudoephedrine)	ALLEGRA-D (fexofenadine/ pseudoephedrine) CLARINEX-D (desloratadine/ pseudoephedrine) CLARITIN-D (loratadine/pseudoephedrine) fexofenadine/ pseudoephedrine (Rx and OTC)	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA	
DRUG CLASS				
		ZYRTEC-D (cetirizine/pseudoephedrine)		
ANTIMIGRAINE AG	ENTS, TRIPTANSAP			
	TRIP	TANS		
	IMITREX NASAL SPRAY(sumatriptan) IMITREX INJECTION (sumatriptan) naratriptan sumatriptan	AMERGE (naratriptan) AXERT (almotriptan) FROVA (frovatriptan) IMITREX tablets (sumatriptan) MAXALT (rizatriptan) MAXALT MLT (rizatriptan) RELPAX (eletriptan) sumatriptan nasal spray/injection ZOMIG (zolmitriptan)	Three (3) day trials of each unique chemical entity of the preferred agents are required before a non-preferred agent will be approved unless one of the exceptions on the PA form is present. Quantity limits apply for this drug class. *AP does not apply to nasal spray or injectable sumatriptan.	
	TRIPTAN CO	DMBINATIONS	or injectable sumatriplan.	
		TREXIMET (sumatriptan/naproxen sodium)		
ANTIPARKINSON'S	S AGENTS (Oral)			
	ANTICHO	LINERGICS		
	benztropine trihexyphenidyl	COGENTIN (benztropine)	Patients starting therapy on drugs in this class must show a documented allergy to all of the preferred agents, in the corresponding class, before a non-preferred agent will be authorized.	
	COMT IN	HIBITORS	processes agent in accommon	
		COMTAN (entacapone) TASMAR (tolcapone)		
	DOPAMINE AGONISTS			
	pramipexole ropinirole	MIRAPEX (pramipexole) MIRAPEX ER (pramipexole) REQUIP (ropinirole) REQUIP XL (ropinirole)	Mirapex, Mirapex ER, Requip, and Requip XL will be approved for a diagnosis of Parkinsonism with no trials of preferred agents required.	
		KINSON'S AGENTS		
	amantadine AP bromocriptine	AZILECT (rasagiline) ELDEPRYL (selegiline)	Amantadine will be approved only for a diagnosis of Parkinsonism.	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	carbidopa/levodopa selegiline STALEVO (levodopa/carbidopa/entacapone)	levodopa/carbidopa ODT LODOSYN (carbidopa) PARCOPA (levodopa/carbidopa) SINEMET (levodopa/carbidopa) ZELAPAR (selegiline)	
ANTIPSYCHOTICS	, ATYPICAL		
	SINGLE IN	GREDIENT	
	clozapine GEODON (ziprasidone) INVEGA (paliperidone) INVEGA SUSTENNA (paliperidone)* risperidone risperidone ODT risperidone solution SEROQUEL (quetiapine) AP (25mg Tablet Only)	ABILIFY (aripiprazole) CLOZARIL (clozapine) FANAPT (iloperidone) FAZACLO (clozapine) LATUDA (lurasidone) olanzapine RISPERDAL (risperidone) RISPERDAL CONSTA (risperidone)* RISPERDAL ODT (risperidone) RISPERDAL SOLUTION (risperidone) SAPHRIS (asenapine) SEROQUEL XR (quetiapine) ZYPREXA (olanzapine) ZYPREXA INTRAMUSCULAR (olanzapine)*	A fourteen (14) day trial of a preferred agent is required for treatment naïve patients before a non-preferred agent will be approved unless one of the exceptions on the PA form is present. Upon discharge, a hospitalized patient stabilized on a non-preferred agent may receive authorization to continue this drug for labeled indications and at recommended dosages. Claims for Seroquel 25 mg will be approved: 1. for a diagnosis of schizophrenia or 2. for a diagnosis of bipolar disorder or 3. when prescribed concurrently with other strengths of Seroquel in order to achieve therapeutic treatment levels. Seroquel 25 mg. will not be approved for use as a sedative hypnotic. All antipsychotic agents require prior authorization for children up to six (6) years of age.



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THED A DELITIO			
THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
DNOG GEAGG			between the ages of 6-17 for irritability associated with autism. Abilify will be prior authorized for MDD if the following criteria are met: 1. The patient is at least 18 years of age. 2. Diagnosis of Major Depressive Disorder (MDD), 3. Evidence of trials of appropriate therapeutic duration (30 days), at the maximum tolerable dose, of at least one agent in two of the following classes: SSRI, SNRI or bupropion in conjunction with Seroquel at doses of 150 mg or more 4. Prescribed in conjunction with an SSRI, SNRI, or bupropion 5. The daily dose does not exceed 15 mg. *All injectable antipsychotic products require clinical prior authorization.
	ATYPICAL ANTIPSYCHO	TIC/SSRI COMBINATIONS	
		SYMBYAX (olanzapine/fluoxetine)	
ANTIVIRALS (Oral)			
	ANTI H	IERPES	
	acyclovir VALTREX (valacyclovir)	famciclovir FAMVIR (famciclovir) valacyclovir ZOVIRAX (acyclovir)	Five (5) day trials each of the preferred agents are required before the non-preferred agents will be authorized unless one of the exceptions on the PA form is present.
		FLUENZA	
	RELENZA (zanamivir) TAMIFLU (oseltamivir)	FLUMADINE (rimantadine) rimantadine amantadine AP	The anti influenza agents will be approved only for a diagnosis of influenza.



