

Formulary changes announced

First quarter changes to the Independent Health Drug formularies are summarized below and are currently in effect unless otherwise noted.

NPG/NPB - Non-Preferred Generic/Brand PG/PB - Preferred Generic/Brand

NF – Non-Formulary SP - Specialty Pharmacy PA - Prior Authorization

ST – Step Therapy

QL – quantity Limits SC – subcutaneous IM - intramuscular inj – injection

tab – tablet cap – capsule soln – solution

susp – suspension MDI-metered-dose inhalation aerosol

ODT – orally-disintegrating tablet

LDD – Specialty Pharmacy Limited Distribution LA - available only at certain pharmacies

G - Generic (T2) for Medicare

Changes to Drug Formulary I (DFI), FEHB Formulary, and Pharmacy Benefit Dimensions Formulary				
Medications Added to Formulary	Medications Non-Formulary 4/1/2023	Changes to Formulary 4/1/2023		
4/1/2023				
Airsupra (budesonide/albuterol) MDI- NPB, ST, AL, QL Cibinqo added PB PA (1/1/2023) Jaypirca (pirtobrutinib) tab- NPB, PA, SP, LA Jesduvroq (daprodustat) tab- TBD NPB, PA if Rx benefit Jylamvo (methotrexate) oral soln- NPB Krazati (adagrasib) tab- NPB, PA, SP, LA Orserdu (elacestrant) tab- NPB, PA, SP, LA Rezlidhia ((olutasidenib) cap- NPB, PA, SP Sunlenca (lenacapavir) tab- PB	Atorvaliq (atorvastatin) oral susp Brenzavvy (bexagliflozin) tab Idacio (adalimumab-aacf) SC inj Iyuzeh (latanoprost) ophthalmic soln Olpruva (sodium phenylbutyrate) oral susp	Bonjesta T2 to NF (7/1/2023) Dificid PA changed to PA except ID and oncology (12/9/2022) Kristalose T3 to NF (7/1/2023) mesalamine 4g enema & cleanser wipe kit T2 to NF (7/1/2023) Millipred T3 to NF (7/1/2023) Noxafil (packets, suspension) PA changed to PA except ID and oncology (11/14/2022) PEG 3350 packets add to T1 ACA (11/21/2022) Saxenda T3PA to NF (7/1/2023) [FEHB 1/1/2024] testosterone topical gels/soln PA changed to PA except endocrinology and urology (1/1/23)		

Changes to Drug Formulary II (DFII), Drug Formulary III (DFIII), and Essential Plan Formulary (EPF)				
Medications Added to Formulary 4/1/2023	Medications Non-Formulary 4/1/2023	Changes to Formulary 4/1/2023		
Airsupra (budesonide/albuterol) MDI- NPB, ST, AL, QL Cibinqo added PB PA (1/1/2023) Jaypirca (pirtobrutinib) tab- NPB, PA, SP, LA Jesduvroq (daprodustat) tab- TBD NPB, PA if Rx benefit Jylamvo (methotrexate) oral soln- NPB Krazati (adagrasib) tab- NPB, PA, SP, LA Orserdu (elacestrant) tab- NPB, PA, SP, LA Rezlidhia ((olutasidenib) capsules- NPB, PA, SP Sunlenca (lenacapavir) tab- PB	Atorvaliq (atorvastatin) oral susp Brenzavvy (bexagliflozin) tab Idacio (adalimumab-aacf) SC inj Iyuzeh (latanoprost) ophthalmic soln Olpruva (sodium phenylbutyrate) oral susp	Bonjesta T2 to NF (7/1/2023) budesonide 3mg DR cap T2 to T1 Kristalose T3 to NF (7/1/2023) mesalamine 4g enema & cleanser wipe kit T2 to NF (7/1/2023) Millipred T3 to NF (7/1/2023) Noxafil (packets, suspension) PA changed to PA except ID and oncology (11/14/2022) PEG 3350 packets add to T1 ACA (11/21/2022) Saxenda T3PA to NF (7/1/2023) sucralfate oral suspension added NPG (12/29/2022) testosterone topical gels/soln PA changed to PA except endocrinology and urology (1/1/23)		

Changes to Medicaid Formulary				
Medications Added to Formulary 4/1/2023	Medications Non-Formulary 4/1/2023	Changes to Formulary 4/1/2023		
Airsupra (budesonide/albuterol) MDI- PB, ST, AL, QL	Atorvaliq (atorvastatin) oral susp	Bonjesta T2 to NF (7/1/2023)		
Sunlenca (lenacapavir) tab- PB	Brenzavvy (bexagliflozin) tab	Noxafil (packets, suspension) PA changed to PA except		
Tamiflu PB (12/9/2022)	Idacio (adalimumab-aacf) SC inj	ID and oncology (11/14/2022)		
	Iyuzeh (latanoprost) ophthalmic soln	testosterone topical gels/soln PA changed to PA except		
	Jaypirca (pirtobrutinib) tab	endocrinology and urology (1/1/23)		
	Jesduvroq (daprodustat) tab-TBD NF if Rx benefit			
	Jylamvo (methotrexate) oral soln- NPB			
	Krazati (adagrasib) tab			
	Olpruva (sodium phenylbutyrate) oral susp			
	Orserdu (elacestrant) tab			
	Rezlidhia ((olutasidenib) cap			

