

Drug Formulary Update, January 2014

Updates to the HealthPartners Drug Formularies are listed below.

Updates for the Commercial Drug Formularies and the Minnesota Health Care Programs (Medicaid and Minnesota Care "State Programs") Drug Formulary are listed first, and changes for the Medicare Drug Formulary are listed in the following section.

Commercial and Minnesota Health Care Programs

The following updates are effective January 1, 2014 unless otherwise noted, and apply to PreferredRx, GenericsPlusRx, and HealthPartners Minnesota Health Care Programs (Medicaid and Minnesota Care "State Programs") Drug Formularies.

Medication	Status	Notes
Afatinib (Gilotrif)	PA	Afatinib will be approved (1) for the first-line treatment of patients with metastatic non-small cell lung cancer (NSCLC) whose tumors have an epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 (L858R) mutations and (2) no other EGFR mutations are present and (3) a requested maximum daily dose of up to 40 mg. Initial coverage is authorized for 3 months, with 3-month extensions for patients with no disease progression. Gilotrif is considered a specialty medication by HealthPartners.
Aztreonam inhaled (Cayston)	PA	Aztreonam inhalation will be approved (1) for use in cystic fibrosis patients known to have <i>Pseudomonas aeruginosa</i> in the lungs and (2) in a quantity up to one 28-day package per month. Cayston is considered a specialty medication by HealthPartners.
Breo (fluticasone/ vilanterol)	NF	Breo is a combination inhaler, for COPD. Alternatives are available, including Advair and Symbicort.
Bromfenac (Prolensa) ophthalmic		Prolensa has been added as a line-extension to Bromday. Prolensa is on formulary for PreferredRx, and is non-formulary for GenericsPlusRx and State Programs.
Cough syrups	NF	These cough syrups have been deleted from formulary, effective April 1 2014. Hydrocodone/ chlorpheniramine Guaifenesin/ pseudoephedrine/ codeine (Cheratussin DAC) Phenylephrine/ codeine/ promethazine (Promethazine VC - codeine) Hydrocodone/ homatropine (Hydromet) Pseudoephedrine/ codeine/ chlorpheniramine There is little clinical need for these cough syrups, and they are more costly than preferred alternatives: guaifenesin/ codeine, codeine/ promethazine, and dextromethorphan/ promethazine.

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Medication	Status		No	tes	
Diclofenac gel (Voltaren gel)	F	Voltaren gel, for osteoarthritis, is on formulary. Step-therapy criteria have been removed.			
Dolutegravir (Tivicay)	F	Dolutegravir is a ne	w HIV medication.		
Dornase alfa inhaled (Pulmozyme)	PA	Dornase alfa inhala and (2) in a quantity Pulmozyme is consi	of up to one 30-un	it carton per month	
Duloxetine (Cymbalta)	F	Duloxetine, for dep therapy criteria hav	·	athic pain, is on forn	nulary. Step-
Mechlorethamine (Valchlor) gel	PA	patients who hav up to two 60 gm to Initial coverage is a	ycosis fungoides-type received prior skirtubes per month. uthorized for 3 mondisease progression.	pe cutaneous T-cell n-directed therapy a ths, with 3-month e	lymphoma in nd (2) at a quantity xtensions for
Nepafenac (Ilevro) ophthalmic		Ilevro has been add Ilevro is on formula GenericsPlusRx and	ry for PreferredRx, a		/ for
Oxycodone ER (OxyContin)	QL	Quantity Limits for OxyContin have been updated. Quantity limits now allow a maximum of 120mg morphine equivalents p (MED), similar to other opioid medications.			
		Strength	New Daily Quantity Limit	New Daily mg limit	MED
		10 mg	4 (no change)	40	60
		15 mg	4 (no change)	60	90
		20 mg	4 (no change)	80	120
		30 mg	2	60	90
		40 mg	2	80	120
		60 mg	0	0	0
		80 mg	0	0	0
		These limits will be sent to affected pro higher doses are no Exceptions allowing Requests should income the risk of addiction	viders and member t affected by this ch higher quantities ca clude a treatment pl	s. Patients previous ange. an be requested if m an, and must includ	sly authorized at nedically necessary.
Ponatinib (Iclusig)	NF	Ponatinib has been	removed from form	ulary, due to marke	t withdrawal.

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Medication	Status	Notes
Simbrinza (brinzolamide/ brimonidine) ophthalmic		Simbrinza has been added as a line-extension. Both brinzolamide and brimonidine are on formulary, and costs are similar. Simbrinza is on formulary for PreferredRx, and is non-formulary for GenericsPlusRx and State Programs.
Tobramycin inhaled	PA	Tobramycin inhalation will be approved (1) for use in cystic fibrosis patients known to have <i>Pseudomonas aeruginosa</i> in the lungs and (2) in a quantity up to one 56-count carton per month. Generic Tobi is preferred. Other forms are considered non-formulary with prior authorization: Tobi Brand, Tobi Podhaler, and Bethkis. Inhaled tobramycin is considered a specialty medication by HealthPartners.
Tricyclic antidepressants amitriptyline doxepin imipramine trimipramine clomipramine	Age	 These tricyclic antidepressants will be limited for the elderly (ages 65 and older). amitriptyline (Elavil), chlordiazepoxide/ amitriptyline (Limbitrol), and perphenazine/ amitriptyline (Etrafon) doxepin (Silenor) imipramine (Tofranil) trimipramine (Surmontil) clomipramine (Anafranil). Clomipramine will be approved for obsessive-compulsive disorder (no additional coverage criteria). These tricyclic antidepressants are anticholinergic with a high rate of side effects in the elderly. Nortriptyline (Pamelor) and desipramine (Norpramin) remain on formulary for all age groups. These limits will be added on April 1. Additional communications are being sent to affected providers and members. Exceptions for these tricyclic antidepressants can be requested if medically necessary.
V-Go	PA	V-Go is a disposable insulin pump, reserved for prescribing by an Endocrinologist to ensure appropriate training.
Vortioxetine (Brintellix)	NF	Vortioxetine is a new antidepressant. Significant advantages aren't clear.