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS ANTIVIRALS (Topical)** ABREVA (docosand) DENAVIR (penciclovir) ABREVA (penciclovir) ARTOPIC DERMATITIS ELIDEL (pimecrolimus)** ELIDEL (pimecrolimus)** PROTOPIC (tacrolimus) PROTOPIC (tacrolimus) Athiry (30) day trial of a preferred medium or high potency topical conticosteroid is required before protopic will be considered; additionally, a thiry (30) day trial of Elide libe to considered; additionally, a thiry (30) day trial of Elide libe to considered; additionally, a thiry (30) day trial of Elide libe considered; unless one of the exceptions on the PA form is present. BETA BLOCKERS (Oral) & MISCELLANEOUS ANTIANGINALS (Oral)** BETA BLOCKERS (Sotalo) BETAPACE (sotalo) BETAPACE (sotalo) BETAPACE (sotalo) BETAPACE (sotalo) BETAPACE (sotalo) CORGARD (nadolo) Delaxiolo Delax				
ARREVA (docosanol) DENAVIR (penciclovir) BETA BLOCKERS acebutolol atenolol betaxolol	DRUG CLASS		NON-PREFERRED AGENTS	PA CRITERIA
ARREVA (docosanol) DENAVIR (penciclovir) BETA BLOCKERS acebutolol atenolol betaxolol	ANTIVIRALS (Topic	cal) ^{ap}		
ELIDEL (pimecrolimus) PROTOPIC (tacrolimus) A thirty (30) day trial of a preferred medium or high potency topical conticosteroid is required before coverage of Elidel will be considered; additionally, a thirty (30) day trial of Elidel is required before Protopic will be considered, unless one of the exceptions on the PA form is present. BETA BLOCKERS BETAPACE (sotalol) CARTROL (caretolol) CARTROL (caretolol) CORGARD (nadolol) metoprolol metoprolol ER nadolol pindolol propranolol propranolol ER sotalol TENARMIN (atenolol) TOPROL XL (metoprolol) ZEBETA (bisoprolol) ZEBETA (bisoprolol) ZEBETA (bisoprolol/HCTZ) nadolol/bendroflumethiazide propranolol/HCTZ			ZOVIRAX (acyclovir)	preferred agents are required before the non-preferred agent will
medium or high potency topical conficosteroid is required before coverage of Elidel will be considered, additionally, a thirty (30) day trial of Elidel is required before Protopic will be considered, additionally, a thirty (30) day trial of Elidel is required before Protopic will be considered, unless one of the exceptions on the PA form is present. BETA BLOCKERS acebutolol acebutolol BETAPACE (sotalol) BILOCADREN (timolol) betaxolol betaxolol betaxolol CARTROL (carteolol) metoprolol ER nadolol metoprolol ER nadolol propranolol propranolol propranolol propranolol propranolol propranolol propranolol propranolol ER sotalol SECTRAL (acebutolol) TENORRIN (atenolol) TOPROL XL (metoprolol) ZEBETA (bisoprolol) ZEBETA (bisoprolol) TENORRIN (atenolol) TOPROL XL (metoprolol) ZEBETA (bisoprolol) TENORRIN (atenolol) TENORRIN (atenolol) TOPROL XL (metoprolol) TENORRIN (atenolol) TENORRIN (atenolol) TOPROL XL (metoprolol) TENORRIN (atenolol) TENORRIC (acenolol/chlorta) TENORRIC (alenolol/chlorta) TENORRIC (alenolol/chlorta) TENORRIC (alenolol/chlorthalidone) ZIAC (bisoprolol/HCTZ) TENORRIC (alenolol/chlorthalidone)	ATOPIC DERMATIT	ΓIS		
acebutolol atenolol betaxolol betaxolol bisoprolol metoprolol ER nadolol propranolol propranolol ER sotalol propranolol BETA BLOCKER/DIURETIC COMBINATION DRUGS BETA BLOCKER/DIURETIC (atenolol/Chlorthalidone) bisoprolol/HCTZ nadolol/bendroflumethiazide propranolol/HCTZ atenolol/Chlorthalidone bisoprolol/HCTZ nadolol/HCTZ acebutolol BETA BLOCKER/DIURETIC (sotalol) BETA BLOCKER/DIURETIC (sotalol) BETA BLOCKER/DIURETIC (atenolol/Chlorthalidone) propranolol/HCTZ nadolol/bendroflumethiazide propranolol/HCTZ nadolol/bendroflumethiazide propranolol/HCTZ acebutolol BETA BLOCKER/DIURETIC (atenolol/Chlorthalidone) propranolol/HCTZ nadolol/bendroflumethiazide propranolol/HCTZ NIDERIDE (propranolol/HCTZ) TENORETIC (atenolol/Chlorthalidone) ZIAC (bisoprolol/HCTZ) ZIAC (bisoprolol/HCTZ) ZIAC (bisoprolol/HCTZ) ZIAC (bisoprolol/HCTZ)		ELIDEL (pimecrolimus) ^{AP}	PROTOPIC (tacrolimus)	medium or high potency topical corticosteroid is required before coverage of Elidel will be considered; additionally, a thirty (30) day trial of Elidel is required before Protopic will be considered, unless one of the exceptions on the
acebutolol atenolol BLOCADREN (timolol) BLOCADREN (timolol) betaxolol bisoprolol CARTROL (carteolol) CORGARD (nadolol) generic formulation of a requested non-preferred agents, including the generic formulation of a requested non-preferred product, are required before one of the non-preferred agents, including the generic formulation of a requested non-preferred product, are required before one of the non-preferred agents will be approved unless one pindolol (INNOPRAN XL (propranolol) agents will be approved unless one pindolol (Propranolol) (BETA BLOCKERS	(Oral) & MISCELLANEOUS ANTIAN	NGINALS (Oral) ^{AP}	
atenolol betaxolol BLOCADREN (timolol) three (3) chemically distinct betaxolol betaxolol BYSTOLIC (nebivolol) preferred agents, including the generic formulation of a requested non-preferred agents, including the generic formulation of a requested non-preferred product, are required metoprolol ER INDERAL LA (propranolol) propranolol ER INNOPRAN XL (propranolol) agents will be approved unless one pindolol (EVATOL (penbutolol) propranolol (EVATOL (penbutolol)) propranolol (EVATOL (penbutolol)) propranolol ER (EVATOL (penbutolol)) propranolol ER (EVATOL (penbutolol)) (EVATOL (penbutolol)) propranolol ER (EVATOL (penbutolol)) (EVATOL		BETA BL	OCKERS	
atenolol/chlorthalidone bisoprolol/HCTZ metoprolol/HCTZ nadolol/bendroflumethiazide propranolol/HCTZ TENORETIC (atenolol/HCTZ) ZIAC (bisoprolol/HCTZ)		atenolol betaxolol bisoprolol metoprolol metoprolol ER nadolol pindolol propranolol propranolol ER sotalol timolol	BLOCADREN (timolol) BYSTOLIC (nebivolol) CARTROL (carteolol) CORGARD (nadolol) INDERAL LA (propranolol) INNOPRAN XL (propranolol) KERLONE (betaxolol) LEVATOL (penbutolol) LOPRESSOR (metoprolol) SECTRAL (acebutolol) TENORMIN (atenolol) TOPROL XL (metoprolol) ZEBETA (bisoprolol)	three (3) chemically distinct preferred agents, including the generic formulation of a requested non-preferred product, are required before one of the non-preferred agents will be approved unless one of the exceptions on the PA form is
bisoprolol/HCTZ INDERIDE (propranolol/HCTZ) metoprolol/HCTZ LOPRESSOR HCT (metoprolol/HCTZ) nadolol/bendroflumethiazide TENORETIC (atenolol/chlorthalidone) propranolol/HCTZ ZIAC (bisoprolol/HCTZ)				
BETA- AND ALPHA-BLOCKERS		bisoprolol/HCTZ metoprolol/HCTZ nadolol/bendroflumethiazide	INDERIDE (propranolol/HCTZ) LOPRESSOR HCT (metoprolol/HCTZ) TENORETIC (atenolol/chlorthalidone)	
		BETA- AND ALF	PHA-BLOCKERS	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS carvedilol labetalol COREG (carvedilol) COREG (Creavedilol) TRANDATE (labetalol) ANTIANGINALS RANEXA (ranolazing) ^{AP} Ranexa will be approved for patients with angina who are also staking a calcium channel blocker, a beta blocker, or a nitrite as single agents or a combination agent containing one of these ingredients. BLADDER RELAXANT PREPARATIONS ^{AP} Oxybutynin FR TOVIAZ (fesoterodine) VESICARE (solifenacin) DETROL (tolterodine) DETROL (a) (tolterodine) DETRO						
Iabetalol COREG CR (carvedilol) TRANDATE (labetalol)		PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA		
Ranexa will be approved for patients with anging who are also taking a calcium channel blocker, a beta blocker, or a nitrite as single agents or a combination agent containing one of these ingredients. BLADDER RELAXANT PREPARATIONS ^{AP} Oxybutynin Carbon Sybutynin Carbon Containing One of these ingredients. ENABLEX (darifenacin) DETROL (tolterodine) DETROL (tolterodine) DETROL (tolterodine) DITROPAN L (oxybutynin) DITROPAN L (oxybutynin) OXYTROL (oxybutynin) OXYTROL (oxybutynin) OXYTROL (oxybutynin) SANCTURA (trospium) Trospium BISPHOSPHONATES alendronate FOSAMAX SOLUTION (alendronate) ACTONEL (risedronate) ACTONEL (risedronate) BONIVA (blandronate) DIDRONEL (elidronate) BONIVA (blandronate) DIDRONEL (elidronate) FOSAMAX TABLETS (alendronate) FOSAMAX PLUS D (alendronate)viramin D) ZOMETA (zoledronic acid) OTHER BONE RESORPTION SUPPRESSION AND RELATED AGENTS MIACALCIN (calcitonin) MIACALCIN (calcitonin) Evista (calcitonin) Evista will be approved for posterior acid invasive breast cancer. Evista will be approved for invasive breast cancer.			COREG CR (carvedilol)			
BLADDER RELAXANT PREPARATIONS ^{AP} Oxybutynin Containing one of these ingredients. BIABLEX (darifenacin) DETROL (tolterodine) DETROL LA (tolterodine) DITROPAN La (toxybutynin) DITROPAN LA (toxybutynin) CATURA KR (trospium) SANCTURA (trospium) SA		ANTIAN	IGINALS			
oxybutynin eR			RANEXA (ranolazine) ^{AP}	patients with angina who are also taking a calcium channel blocker, a beta blocker, or a nitrite as single agents or a combination agent		
DETROL (folterodine) TOVIAZ (fesoterodine) VESICARE (solifenacin) DETROL A (totterodine) DITROPAN XL (oxybutynin) DITROPAN XL (oxybutynin) DITROPAN XL (oxybutynin) DITROPAN XL (oxybutynin) SELNIQUE (oxybutynin) SANCTURA XR (trospium) SANCTURA XR (trospium) SANCTU	BLADDER RELAXA	ANT PREPARATIONS AP				
BISPHOSPHONATES alendronate FOSAMAX SOLUTION (alendronate) ACTONEL (risedronate) ACTONEL with CALCIUM (risedronate/calcium) ATELVIA (risedronate) BONIVA (ibandronate) DIDRONEL (etidronate) FOSAMAX TABLETS (alendronate) FOSAMAX PLUS D (alendronate/vitamin D) ZOMETA (zoledronic acid) OTHER BONE RESORPTION SUPPRESSION AND RELATED AGENTS acicitonin Calcitonin EVISTA (raloxifene) FORTICAL (calcitonin) BPH AGENTS A 30-day trial of the preferred agent is required before a non-preferred agent will be approved. B calcitonin Evista will be approved for postmenopausal women with osteoporosis or at high risk for invasive breast cancer.		oxybutynin ER TOVIAZ (fesoterodine)	DETROL (tolterodine) DETROL LA (tolterodine) DITROPAN (oxybutynin) DITROPAN XL (oxybutynin) GELNIQUE (oxybutynin) OXYTROL (oxybutynin) SANCTURA (trospium) SANCTURA XR (trospium)	chemically distinct preferred agents is required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is		
BISPHOSPHONATES alendronate FOSAMAX SOLUTION (alendronate) ACTONEL (risedronate) ACTONEL with CALCIUM (risedronate/calcium) ATELVIA (risedronate) BONIVA (ibandronate) DIDRONEL (etidronate) FOSAMAX TABLETS (alendronate) FOSAMAX PLUS D (alendronate/vitamin D) ZOMETA (zoledronic acid) OTHER BONE RESORPTION SUPPRESSION AND RELATED AGENTS MIACALCIN (calcitonin) WIACALCIN (calcitonin) Calcitoninin EVISTA (raloxifene) FORTICAL (calcitonin) EVISTA (raloxifene) FORTICAL (calcitonin) BPH AGENTS ^{AP} A 30-day trial of the preferred agent is required before a non-preferred agent will be approved. Evista will be approved for postmenopausal women with osteoporosis or at high risk for invasive breast cancer.	BONE RESORPTIO	N SUPPRESSION AND RELATED	AGENTS			
FOSAMAX SOLUTION (alendronate) ACTONEL WITH CALCIUM (risedronate/calcium) ATELVIA (risedronate) BONIVA (ibandronate) DIDRONEL (etidronate) FOSAMAX TABLETS (alendronate/vitamin D) ZOMETA (zoledronic acid) OTHER BONE RESORPTION SUPPRESSION AND RELATED AGENTS MIACALCIN (calcitonin) Calcitonin EVISTA (raloxifene) FORTEO (teriparatide) FORTICAL (calcitonin) BPH AGENTS ^{AP} is required before a non-preferred agent will be approved. Is required before a non-preferred agent will be approved. Sequence agent will be approved. Evista will be approved for postmenopausal women with osteoporosis or at high risk for invasive breast cancer.	DONE RECORD THE					
OTHER BONE RESORPTION SUPPRESSION AND RELATED AGENTS MIACALCIN (calcitonin) calcitonin EVISTA (raloxifene) FORTEO (teriparatide) FORTICAL (calcitonin) Colitorin Evista will be approved for postmenopausal women with osteoporosis or at high risk for invasive breast cancer.			ACTONEL WITH CALCIUM (risedronate/calcium) ATELVIA (risedronate) BONIVA (ibandronate) DIDRONEL (etidronate) FOSAMAX TABLETS (alendronate) FOSAMAX PLUS D (alendronate/vitamin D)	is required before a non-preferred		
EVISTA (raloxifene) FORTEO (teriparatide) FORTICAL (calcitonin) postmenopausal women with osteoporosis or at high risk for invasive breast cancer.		OTHER BONE RESORPTION SUPP	PRESSION AND RELATED AGENTS			
		MIACALCIN (calcitonin)	EVISTA (raloxifene) FORTEO (teriparatide)	postmenopausal women with osteoporosis or at high risk for		
	BPH AGENTSAP	BPH AGENTS ^{AP}				
		5-ALPHA-REDUCTA	SE (5AR) INHIBITORS			