Changes to Medicare Formulary			
Medications Added to Formulary	Medications Non-Formulary 4/1/2023	Changes to Formulary 4/1/2023	
4/1/2023			
Jaypirca (pirtobrutinib) tab- NPB, PA, SP, LA Jesduvroq (daprodustat) tab- TBD NPB, PA if Rx benefit Jylamvo (methotrexate) oral soln- NPB Krazati (adagrasib) tab- NPB, PA, SP, LA Orserdu (elacestrant) tab- NPB, PA, SP, LA Rezlidhia ((olutasidenib) capsules- NPB, PA, SP	Atorvaliq (atorvastatin) oral susp Brenzavvy (bexagliflozin) tab Idacio (adalimumab-aacf) SC inj Iyuzeh (latanoprost) ophthalmic soln Olpruva (sodium phenylbutyrate) oral susp	amlodipine/hydrochlorothiazide/valsartan added to T2 (3/1/2023) dipyridamole lowered from T4 to T2 effective 1/1/23	

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COVID-19 Emergency Use Authorizations (EUA):

These EUAs are covered as medical drugs under specially created administration codes for Medicaid and Commercial members. Providers buy-and-bill Independent Health directly. These EUAs are covered under Original Medicare (Part A or Part B) for those members.

- On 11/30/22, the FDA withdrew the EUA for **bebtelovimab** in all areas because of evidence the drug would be ineffective against major circulating variants of COVID-19. Commercial distribution has been paused but could restart if future variants become susceptible once again.
- On 12/8/22, the EUAs for both the **Pfizer-BioNTech COVID-19 vaccine** and the **Moderna COVID-19 vaccine** were amended to authorize a single Omicron BA.4- and BA.5-specific booster dose after the completion of any primary COVID-19 vaccine series in very young children. For the Pfizer-BioNTech vaccine, the authorization is for individuals 6 months of age to 4 years of age. For the Moderna vaccine, the authorization is for individuals 6 months of age to 5 years of age. The CDC recommended these treatments on 12/9/22.
- On 12/21/22, the EUA for **Actemra** (tocilizumab) for treatment of hospitalized adult patients with COVID-19 who are receiving systemic corticosteroids and require supplemental oxygen, non-invasive or invasive mechanical ventilation, or extracorporeal membrane oxygenation (ECMO) was changed to a full approval. The EUA was previously in place for patients 2 years of age and older. Actemra remains authorized for patients who fit these criteria 2 to 18 years of age.
- On 1/26/23, the FDA revised the EUA for Evusheld (tixagevimab/cilgavimab) to limit its use to when the combined frequency of non-susceptible SARS-CoV-2 variants nationally is ≤90%. Based on this revision, it is not currently authorized for use anywhere in the US until further FDA notice.
- On 2/1/2023, the EUAs for both Lagevrio (molnupiravir) and Plaxlovid (nirmatrelvir/ritonavir) were changed to remove the requirement for positive test results to prescribe these products. The FDA continues to recommend that providers use direct SARS-CoV-2 viral testing to help diagnose COVID-19.

Informational:

- On 11/22/22, the U.S. Food and Drug Administration released a Drug Safety Communication regarding **Prolia** (denosumab). The FDA announced it was investigating a risk of severe hypocalcemia in patients with advanced kidney disease on dialysis. While patients using Prolia and on dialysis have not been directed to stop this medication at this time, they should be alerted of possible signs and symptoms of hypocalcemia and healthcare professionals should monitor these patients more closely.
- On 11/29/22, the sponsor of **Tecentriq** (atezolizumab) announced it was voluntarily removing the accelerated indication for locally advanced/metastatic urothelial cell cancer after confirmatory trial results failed to show a significant benefit. Its indications for non-small cell lung cancer, small cell lung cancer, hepatocellular carcinoma, and melanoma remain.
- On 1/12/23, the US Drug Enforcement Administration (DEA) and Substance Abuse and Mental Health Services Administration announced the elimination of the X-waiver requirement for **buprenorphine** prescribing, effective immediately. All future prescriptions for buprenorphine only require a DEA number, not a special number starting with X, and the limits on the number of patients who may be treated under a particular X-number have been removed. These requirements were considered a barrier to more widespread access of buprenorphine for treating opioid use disorder.
- On 2/9/2023, the accelerated approval of Jemperli (dostarlimab) for endometrial cancer that has progressed on or following prior
 treatment with a platinum-containing regimen in any setting and are not candidates for curative surgery or radiation was changed to a
 full approval.