Medicare Drug Formulary

These changes are effective January 1, 2014.

Medication	Status	Notes
Afatinib (Gilotrif)	T5 PA	Covered Use: ALL FDA-APPROVED INDICATIONS, NOT OTHERWISE EXCLUDED FROM PART D.
		Required Medical Information: FOR NEW START PATIENTS: INITIAL CRITERIA - (1) DIAGNOSIS OF METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC) WITH TUMORS THAT HAVE EPIDERMAL GROWTH FACTOR RECEPTOR (EGFR) EXON 19 DELETIONS OR EXON 21 (L858R) SUBSTITUTION MUTATIONS AS DETECTED BY AN FDA-APPROVED TEST OR AT A CLIA-APPROVED FACILITY AND (2) NO OTHER EGFR MUTATIONS ARE PRESENT. RENEWAL CRITERIA - DOCUMENTATION EVERY 3 MONTHS THAT THERE HAS BEEN NO DISEASE PROGRESSION.
		Coverage Duration: THREE MONTHS, WITH APPROVAL EVERY THREE MONTHS IF RENEWAL CRITERIA ARE MET.
		Gilotrif was added August 30 2013, to meet Medicare new drug guidelines.
Degarelix (Firmagon)	T4	Prior authorization limits have been removed.
Diclofenac gel (Voltaren gel)	Т3	Step-therapy criteria have been removed.
Dolutegravir (Tivicay)	Т3	Dolutegravir is a new HIV medication. Tivicay was added August 30 2013, to meet Medicare new drug guidelines.
Duloxetine (Cymbalta)	Т3	Step-therapy criteria have been removed.
Mechlorethamine (Valchlor) gel	T5 PA	Covered Use: ALL FDA-APPROVED INDICATIONS, NOT OTHERWISE EXCLUDED FROM PART D.
		Coverage Duration: THREE MONTHS, WITH APPROVAL EVERY THREE MONTHS IF RENEWAL CRITERIA ARE MET.
		Valchlor was added October 25 2013, to meet Medicare new drug guidelines.
Pertuzumab (Perjeta)	T5 PA	Updated PA to include new neoadjuvant labeled indication.
		Covered Use: ALL FDA-APPROVED INDICATIONS, NOT OTHERWISE EXCLUDED FROM PART D.
		Required Medical Information: FOR NEW START PATIENTS: INITIAL CRITERIA - DIAGNOSIS OF AN FDA-APPROVED INDICATION, NOT OTHERWISE EXCLUDED FROM PART D. RENEWAL CRITERIA - DOCUMENTATION THAT DISEASE PROGRESSION HAS NOT OCCURRED.
		Coverage Duration: SIX MONTHS, WITH APPROVAL EVERY SIX MONTHS IF RENEWAL CRITERIA ARE MET.
Ponatinib (Iclusig)		Removed from formulary, due to market withdrawal.

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Medication	Status	Notes
Vincristine liposomal (Marqibo)	T5 PA	Covered Use: ALL FDA-APPROVED INDICATIONS, NOT OTHERWISE EXCLUDED FROM PART D. Required Medical Information: FOR NEW START PATIENTS: (1) DIAGNOSIS OF AN FDA-APPROVED INDICATION, NOT OTHERWISE EXCLUDED FROM PART D, OR (2) DOCUMENTATION THAT A PATIENT IS CURRENTLY RECEIVING OR HAS PREVIOUSLY RECEIVED AND BENEFITED FROM THE USE OF THIS MEDICATION FOR THE TREATMENT OF CANCER. Coverage Duration: CONTRACT YEAR, BALANCE OF CONTRACT YEAR, OR FDA-APPROVED DURATION, WHICHEVER IS APPLICABLE FOR USE. Marqibo was added September 2 2013, to meet Medicare new drug
Vortioxetine (Brintellix)	T3 PA	guidelines. FOR NEW START PATIENTS: (1) DIAGNOSIS OF AN FDA-APPROVED INDICATION, NOT OTHERWISE EXCLUDED FROM PART D, AND (2) DOCUMENTATION OF AN INADEQUATE RESPONSE OR MEDICAL CONTRAINDICATION TO TWO PREFERRED ALTERNATIVE ANTIDEPRESSANTS: CITALOPRAM, ESCITALOPRAM, FLUOXETINE, PAROXETINE, SERTRALINE, VENLAFAXINE, OR CYMBALTA. Brintellix was added November 30 2013, to meet Medicare new drug guidelines.

For Medicare: T1, covered generic T5, specialty

T2, covered generic T3

T3, covered Brand

T4, covered Brand

Formulary Information and Requests

Formulary Information is available at HealthPartners.com/ Provider/ <u>Pharmacy Services</u>, including the <u>Drug</u> Formularies.

Pharmacy Customer Service is available to providers (physicians and pharmacies) by fax, phone, and mail.

- Fax submission of coverage requests is preferred: 952-853-8700 or 1-888-883-5434.
- Telephone service is available: 952-883-5813 or 1-800-492-7259. HealthPartners Pharmacy Customer Service is available from 8AM 6PM CST, Monday through Friday. After hours calls are answered by our Pharmacy Benefit Manager.
- Mail: HealthPartners Pharmacy Services, 8170 33rd Avenue South, PO Box 1309, Mpls, MN 55440.