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	AVODART (dutasteride) finasteride	PROSCAR (finasteride)	Thirty (30) day trials each of at least two (2) chemically distinct preferred agents, including the generic formulation of a requested non-preferred agent, are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present.
	ALPHA B	LOCKERS	
	doxazosin tamsulosin terazosin	alfuzosin CARDURA (doxazosin) CARDURA XL (doxazosin) FLOMAX (tamsulosin) HYTRIN (terazosin) RAPAFLO (silodosin) UROXATRAL (alfuzosin)	
	5-ALPHA-REDUCTASE (5AR) INHIBIT	ORS/ALPHA BLOCKER COMBINATION	
		JALYN (dutasteride/tamsulosin)	Thirty (30) day trials of dutasteride and tamsulosin concurrently are required before the non-preferred agent will be approved.
BRONCHODII ATO	RS & RESPIRATORY DRUGS		
		LINERGIC	
	ATROVENT HFA (ipratropium) ipratropium SPIRIVA (tiotropium)		Thirty (30) day trials each of the preferred agents in the corresponding group are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present.
	ANTICHOLINERGIC-BETA	AGONIST COMBINATIONS	
	COMBIVENT (albuterol/ipratropium)	albuterol/ipratropium DUONEB (albuterol/ipratropium)	For severely compromised patients, albuterol/ipratropium will be approved if the combined volume of albuterol and ipratropium nebules is inhibitory.



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	PDE4 IN	HIBITOR	
		DALIRESP (roflumilast)	Daliresp will be approved when the following criteria are met: 1. Patient is ≥ forty (40) years of age and 2. Diagnosis of severe chronic obstructive pulmonary disease (COPD) associated with chronic bronchitis and multiple exacerbations requiring systemic glucocorticoids in the preceding six (6) months and 3. Concurrent therapy with an inhaled corticosteroid and longacting bronchodilator and evidence of compliance and 4. No evidence of moderate to severe liver impairment (Child-Pugh Class B or C) and 5. No concurrent use with strong cytochrome P450 inhibitors (rimampicin, phenobarbital, carbamazepine or phenytoin).
	-	N SOLUTION	
	albuterol 2.5mg/0.5mL	ACCUNEB (albuterol)** albuterol 0.63mg & 1.25mg/3mL ^{AP} BROVANA (arformoterol) levalbuterol metaproterenol PERFOROMIST (formoterol) XOPENEX (levalbuterol)	Thirty (30) day trials each of the chemically distinct preferred agents in their corresponding groups are required before a non-preferred agent in that group will be authorized unless one of the exceptions on the PA form is present. **No PA is required for ACCUNEB for children up to 5 years of age.
INHALERS, LONG-ACTING			. , , ,
	FORADIL (formoterol) SEREVENT (salmeterol)	ARCAPTA (indacaterol maleate) ^{NR}	
	INHALERS, SI	HORT-ACTING	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	MAXAIR (pirbuterol) PROAIR HFA (albuterol) PROVENTIL HFA (albuterol) VENTOLIN HFA (albuterol)	XOPENEX HFA (levalbuterol)	Xopenex Inhalation Solution will be approved for 12 months for a diagnosis of asthma or COPD for patients on concurrent asthma controller therapy (either oral or inhaled) with documentation of failure on a trial of albuterol or documented intolerance of albuterol, or for concurrent diagnosis of heart disease.
	OF	RAL	
	albuterol terbutaline	metaproterenol VOSPIRE ER (albuterol)	
CALCIUM CHANNE	EL BLOCKERS ^{AP}		
	LONG-	ACTING	
	amlodipine	ADALAT CC (nifedipine)	Fourteen (14) day trials each of the
	diltiazem XR, XT felodipine ER nifedipine ER nisoldipine verapamil ER	CALAN SR (verapamil) CARDENE SR (nicardipine) CARDIZEM CD, LA, SR (diltiazem) COVERA-HS (verapamil) DILACOR XR (diltiazem) DYNACIRC CR (isradipine) ISOPTIN SR (verapamil) NORVASC (amlodipine) PLENDIL (felodipine) PROCARDIA XL (nifedipine) SULAR (nisoldipine) TIAZAC (diltiazem) VERELAN/VERELAN PM (verapamil)	preferred agents are required before a non-preferred agent will be approved unless one of the exceptions on the PA form is present.
	SHORT	-ACTING	
	diltiazem verapamil	CALAN (verapamil) CARDENE (nicardipine) CARDIZEM (diltiazem) DYNACIRC (isradipine) isradipine nicardipine nimodipine nifedipine NIMOTOP (nimodipine) PROCARDIA (nifedipine)	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA	
CEPHALOSPORIN	S AND RELATED ANTIBIOTICS (Or	al) ^{ap}		
	BETA LACTAMS AND BETA LACTAM/BETA	A-LACTAMASE INHIBITOR COMBINATIONS		
	amoxicillin/clavulanate	amoxicillin/clavulanate ER AUGMENTIN XR (amoxicillin/clavulanate) MOXATAG (amoxicillin)	A five (5) day trial of the preferred agent is required before a non-preferred agent is authorized unless one of the exceptions on the PA form is present.	
	CEPHALO	SPORINS		
	cefaclor cefadroxil cefdinir cefditoren cefpodoxime cefprozil cefuroxime cephalexin SPECTRACEF (cefditoren)	CECLOR (cefaclor) CEDAX (ceftibuten) CEFTIN (cefuroxime) CEFZIL (cefprozil) DURICEF (cefadroxil) KEFLEX (cephalexin) OMNICEF (cefdinir) PANIXINE (cephalexin) RANICLOR (cefaclor) SUPRAX (cefixime) VANTIN (cefpodoxime)		
COUGH & COLD/15	ST GENERATION ANTIHISTAMINES			
	ANTIHISTAMINES	, 1 ST GENERATION		
	chlorpheniramine clemastine diphenhydramine		See posted list of covered NDCs.	
	dextromethorphan HBR/promethazine		See posted list of covered NDCs.	
	ANTIHISTAMINE-ANTITUSSIVE-DECONGESTANT COMBINATIONS			
	brompheniramine/dextromethorphan HBR/pseudoephedrine chlorpheniramine/dextromethorphan/ pseudoephedrine		See posted list of covered NDCs.	
	ANTITUSSIVE-N	ION-NARCOTIC		