Products removed from market:

On 11/22/22, the sponsor of Blenrep (belantamab mafodotin-blmf) announced it was beginning the process of removing the drug
from the market after confirmatory trials failed to meet requirements to convert the accelerated approval to a full approval. It was
given an accelerated approval in August 2020 as monotherapy for the treatment of relapsed/refractory multiple myeloma after receipt
of at least four prior therapies. The sponsor will be continuing clinical trials as it believes the drug will eventually show clinical benefit
as survival data matures further.

Line extensions:

Adds:

- Austedo (deutetrabenazine) new titration pack added to each LOB following current coverage
- Oxbryta (voxelotor) new 300 mg strength added to each LOB following current coverage
- Ozempic (semaglutide) new concentration added to each LOB following current coverage
- Skyrizi (risankizumab) new strength added to each LOB following current coverage
- Turalio (pexidartinib) new 125 mg capsules added to each LOB following current coverage

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Drugs with new indications:

Brand name	Generic name	New indication(s)	Coverage changes
Actemra	tocilizumab	COVID-19 hospitalized adults	n/a
Adacel	tetanus, diphtheria, pertussis	pertussis in infants (given during pregnancy)	n/a
Brukinsa	zanubrutinib	chronic or small lymphocytic lymphoma	Policy updates
Cibinqo	abrocitinib	indication expanded down to 12 years of age	AL changes
Cytalux	pafolacianine	identification of lung cancer lesions	n/a
Enjaymo	sutimlimab-jome	transfusion-independent cold agglutinin disease	Policy changes
Eylea	aflibercept	retinopathy of prematurity	Policy updates
Ibrance	palbociclib	indication expanded to include all female patients	Policy changes
Keytruda	pembrolizumab	single-agent NSCLC indication	Policy changes
Pemfexy	pemetrexed	NSCLC in combo with platinum/pembrolizumab	Policy updates
Precedex	dexmedetomidine	procedural sedation expanded down to 1 month of age	n/a
Revatio	sildenafil	age range expanded down to 1 years of age and older	AL changes
Takhzyro	landelumab-flyo	age range expanded down to 2 years of age and older	AL changes
Tascenso	fingolimod ODT	age range expanded to adults	AL changes
Tecentriq	atezolizumab	unresectable or metastatic alveolar soft part sarcoma	Policy updates
Tukysa	tucatinib	HER-2(+) colorectal cancer w/trastuzumab	Policy updates
Tymlos	abaloparatide	indication expanded to men	Policy changes
Udenyca	pegfilgrastim-cbqv	Hematopoietic Subsyndrome of Acute Radiation Syndrome	Policy updates
Vraylar	cariprazine	adjunct for major depressive disorder in adults	Policy updates
Xeloda	capecitabine	various, including pancreatic cancer	Policy updates

Medical:

- Adstiladrin (nadofaragene firadenovec-vncg) intravesical suspension: medical PA
- Briumvi (ublituximab-xiiy) IV inj- medical PA SP
- Elahere (mirvetuximab soravtansine-gynx) IV inj- medical PA
- Hemgenix (etranacogene dezaparvovec-drlb) IV susp- medical PA SP LA
- Jesduvroq (daprodustat) tab- TBD: medical bundled with dialysis
- Leqembi (lecanemab-irmb) IV inj- Block for inconclusive clinical benefit; FEHB: medical PA
- Lunsumio (mosunetuzumab-axgb) IV inj- medical PA
- NexoBrid (anacaulase-bcdb) topical gel- medical
- Rebyota (fecal microbiota, live-jslm) rectal suspension- medical PA
- Rykindo (risperidone) IM inj- medical PA
- Sezaby (phenobarbital) IV inj- medical
- Sunlenca (lenacapavir) SC inj- medical
- Tzield (teplizumab-mzvw) IV inj- medical PA SP LA
- Vibrant cap as medical device- block as experimental intervention
- Xenoview (xenon Xe-129 hyperpolarized) inhalant contrast agent- medical

New generics:

Brands now non-formulary unless otherwise indicated. For Medicaid, generics are generally left as non-formulary if their respective brands were non-formulary.

		Generic tier placement/utilization management			
Brand name	Generic name	Commercial/ FEHB	Exch/Small/EBP	Medicaid	Medicare Indiv/PDP
Adrenalin	epinephrine	Medical	Medical	Medical	Medical
Cambia	diclofenac powder packs	NF	NF	NF	NF/NF
Denavir	penciclovir	NF	NF	NF	NF/NF
Esbriet caps	pirfenidone	T3 PA	T3 PA	NF	NF/NF
Hetlioz	tasimelteon	T3 PA	T3 PA	NF	T5/T1
Keveyis	dichlorphenamide	T3 PA	T3 PA	NF	T5 PA/T1 PA
Latuda	lurasidone	T1	T1	T1	T5/T1
Mirvaso	brimonidine topical	T3 PA	T3 PA	NF	T4/T1
Trokendi XR	topiramate	T3 PA	T3 PA	NF	NF/NF
Zioptan	tafluprost	NF	NF	NF	NF/NF

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