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA	
	DELSYM (dextromethorphan polistirex)		See posted list of covered NDCs.	
	DECONG	ESTANTS		
	phenylephrine pseudoephedrine		See posted list of covered NDCs.	
	ANTITUSSIVES/I	EXPECTORANTS		
	guaifenesin guaifenesin/dextromethorphan		See posted list of covered NDCs.	
	DECONGESTANT-ANTIHISTAMINE-	ANTICHOLINERGIC COMBINATIONS		
	pseudoephedrine/chlorpheniramine/ scopolamine syrup		See posted list of covered NDCs.	
	DECONGESTANT-ANTIHIS	STAMINE COMBINATIONS		
	phenylephrine HCL/chlorpheniramine maleate syrup/drops phenylephrine HCL/promethazine syrup		See posted list of covered NDCs.	
	NARCOTIC ANTITUSSIVE-EX	PECTORANT COMBINATION		
CYTOKINE & CAM				
	ENBREL (etanercept) HUMIRA (adalimumab)	CIMZIA (certolizumab/pegol) KINERET (anakinra) ORENCIA (abatacept) SUBCUTANEOUS SIMPONI (golimumab)	Thirty day trials of each of the preferred agents are required before a non-preferred agent will be approved. See additional criteria for treatment	
			of psoriasis or psoriatic arthritis at http://www.dhhr.wv.gov/bms/Pharm acy/Pages/pac.aspx	
ERYTHROPOIESIS	ERYTHROPOIESIS STIMULATING PROTEINS ^{CL}			
	PROCRIT (rHuEPO)	ARANESP (darbepoetin) EPOGEN (rHuEPO)	A thirty (30) day trial of the preferred agent is required before a non-preferred agent will be approved.	
			Prior authorization will be given for the erythropoesis agents if the following criteria are met:	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA		
			 Hemoglobin or Hematocrit less than 10/30 respectively. For renewal, hemoglobin or hematocrit levels greater than 12/36 will require dosage reduction or discontinuation. Exceptions will considered on an individual basis after medical documentation is reviewed. (Laboratory values must be dated within six (6) weeks of request.) Transferrin saturation ≥ 20%, ferritin levels ≥100 mg/ml, or on concurrent therapeutic iron therapy. (Laboratory values must be dated within three (3) weeks of request. For re-authorization, transferrin saturation or ferritin levels are not required if the patient has been responsive to the erythropoietin agent. For HIV-infected patients, endogenous serum erythropoietin level must be ≤ 500mU/ml to initiate therapy. No evidence of untreated GI bleeding, hemolysis, or Vitamin B-12, iron or folate deficiency. 		
FLUOROQUINOLO	FLUOROQUINOLONES (Oral) ^{AP}				
	CIPRO (ciprofloxacin) Suspension ciprofloxacin ciprofloxacin ER levofloxacin	AVELOX (moxifloxacin) CIPRO (ciprofloxacin) Tablets CIPRO XR (ciprofloxacin) FACTIVE (gemifloxacin) FLOXIN (ofloxacin) LEVAQUIN (levofloxacin) NOROXIN (norfloxacin) ofloxacin	A five (5) day trial of one of the preferred agents is required before a non-preferred agent will be approved unless one of the exceptions on the PA form is present.		



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
		PROQUIN XR (ciprofloxacin)	
GENITAL WARTS	AGENTS		
	ALDARA (imiquimod)	CONDYLOX (podofilox) imiquimod podofilox VEREGEN (sinecatechins) ZYCLARA (imiquimod)	A thirty (30) day trial of the preferred agent is required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present. Zyclara will be approved for a diagnosis of actinic keratosis.
GLUCOCORTICOID	DS (Inhaled) ^{AP}		
		ORTICOIDS	
	AEROBID (flunisolide) AEROBID-M (flunisolide) ASMANEX (mometasone) FLOVENT HFA (fluticasone) FLOVENT Diskus (fluticasone) QVAR (beclomethasone)	ALVESCO (ciclesonide) budesonide PULMICORT (budesonide)*	Thirty (30) day trials each of the preferred agents are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present. Pulmicort Respules do not require a prior authorization for children through 8 years of age or for individuals unable to use an MDI. When children who have been stabilized on Pulmicort Respules reach age 9, prescriptions for the Pulmicort inhaler will be authorized for them. *For children less than 9 years of age and for those who meet the PA requirements, brand Pulmicort is preferred over the generic.
		HODILATOR COMBINATIONS	
	ADVAIR (fluticasone/salmeterol) ADVAIR HFA (fluticasone/salmeterol) DULERA (mometasone/formoterol) SYMBICORT(budesonide/formoterol)		



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
GLUCOCORTICOID	OS (Topical)		
		HIGH POTENCY	
	betamethasone dipropionate cream/ointment betamethasone dipropionate/propylene glycol betamethasone valerate ointment clobetasol propionate cream/gel/ointment/solution clobetasol propionate/emollient desoximetasone cream/gel/ointment fluocinonide halobetasol propionate triamcinolone acetonide 0.5%	amcinonide APEXICON (diflorasone diacetate) APEXICON E (diflorasone diacetate) betamethasone dipropionate gel clobetasol propionate foam CLOBEX (clobetasol propionate) CORMAX (clobetasol propionate) diflorasone diacetate diflorasone diacetate/emollient DIPROLENE (betamethasone dipropionate/propylene glycol) DIPROLENE AF (betamethasone dipropionate/propylene glycol) DIPROSONE (betamethasone dipropionate) fluocinonide/emollient halcinonide HALOG (halcinonide) KENALOG 0.5% (triamcinolone acetonide) LIDEX (fluocinonide) LUXIQ (betamethasone valerate) OLUX (clobetasol propionate/emollient) PSORCON (diflorasone diacetate) TEMOVATE (clobetasol propionate) TEMOVATE-E (clobetasol propionate) ULTRAVATE (halobetasol propionate) VANOS (fluocinonide)	Five day trials of one form of each preferred unique active ingredient in the corresponding potency group are required before a non-preferred agent will be approved.
		POTENCY	
	betamethasone dipropionate lotion betamethasone valerate cream desoximetasone 0.05%cream fluocinolone acetonide 0.025% fluticasone propionate hydrocortisone valerate	ARISTOCORT (triamcinolone) betamethasone valerate lotion BETA-VAL (betamethasone valerate) CLODERM (clocortolone pivalate) CORDRAN/CORDRAN SP (flurandrenolide) CUTIVATE (fluticasone propionate)	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	mometasone furoate triamcinolone acetonide 0.025% and 0.1%	DERMATOP (prednicarbate) ELOCON (mometasone furoate) hydrocortisone butyrate hydrocortisone butyrate/emollient KENALOG 0.1% (triamcinolone acetonide) LOCOID (hydrocortisone butyrate) LOCOID LIPOCREAM (hydrocortisone butyrate/emollient) prednicarbate TOPICORT LP (desoximetasone) TRIDERM (triamcinolone acetonide) WESTCORT (hydrocortisone valerate)	
	LOW PO	OTENCY	
	desonide fluocinolone acetonide 0.01% hydrocortisone 0.5%, 1%, 2.5% hydrocortisone acetate 0.5%, 1% (Rx & OTC)	ACLOVATE (alclometasone dipropionate) alclometasone dipropionate CAPEX (fluocinolone acetonide) DERMA-SMOOTHE FS (fluocinolone acetonide) DESONATE (desonide) DESOWEN (desonide) LOKARA (desonide) PANDEL (hydrocortisone probutate) VERDESO (desonide)	
GROWTH HORMO	NE ^{CL}		
	GENOTROPIN (somatropin) NORDITROPIN NORDIFLEX (somatropin) NORDITROPIN FLEXPRO (somatropin) NUTROPIN AQ NUSPIN (somatropin)	HUMATROPE (somatropin) INCRELEX (mecasermin) NUTROPIN (somatropin) NUTROPIN AQ (somatropin) OMNITROPE (somatropin) SAIZEN (somatropin) SEROSTIM (somatropin) TEV-TROPIN (somatropin) ZORBTIVE (somatropin)	The preferred agents must be tried before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present. Patients already on a non-preferred agent will receive authorization to continue therapy on that agent for the duration of the existing PA.
H. PYLORI COMBI	NATION TREATMENTS		the duration of the existing 17t.
	Please use individual components: preferred PPI (Dexilant, omeprazole or pantoprazole) amoxicillin tetracycline	HELIDAC (bismuth/metronidazole/tetracycline) PREVPAC (lansoprazole/amoxicillin/clarithromycin) PYLERA	A trial of all the individual preferred components (with Dexilant, omeprazole or pantoprazole) at the recommended dosages, frequencies and duration is required before the



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	metronidazole clarithromycin bismuth	(bismuth/metronidazole/tetracycline)	brand name combination packages will be approved unless one of the exceptions on the PA form is present.
HEPATITIS B TREA	ATMENTS		
	EPIVIR HBV (lamivudine) HEPSERA (adefovir) TYZEKA (telbivudine)	BARACLUDE (entecavir)	A thirty (30) day trial of one of the preferred agents is required before the non-preferred agent will be authorized unless one of the exceptions on the PA form is present.
HEPATITIS C TREA	ATMENTSCL		
	INCIVEK (telaprevir) ^{CL} PEGASYS (pegylated interferon) PEG-INTRON (pegylated interferon) ribavirin VICTRELIS (boceprevir) ^{CL}	COPEGUS (ribavirin) INFERGEN (consensus interferon) REBETOL (ribavirin) RIBAPAK DOSEPACK (ribavirin) RIBASPHERE (ribavirin)	Patients starting therapy in this class must try the preferred agent of a dosage form before a non-preferred agent of that dosage form will be authorized. See additional criteria for Incivek and Victrelis at http://www.dhhr.wv.gov/bms/Pharmacy/Pages/pac.aspx
HYPERURICEMIA A	AND GOUT AGENTS		<u> </u>
	ANTIMI	тотісѕ	
		COLCRYS (colchicine)*	A thirty-day trial of one of the preferred agents for the prevention of gouty arthritis attacks (colchicine/probenecid, probenecid, or allopurinol) is required before a non-preferred agent will be approved unless one of the exceptions on the PA form is present. *In the case of acute gouty attacks, a 10-day supply (20 tablets) of Colcrys will be approved per 90



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA		
	ANTIMITOTIC-URICOSURIC COMBINATION				
	colchicine/probenecid				
		DSURIC			
	probenecid				
	XANTHINE OXID	ASE INHIBITORS			
	allopurinol	ULORIC (febuxostat) ZYLOPRIM (allopurinol)			
HYPOGLYCEMICS	, INCRETIN MIMETICS/ENHANCER	S			
	INJEC	TABLE			
		BYETTA (exenatide) SYMLIN (pramlintide) VICTOZA (liraglutide)	Byetta, Symlin, and Victoza will be subject to the following clinical edits: Byetta and Victoza will be approved with a previous history of a thirty (30) day trial of an oral agent (sulfonylurea, thiazolindinedione (TZD) and/ or metformin) and no evidence of concurrent insulin therapy. Symlin- History of insulin utilization in the past 90 days. No gaps in insulin therapy greater than 30 days.		
	ORAL ^{Ar}				
	JANUMET (sitagliptin/metformin) JANUVIA (sitagliptin) JUVISYNC (sitagliptin/simvastatin) KOMBIGLYZE XR (saxagliptin/metformin) ONGLYZA (saxagliptin) TRADJENTA (linagliptin)		Januvia/Janumet/Juvisync, Onglyza/Kombiglyze XR and Tradjenta will be subject to the following edits: 1.Previous history of a 30-day trial of an oral agent (sulfonylurea, thiazolidinedione (TZD) or metformin)		



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA	
			2.Onglyza/Kombiglyze XR will not be approved for concurrent use with insulin. 3.Januvia/Janumet /Juvisync will be approved for concurrent use with insulin for six (6) month intervals. For re-authorization, HgBA1C levels must be less than or equal (≤) to 7. Current laboratory values must be submitted.	
HYPOGLYCEMICS	, INSULINS			
	HUMALOG (insulin lispro) vials HUMALOG PEN/KWIKPEN (insulin lispro) HUMALOG MIX (insulin lispro/lispro protamine) vials only HUMULIN (insulin) vials only LANTUS (insulin glargine) all forms LEVEMIR (insulin detemir) all forms NOVOLIN (insulin) all forms NOVOLOG (insulin aspart) all forms NOVOLOG MIX all forms (insulin aspart/aspart protamine)	APIDRA (insulin glulisine) ^{AP} HUMALOG MIX PENS (insulin lispro/lispro protamine) HUMULIN PEN (insulin)	 To receive Apidra, patients must meet the following criteria: be 4 years or older; be currently on a regimen including a longer-acting or basal insulin. have had a trial of a similar preferred agent, Novolog or Humalog, with documentation that the desired results were not achieved. 	
HYPOGLYCEMICS				
	-	TINIDES	A 11: ((22) 1 () ()	
	PRANDIN (repaglinide) STARLIX (nateglinide)	nateglinide	A thirty (30) day trial of the preferred agent is required before a non-preferred agent will be authorized, unless one of the exceptions on the PA form is present.	
		PRANDIMET (repaglinide/metformin)		
HYPOGLYCEMICS	, TZDS			
	THIAZOLID	INEDIONES		



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	ACTOS (pioglitazone)	AVANDIA (rosiglitazone) ^{AP}	Treatment naïve patients require a two (2) week trial of Actos before Avandia will be authorized, unless one of the exceptions on the PA form is present.
	TZD COME	BINATIONS	
		ACTOPLUS MET (pioglitazone/ metformin) ACTOPLUS MET XR (pioglitazone/ metformin) AVANDAMET (rosiglitazone/metformin) AVANDARYL (rosiglitazone/glimepiride) DUETACT (pioglitazone/glimepiride)	Patients are required to use the components of Actoplus Met and Duetact separately. Exceptions will be handled on a case-by-case basis.
IMPETIGO AGENTS	S (Topical)		
	bacitracin gentamicin sulfate mupirocin	ALTABAX (retapamulin) BACTROBAN (mupirocin) CORTISPORIN (bacitracin/neomycin/polymyxin/HC)	Ten (10) day trials of at least one preferred agent, including the generic formulation of a requested non-preferred agent, are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present.
INTRANASAL RHIN	IITIS AGENTS ^{AP}		
		LINERGICS	
	ipratropium	ATROVENT(ipratropium)	Thirty (30) day trials of the preferred nasal anti-cholinergic, an antihistamine, and corticosteroid groups are required before a non-preferred anti-cholinergic will be approved unless one of the exceptions on the PA form is present.
	ANTIHIS	TAMINES	·
	ASTELIN (azelastine) PATANASE (olopatadine)	ASTEPRO (azelastine) azelastine	Thirty (30) day trials of both preferred intranasal antihistamines and a thirty (30) day trial of one of the preferred intranasal corticosteroids are required before the non-preferred agent will be approved unless one of the exceptions on the PA form is



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
			present.
	CORTICO	STEROIDS	
	fluticasone propionate NASACORT AQ (triamcinolone) NASONEX (mometasone)	BECONASE AQ (beclomethasone) flunisolide FLONASE (fluticasone propionate) NASALIDE (flunisolide) NASAREL (flunisolide) OMNARIS (ciclesonide) RHINOCORT AQUA (budesonide) triamcinolone VERAMYST (fluticasone furoate)	Thirty (30) day trials of each preferred agent in the corticosteroid group are required before a non-preferred corticosteroid agent will be authorized unless one of the exceptions on the PA form is present.
LEUKOTRIENE MO	DIFIERS		
	ACCOLATE (zafirlukast) SINGULAIR (montelukast)	zafirlukast ZYFLO (zileuton)	Thirty (30) day trials each of the preferred agents are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present.
LIPOTROPICS, OTI	HER (Non-statins) ^{AP}		
	BILE ACID SE	QUESTRANTS	
	cholestyramine colestipol WELCHOL (colesevelam) ^{AP}	COLESTID (colestipol) QUESTRAN (cholestyramine)	A twelve (12) week trial of one of the preferred agents is required before a non-preferred agent in the corresponding category will be authorized. Welchol will be approved for add-on therapy for type 2 diabetes when there is a previous history of a 30-day trial of an oral agent (sulfonylurea, thiazolidinedione (TZD) or metformin).
	CHOLESTEROL ABSO	ORPTION INHIBITORS	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
		ZETIA (ezetimibe)	Zetia, as monotherapy, will only be approved for patients who cannot take statins or other preferred agents. AP does not apply.
			Zetia will be approved for add-on therapy only after an insufficient response to the maximum tolerable dose of a statin after 12 weeks of therapy. AP does not apply.
	FATTY	ACIDS	
	LOVAZA (omega-3-acid ethyl esters) ^{AP}		Lovaza will be approved when the patient is intolerant or not responsive to, or not a candidate for nicotinic acid or fibrate therapy.
	FIBRIC ACID	DERIVATIVES	
	fenofibrate 54mg & 160mg fenofibrate micronized 67mg, 134mg & 200mg gemfibrozil TRICOR (fenofibrate nanocrystallized) TRILIPIX (fenofibric acid)	ANTARA (fenofibrate) FENOGLIDE (fenofibrate) FIBRICOR (fenofibric acid) fenofibrate nanocrystallized 145mg LIPOFEN (fenofibrate) LOFIBRA (fenofibrate) LOPID (gemfibrozil) TRIGLIDE (fenofibrate)	
	NIA	CIN	
	niacin NIASPAN (niacin)	NIACELS (niacin) NIACOR (niacin) NIADELAY (niacin) SLO-NIACIN (niacin)	
LIPOTROPICS, ST.	ATINS ^{AP}		
	STA	TINS	
	CRESTOR (rosuvastatin) LESCOL (fluvastatin) LESCOL XL (fluvastatin) LIPITOR (atorvastatin) lovastatin pravastatin simvastatin ^{CL*}	ALTOPREV (lovastatin) atorvastatin LIVALO (pitavastatin) MEVACOR (lovastatin) PRAVACHOL (pravastatin) ZOCOR (simvastatin)	Twelve (12) week trials each of two (2) of the preferred statins, including the generic formulation of a requested non-preferred agent, are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

STATIN COMBINATIONS ADVICOR (lovastatin/niacin) CADUET (atorvastatin/amlodipine) SIMCOR (simvastatin/niacin ER) VYTORIN (simvastatin/ ezetimibe) VYTORIN (sinvastatin/ ezetimibe) VYTORIN (simvastatin/ ezetimibe) VYTORIN (sinvastatin/ ezetimibe) Vytorin 801 set in sufficient response to the maximum tolerable dose of Lipitor (trosustatin) exit in sufficient response to the maximum tolerable dose of Lipitor (trosustatin) execution of the exceptions on the PA form is present. Five (5) day trials each of the preferred agent will be authorized under sufficient response to the maximum tolerable dose of Lipitor (trosustatin) exerting the preferred agent sufficient response to the maximum tolerable dose of Lipitor (trosustatin) exerting the sufficient response to the maximum tolerable dos	THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
STATIN COMBINATIONS ADVICOR (lovastatin/niacin) CADUET (atorvastatin/amlodipine) SIMCOR (simvastatin/hiacin ER) VYTORIN (simvastatin/ ezetimibe) CL* VYtorin will be approved only after an insufficient response to the maximum tolerable dose of Lipitor (atorvastatin) or Crestor (rosuvastatin) after 12 weeks, unless one of the exceptions on the PA form is present. *Vytorin 80/10mg tablets will require a clinical PA MACROLIDES KETEK (telithromycin) **Requests for telithromycin will be authorized if there is documentation of the use of any antibiotic within the past 28 days. **MACROLIDES** **BIAXIN (clarithromycin) clarithromycin erythromycin erythromycin ERYED (erythromycin) ERYPED (erythro				present. *Zocor/simvastatin 80mg tablets will
ADVICOR (lovastatin/niacin) CADUET (atorvastatin/amlodipine) SIMCOR (simvastatin/amlodipine) SIMCOR (simvastatin/amlodipine) SIMCOR (simvastatin/amlodipine) SIMCOR (simvastatin/niacin ER) WACROLIDES (Oral) KETOLIDES KETEK (telithromycin) BIAXIN (clarithromycin) clarithromycin erythromycin erythromycin E.E.S. (erythromycin) ERYPED (erythromycin)				require a clinical PA
CADUET (atorvastatin/aniodipine) SIMCOR (simvastatin/niacin ER) ### Acrolides Macrolides Macrolides				
KETOLIDES MACROLIDES BIAXIN (clarithromycin) Five (5) day trials each of the preferred agents are required before a non-preferred agent will be authorized unless one of the preferred agent will be authorized unless one of the preferred agent will be authorized unless one of the E.E.S. (erythromycin ethylsuccinate) E-MYCIN (erythromycin) ERYPED (erythromycin ethylsuccinate) ERYPED (erythromycin) ERY-TAB (erythromycin) ERY-TAB (erythromycin)		CADUET (atorvastatin/amlodipine)	VYTORIN (simvastatin/ ezetimibe)	an insufficient response to the maximum tolerable dose of Lipitor (atorvastatin) or Crestor (rosuvastatin) after 12 weeks, unless one of the exceptions on the PA form is present. *Vytorin 80/10mg tablets will
KETEK (telithromycin) Requests for telithromycin will be authorized if there is documentation of the use of any antibiotic within the past 28 days. MACROLIDES BIAXIN (clarithromycin)	MACROLIDES/KET	OLIDES (Oral)		
MACROLIDES azithromycin clarithromycin erythromycin erythromycin E.E.S. (erythromycin) ERYC (erythromycin) ERYC (erythromycin) ERY-TAB (erythromycin) azithrorized if there is documentation of the use of any antibiotic within the past 28 days. Five (5) day trials each of the preferred agents are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present.		КЕТО		
azithromycin clarithromycin erythromycin BIAXIN (clarithromycin) Erythromycin BIAXIN XL (clarithromycin) clarithromycin ER DIFICID (fidaxomicin) ^{NR} E.E.S. (erythromycin ethylsuccinate) E-MYCIN (erythromycin) ERYC (erythromycin) ERYPED (erythromycin) ERY-TAB (erythromycin) ERY-TAB (erythromycin) Erythromycin			KETEK (telithromycin)	authorized if there is documentation of the use of any antibiotic within
clarithromycin erythromycin BIAXIN XL (clarithromycin) preferred agents are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present. ENYCIN (erythromycin) ERYPED (erythromycin) ethylsuccinate) ERY-TAB (erythromycin)		MACRO	OLIDES	
ERYTHROCIN (erythromycin stearate) erythromycin estolate PCE (erythromycin) ZITHROMAX (azithromycin) ZMAX (azithromycin)		clarithromycin	BIAXIN XL (clarithromycin) clarithromycin ER DIFICID (fidaxomicin) E.E.S. (erythromycin ethylsuccinate) E-MYCIN (erythromycin) ERYC (erythromycin) ERYPED (erythromycin ethylsuccinate) ERY-TAB (erythromycin) ERYTHROCIN (erythromycin stearate) erythromycin estolate PCE (erythromycin) ZITHROMAX (azithromycin)	preferred agents are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is
MULTIPLE SCLEROSIS AGENTS ^{CL, AP}	MULTIPLE SCLERO	OSIS AGENTSCL, AP		



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THEDADELITIO			
THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	INTERF	ERONS	
	AVONEX (interferon beta-1a) BETASERON (interferon beta-1b) REBIF (interferon beta-1a)	EXTAVIA (interferon beta-1b)	A 30-day trial of a preferred agent will be required before a non-preferred agent will be approved.
	NON-INTE		
	COPAXONE (glatiramer)	AMPYRA (dalfampridine) CL* GILENYA (fingolimod) CL** TYSABRI (natalizumab)***	A 30-day trial of the preferred agent will be required before a non-preferred agent will be approved. *Amypra will be prior authorized if the following conditions are met: 1. Diagnosis of multiple sclerosis 2. No history of seizures 3. No evidence of moderate or severe renal impairment 4. Initial prescription will be approved for 30 days only. ** Gilenya: PA Criteria 1) A diagnosis of a relapsing form of multiple sclerosis AND 2) Medication is prescribed by a neurologist AND 3) History of a thirty (30) trial of one of the preferred agents for multiple sclerosis unless one of the exceptions on the PA form is present AND 4) Dosage is limited to one tablet per day. (AP does not apply.) ***Tysabri will only be approved for members who are enrolled in the TOUCH Prescribing Program. AP does not apply.
MUSCLE RELAXAN	NTS (Oral) ^{AP}		



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA		
	ACUTE MUSCULOSKELETAL RELAXANT AGENTS				
	chlorzoxazone cyclobenzaprine methocarbamol	AMRIX (cyclobenzaprine) carisoprodol carisoprodol/ASA carisoprodol/ASA/codeine cyclobenzaprine ER FEXMID (cyclobenzaprine) FLEXERIL (cyclobenzaprine) metaxalone methocarbamol/ASA orphenadrine orphenadrine/ASA/caffeine PARAFON FORTE DSC (chlorzoxazone) ROBAXIN (methocarbamol) SKELAXIN (metaxalone) SOMA (carisoprodol) SOMA COMPOUND (carisoprodol/ASA) SOMA COMP w/ COD (carisoprodol/ASA/ codeine) F AGENTS USED FOR SPASTICITY DANTRIUM (dantrolene) ZANAFLEX (tizanidine)	Thirty (30) day trials of the preferred acute musculoskeletal relaxants are required before a non-preferred acute musculoskeletal agent will be approved, with the exception of carisoprodol. Thirty (30) day trials of the preferred acute musculoskeletal relaxants and Skelaxin are required before carisoprodol will be approved. Thirty (30) day trials of the preferred skeletal muscle relaxants		
NSAIDS ^{AP}	tizanidine	ZANAFLEX (IIZanidine)	associated with the treatment of spasticity (are required before non-preferred agents will be approved unless one of the exceptions on the PA form is present.		
NSAIDS	Nov on				
		LECTIVE			
	diclofenac etodolac fenoprofen flurbiprofen ibuprofen (Rx and OTC) INDOCIN (indomethacin) (suspension only) indomethacin ketorolac naproxen (Rx only) oxaprozin sulindac	ADVIL (ibuprofen) ANAPROX (naproxen) ANSAID (flurbiprofen) CAMBIA (diclofenac) CATAFLAM (diclofenac) CLINORIL (sulindac) DAYPRO (oxaprozin) FELDENE (piroxicam) INDOCIN (indomethacin) ketoprofen ketoprofen ER	Thirty (30) day trials of each of the preferred agents are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present.		



EFFECTIVE 01/01/12 Version 2012.4

This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
		LODINE (etodolac) meclofenamate mefenamic acid MOTRIN (ibuprofen) nabumetone NALFON (fenoprofen) NAPRELAN (naproxen) NAPROSYN (naproxen) NUPRIN (ibuprofen) ORUDIS (ketoprofen) piroxicam PONSTEL (meclofenamate) SPRIX (ketorolac) tolmetin VOLTAREN (diclofenac) ZIPSOR (diclofenac potassium)	
	NSAID/GI PROTECT	ANT COMBINATIONS	
		ARTHROTEC (diclofenac/misoprostol) VIMOVO (naproxen/esomeprazole)	
	COX-II SI	ELECTIVE	
	meloxicam	CELEBREX (celecoxib) CL MOBIC (meloxicam)	Requests for COX-2 Inhibitor agents will be authorized if the following criteria are met: Agent is requested for treatment of a chronic condition, and a. Patient is greater than or equal to 70 years of age, or b. Patient is currently on anticoagulation therapy, or c. Patient has a history or risk of a serious GI complication.
OPHTHALMIC ANT	IBIOTICS (FLUOROQUINOLONES	& SELECT MACROLIDES) ^{AP}	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	ciprofloxacin MOXEZA (moxifloxacin) ofloxacin VIGAMOX (moxifloxacin) **The American Academy of Ophthalmology guidelines on treating bacterial conjunctivitis recommend as first line treatment options: erythromycin ointment, sulfacetamide drops, or polymyxin/trimethoprim drops. Alternative treatments include bacitracin ointment, sulfacetamide ointment, polymyxin/bacitracin ointment, fluoroquinolone drops, or azithromycin drops. All generic forms of ophthalmic erythromycin, sulfacetamide, and polymyxin/trimethoprim, polymyxin/bacitracin and bacitracin are preferred.	AZASITE (azithromycin) BESIVANCE (besifloxacin) CILOXAN (ciprofloxacin) levofloxacin OCUFLOX (ofloxacin) QUIXIN (levofloxacin) ZYMAXID (gatifloxacin)	Five (5) day trials of each of the preferred agents are required before non-preferred agents will be authorized unless one of the exceptions on the PA form is present. **A prior authorization is required for the fluoroquinolone agents for patients under 21 years of age unless there has been a trial of a first line treatment option within the past 10 days.
OPHTHALMIC ANT	I-INFLAMMATORIES		
	flurbiprofen ketorolac 0.4% NEVANAC (nepafenac)	ACULAR LS (ketorolac) ACUVAIL 0.45% (ketorolac tromethamine) AP BROMDAY (bromfenac) diclofenac AP DUREZOL (difluprednate) AP LOTEMAX (loteprednol) VEXOL (rimexolone) XIBROM (bromfenac)	Five (5) day trials of each of the preferred ophthalmic anti-inflammatory agents are required before nonpreferred agents will be authorized unless one of the exceptions on the PA form is present.
OPHTHALMICS FO	R ALLERGIC CONJUNCTIVITIS		
	ALAWAY (ketotifen) ALREX (loteprednol) cromolyn ketorolac 0.5% PATADAY (olopatadine) PATANOL (olopatadine) ZADITOR OTC (ketotifen)	ACULAR (ketorolac) ALAMAST (pemirolast) AP ALOCRIL (nedocromil) AP ALOMIDE (lodoxamide) AP azelastine BEPREVE (bepotastine) AP CROLOM (cromolyn) AP DUREZOL (difuprednate) NR ELESTAT (epinastine) AP EMADINE (emedastine) AP epinastine ketotifen	Thirty (30) day trials of each of three (3) of the preferred agents are required before non-preferred agents will be authorized, unless one of the exceptions on the PA form is present.



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
		LASTACAFT (alcaftadine) OPTICROM (cromolyn) OPTIVAR (azelastine) ZYRTEC ITCHY EYE (ketotifen) AP	
OPHTHALMICS, GL	AUCOMA AGENTS		
	COMBINATI	ON AGENTS	
	COMBIGAN (brimonidine/timolol) dorzolamide/timolol	COSOPT (dorzolamide/timolol)	Authorization for a non-preferred agent will only be given if there is an allergy to the preferred agents.
	BETA BL	OCKERS	
	betaxolol BETOPTIC S (betaxolol) carteolol levobunolol metipranolol timolol	BETAGAN (levobunolol) BETIMOL (timolol) ISTALOL (timolol) OPTIPRANOLOL (metipranolol) TIMOPTIC (timolol)	
	CARBONIC ANHYD	DRASE INHIBITORS	
	AZOPT (brinzolamide) dorzolamide	TRUSOPT (dorzolamide)	
	PARASYMPA ⁻	THOMIMETICS	
	CARBOPTIC (carbachol) ISOPTO CARBACHOL (carbachol) PHOSPHOLINE IODIDE (echothiophate iodide) pilocarpine	ISOPTO CARPINE (pilocarpine) PILOPINE HS (pilocarpine)	
	PROSTAGLAN	IDIN ANALOGS	
	latanoprost LUMIGAN (bimatoprost) TRAVATAN-Z (travoprost)	XALATAN (latanoprost)	
		DMIMETICS	
	ALPHAGAN P (brimonidine) brimonidine 0.2% dipivefrin	brimonidine 0.15% PROPINE (dipivefrin)	
OTIC FLUOROQUIN	NOLONESAP		



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA	
	CIPRODEX (ciprofloxacin/dexamethasone)* ofloxacin	CIPRO HC (ciprofloxacin/hydrocortisone) CETRAXAL 0.2% SOLUTION (ciprofloxacin) FLOXIN (ofloxacin)	Five (5) day trials of each of the preferred agents are required before a non-preferred agent will be approved unless one of the exceptions on the PA form is present. *Ciprodex is limited to patients 8 years of age and younger. Age exceptions will be handled on a case-by-case basis.	
PANCREATIC ENZ	YMES ^{AP}			
	CREON ZENPEP	PANCREAZE PANCRELIPASE 5000	A thirty (30) day trial of a preferred agent is required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present.	
			Non-preferred agents will be approved for members with cystic fibrosis.	
PARATHYROID AG	GENTS ^{ap}			
	calcitriol HECTOROL (doxercalciferol) vitamin d 2 (ergocalciferol) (Rx and OTC)* vitamin d 3 (cholecalciferol) (Rx and OTC)* ZEMPLAR (paricalcitol)	DRISDOL (ergocalciferol) ROCALTROL (calcitriol) SENSIPAR (cinacalcet)	A thirty (30) day trial of a preferred agent will be required before a non-preferred agent will be approved. *See Covered List	
PEDICULICIDES/SO	CABICIDES (Topical) ^{AP}		See Covered List	
	OVIDE (malathion) permethrin (Rx and OTC) pyrethrins-piperonyl butoxide	EURAX (crotamiton) lindane malathion 0.5% lotion NATROBA (spinosad) ULESFIA 5% LOTION (benzyl alcohol)	Trials of the preferred agents (which are age and weight appropriate) are required before non-preferred agents will be approved unless one of the exceptions on the PA form is present.	
PHOSPHATE BINDERS ^{AP}				
	FOSRENOL (lanthanum) RENAGEL (sevelamer) RENVELA (sevelamer carbonate)	calcium acetate ELIPHOS (calcium acetate) PHOSLYRA (calcium acetate) ^{NR}	Thirty (30) day trials of at least two preferred agents are required unless one of the exceptions on the PA form is present.	
PLATELET AGGREGATION INHIBITORS ^{AP}				



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	AGGRENOX (dipyridamole/ASA) cilostazol PLAVIX (clopidogrel)	BRILINTA (ticagrelor) ^{NK} dipyridamole EFFIENT (prasugrel) PERSANTINE (dipyridamole) PLETAL (cilostazol) TICLID (ticlopidine) ticlopidine	A thirty (30) day trial of a preferred agent is required before a non-preferred agent will be approved unless one of the exceptions on the PA form is present. Effient will be approved for acute coronary syndrome when it is to be managed by acute or delayed percutaneous coronary intervention (PCI). Three -day emergency supplies of Effient are available when necessary
PRENATAL VITAM	IINS		
	prenatal vitamin 27 w/calcium/ferrous fumarate/folic acid prenatal vitamins 28 w/calcium/iron ps complex/folic acid prenatal vitamins/ferrous fumarate/docusate/folic acid prenatal vitamins/ferrous fumarate/folic acid prenatal vitamins/ferrous fumarate/folic acid/selenium prenatal vitamins/iron, carbonyl/folic acid/prenatal vitamin no. 15/iron, carbonyl/folic acid/docusate sod prenatal vitamin no. 16/iron, carbonyl/folic acid/docusate sod prenatal vitamin no. 17/iron, carbonyl/folic acid/docusate sod prenatal vitamin no. 18/iron, carbonyl/folic acid/docusate sod prenatal vitamin w-o calcium/ferrous fumarate/folic acid prenatal vitamin w-o vit a/fe carbonyl-fe fumarate/fa	CARENATAL DHA CITRANATAL DHA COMBI RX FOLBECAL DUET/DUET DHA FOLTABS PLUS DHA NATACHEW NATAFORT NATELLE PLUS W/DHA NEEVO NOVANATAL OB-NATAL ONE OPTINATE PRECARE/PRECARE PREMIER PREMESIS PRENATAL RX PRENATAL RX PRENATAL RX PRENATAL U prenatal vitamins/ferrous bis-glycinate chelate/folic acid prenatal vitamins/iron, carbonyl/omega- 3/FA/fat combo no. 1 prenatal vitamins comb no. 20/iron bisgly/folic acid/DHA prenatal vitamins no. 22/iron, carbonyl/FA/docusate/DHA prenatal vitamins w-CA, FE, FA (<1 mg) prenatal vitamins w-o calcium/iron ps	See posted list of covered NDCs. See posted list of covered NDCs.



EFFECTIVE 01/01/12 Version 2012.4

This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC	DEFERRED AGENTS	NON PREFERRED AGENTS	DA ODITEDIA
DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
		complex/FA prenatal vitamins w-o CA no. 5/ferrous fumarate/folic acid prenatal vitamins CMB w-o CA no. 2 prenatal vitamins w-o calcium no. 9/iron/folic acid PRENATE DHA/PRENATE ELITE PRENAVITE PRENEXA PRIMACARE RENATE/RENATE DHA SELECT-OB TANDEM DHA/TANDEM OB	
PROTON PUMP INI			
	DEXILANT (dexlansoprazole) omeprazole pantoprazole	ACIPHEX (rabeprazole) lansoprazole NEXIUM (esomeprazole) NEXIUM PACKETS (esomeprazole) omeprazole/sodium bicarbonate PREVACID capsules (lansoprazole) (Rx and OTC) PREVACID Solu-Tabs (lansoprazole) PRILOSEC (omeprazole) PROTONIX (pantoprazole) ZEGERID OTC (omeprazole)	Sixty (60) day trials of each of the preferred agents, inclusive of a concurrent thirty (30) day trial at the maximum dose of an H₂ antagonist are required before a non-preferred agent will be approved unless one of the exceptions on the PA form is present Prior authorization is not required for Prevacid Solu-Tabs for patients ≤8 years of age.
PSORIATIC AGENT	S - TOPICAL		
	calcipotriene ointment DOVONEX (calcipotriene) TAZORAC (tazarotene)	calcipotriene solution TACLONEX (calcipotriene/betamethasone) VECTICAL (calcitriol)	Thirty (30) day trials of two (2) preferred unique chemical entities are required before non-preferred agents will be approved unless one of the exceptions on the PA form is present.
PULMONARY ANT	HYPERTENSIVES - ENDOTHELIN		
	LETAIRIS (ambrisentan)	TRACLEER (bosentan)	Letairis will be approved for the treatment of pulmonary artery hypertension (PAH) World Health Organization (WHO) Group I to improve exercise ability and decrease the rate of clinical



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	HYPERTENSIVES – PDE5s ^c ∟		deterioration. Tracleer will be approved for the treatment of pulmonary artery hypertension (PAH) (WHO Group I) in patients with World Health Organization (WHO) Class II, III, or IV symptoms to improve exercise capacity and decrease the rate of clinical deterioration AND when there has been a failure with Letairis.
I OZMONANI ANII	ADCIRCA (tadalafil)		
	REVATIO (sildenafil)		
PULMONARY ANTI	HYPERTENSIVES - PROSTACYCL	INS ^{CL}	
	epoprostenol VENTAVIS (iloprost)	FLOLAN (epoprostenol) REMODULIN (treprostinil sodium) TYVASO (treprostinil)	Ventavis will only be approved for the treatment of pulmonary artery hypertension (WHO Group 1) in patients with NYHA Class III or IV symptoms. Remodulin and Tyvaso will be approved only after a 30-day trial of Ventavis unless one of the exceptions on the PA form is present.
SEDATIVE HYPNO			
		AZEPINES	
	temazepam	DALMANE (flurazepam) DORAL (quazepam) estazolam flurazepam HALCION (triazolam) RESTORIL (temazepam) triazolam	Fourteen (14) day trials of the preferred agents in both categories are required before a non-preferred agent will be authorized unless one of the exceptions on the PA form is present.
	отн	IERS	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
DNOG GLAGO	zolpidem	AMBIEN (zolpidem) AMBIEN CR (zolpidem) chloral hydrate EDLUAR SL (zolpidem) LUNESTA (eszopiclone) ROZEREM (ramelteon) SILENOR (doxepin) SOMNOTE (chloral hydrate) SONATA (zaleplon) zaleplon zolpidem tartrate ER ZOLPIMIST SPRAY (zolpidem	
STIMULANTS AND	RELATED AGENTS		
	AMPHE	TAMINES	
	amphetamine salt combination dextroamphetamine VYVANSE (lisdexamfetamine)	ADDERALL (amphetamine salt combination) ADDERALL XR (amphetamine salt combination) amphetamine salt combination ER DESOXYN (methamphetamine) DEXEDRINE (dextroamphetamine) DEXTROSTAT (dextroamphetamine) methamphetamine PROCENTRA (dextroamphetamine) NR	Members currently utilizing Adderall XR as of 1/1/2012 may continue use until 6/30/2012. Except for Strattera, PA is required for adults >18 years. One of the preferred agents in each group (amphetamines and nonamphetamines) must be tried for thirty (30) days before a nonpreferred agent will be authorized. Thirty (30) day trials of at least three (3) antidepressants are required before amphetamines will be approved for depression. Provigil will only be approved for patients >16 years of age with a diagnosis of narcolepsy.
		PHETAMINE	
	CONCERTA (methylphenidate) DAYTRANA (methylphenidate) FOCALIN (dexmethylphenidate) FOCALIN XR (dexmethylphenidate) guanfacine	dexmethylphenidate KAPVAY ER (clonidine) METADATE ER (methylphenidate) methylphenidate ER (Generic Concerta) NUVIGIL (armodafinil)	Strattera will not be approved for concurrent administration with amphetamines or methylphenidates, except for 30 days or less for tapering purposes.



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	INTUNIV (guanfacine extended-release) METADATE CD (methylphenidate) methylphenidate methylphenidate ER STRATTERA (atomoxetine)	pemoline PROVIGIL (modafinil) RITALIN (methylphenidate) RITALIN LA (methylphenidate) RITALIN-SR (methylphenidate)	Strattera is limited to a maximum of 100mg per day. Kapvay will be approved if the following criteria are met: 1. Fourteen (14) day trials of at least one preferred product from the amphetamine and non-amphetamine class and 2. A fourteen (14) day trial of Strattera and 3. A fourteen (14) day trial of clonidine (for Kapvay) unless one of the exceptions on the PA form is present or 4. In cases of a diagnosis of Tourette's syndrome, tics, autism or disorders included in the autism spectrum, only a fourteen (14) day trial of clonidine (for Kapvay) is required for approval.
TETRACYCLINES ^A	P		
	doxycycline hyclate minocycline capsules tetracycline	ADOXA (doxycycline monohydrate) demeclocycline* DORYX (doxycycline hyclate) doxycycline hyclate delayed release doxycycline monohydrate DYNACIN (minocycline) MINOCIN (minocycline) minocycline SR capsules minocycline tablets MONODOX (doxycycline monohydrate) ORACEA (doxycycline monohydrate) SOLODYN (minocycline) SUMYCIN (tetracycline) VIBRAMYCIN SYRUP (doxycycline calcium) VIBRAMYCIN (doxycycline hyclate) VIBRAMYCIN (doxycycline monohydrate) VIBRAMYCIN (doxycycline monohydrate) VIBRAMYCIN (doxycycline monohydrate) VIBRA-TABS (doxycycline hyclate)	A ten-day trial of each of the preferred agents is required before a non-preferred agent will be approved. *Demeclocycline will be approved for conditions caused by susceptible strains of organisms designated in the product information supplied by the manufacturer. A C&S report must accompany this request. *Demeclocycline will also be approved for SIADH.



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
ULCERATIVE COL	ITIS AGENTS ^{AP}		
		RAL	
	APRISO (mesalamine) ASACOL (mesalamine) 400mg COLAZAL (balsalazide) DIPENTUM (olsalazine) PENTASA (mesalamine) 250mg sulfasalazine	ASACOL HD (mesalamine) 800mg AZULFIDINE (sulfasalazine) balsalazide LIALDA (mesalamine) PENTASA (mesalamine) 500mg	Thirty (30) day trials of each of the preferred agents of a dosage form must be tried before a non-preferred agent of that dosage form will be authorized unless one of the exceptions on the PA form is present.
		CTAL	
	CANASA (mesalamine) mesalamine	SF ROWASA (mesalamine)	
VAGINAL ANTIBAC	CTERIALS		
	clindamycin cream METROGEL (metronidazole)	AVC (sulfanilamide) CLEOCIN CREAM (clindamycin) CLEOCIN OVULE (clindamycin) CLINDESSE (clindamycin) metronidazole VANDAZOLE (metronidazole)	A trial, the duration of the manufacturer's recommendation, of each of the preferred agents is required before a non-preferred agent will be approved unless one of the exceptions on the PA form is present.
MISC BRAND/GEN	ERIC		
	CLON	NIDINE	
	CATAPRES-TTS (clonidine) clonidine tablets	clonidine patch NEXICLON XR (clonidine) CATAPRES TABLETS (clonidine)	A thirty (30) day trial of each preferred unique chemical entity in the corresponding therapeutic category is required before a non-preferred agent will be authorized.
MEGESTROL			
	MEGACE ES (megestrol) megestrol	MEGACE (megestrol)	
SUBLINGUAL NITROGLYCERIN			
	nitroglycerin sublingual NITROSTAT SUBLINGUAL (nitroglycerin)	NITROLINGUAL (nitroglycerin) NITROMIST (nitroglycerin)	



This is not an all-inclusive list of available covered drugs and includes only managed categories. Refer to cover page for complete list of rules governing this PDL.

THERAPEUTIC DRUG CLASS	PREFERRED AGENTS	NON-PREFERRED AGENTS	PA CRITERIA
	OCTRE	EOTIDE	
	SANDOSTATIN (octreotide)	octreotide	
	EPINE	PHRINE	
	TWINJECT (epinephrine) EPIPEN (epinephrine)		
	ORAL CONT	RACEPTIVES	
	YASMIN (ethinyl estradiol/drospirenone)	BEYAZ (ethinyl estradiol/drospirenone/levomefolate) Gianvi (ethinyl estradiol/drospirenone) Ocella (ethinyl estradiol/drospirenone) YAZ (ethinyl estradiol/drospirenone)	
SUBSTANCE ABUSE TREATMENTS			
	SUBOXONE (buprenorphine) FILM ^{CL}	SUBOXONE (buprenorphine) TABS ^{CL}	Suboxone PA criteria is available at http://www.dhhr.wv.gov/bms/Pharmacy/Pages/pac.aspx