Formulary Updates

DEFINITIONS Formulary These drugs are included in Mass General Brigham's covered drug list. **Non-Formulary** These drugs are not included in Mass General Brigham's formulary. The plan would only cover formulary alternatives. Providers can request Non-Formulary drugs as an exception, and the plan would require trial of all appropriate formulary alternatives prior to approving coverage of a Non-Formulary drug. If a Non-Formulary drug is approved, the member's cost sharing would be the highest tier. Preferred These drugs are on Mass General Brigham's formulary and offer a lower cost to members. **Non-Preferred** These drugs are on Mass General Brigham's formulary but offer a higher cost to members. Excluded Mass General Brigham does not cover these drugs. Members will receive a denial for all Excluded drug requests.

Updates for MassHealth Members

Effective 03/04/2024

The following generic medications will become non-preferred. Please use the brand name alternative(s):

Generic Medication	Brand Name Alternative
risperidone 12.5 mg, 25 mg, 37.5 mg, 50 mg	Risperdal Consta
extended-release intramuscular injection	

The following brand name medications will become non-preferred. Approval will require a trial of its generic medication:

Brand Name	Generic Medication
Lotronex tablet	Alosetron tablet

Effective 03/05/2024 – Reminders

	As of Dec 2023, GSK discontinued branded Flovent HFA (all strengths) and Flovent Diskus (all strengths).
	Generic Flovent (<i>fluticasone proprionate inhalation <u>aerosol</u></i>) will require a prior authorization if member is ≥ 5 years of age.
Respiratory Agents – Inhaled	Generic Flovent (<i>fluticasone proprionate inhalation <u>powder</u></i>) will require a prior authorization.
	Current members that have been grandfathered through 03/04/2024 will now be subject to a prior authorization as of 03/05/2024 .

Effective 03/04/2024

The following changes are being made to the listed medications to be in compliance with the MassHealth UPPL (Unified Pharmacy Product List):

Amyotrophic Lateral Sclerosis Agents	Qalsody 100mg/15ml solution will be added to the medical benefit with prior authorization.
Anesthetics – Topical	Iheezo 3% gel (<i>chloroprocaine ophthalmic</i>) has been added to the pharmacy benefit with prior authorization.
Anti-Allergy & Anti- Inflammatory Agents	Miebo (<i>perfluorohexyloctane</i>) has been added to the pharmacy benefit with prior authorization and quantity limit of 3 mL per 30 days.
Antibiotics - Injectables	 Xacduro (sulbactam/durlobactam injection) will be added to the pharmacy benefit with prior authorization and will be available without prior authorization on the medical benefit. Note: This medication will not be part of the federal rebate program. Additional restrictions may apply.
Anticonvulsants	Sezaby (<i>phenobarbital 100mg vial</i>) will be available <u>without</u> prior authorization on the medical benefit.



Antidepressants	 Zurzuvae (<i>zuranolone</i>) has been added to the pharmacy benefit with prior authorization and the following quantity limits: 30mg – 14 capsules per 45 days 20mg and 25mg – 28 capsules per 45 days Ketalar (<i>ketamine injection</i>) will only be available on the medical benefit with prior authorization:
Antidiabetic Agents: Non-Insulin and Combo Products	New drug, Inpefa (sotagliflozin), was added to the pharmacy benefit with prior authorization and quantity limit of 30 tablets per 30 days.
Anti-diarrhea Agents	Opium tincture will <u>remain</u> on the pharmacy benefit with prior authorization and additional quantity limit of 72 mL per 30 days.
Antifungals – Oral and Injectable	 The following updates has been made to Rezzayo (<i>rezafungin injection</i>): added to the pharmacy benefit with prior authorization and quantity limit of ≤ 6 vials for one course of therapy added to the medical benefit with prior authorization
Antimalarials	The criteria for off-label diagnoses for Daraprim (pyrimethamine) were updated to ensure it is being used in combination with an additional anti- infective agent.
Anti-Obesity Agents	The criteria was updated to reflect that overweight is an acceptable diagnosis.
Antipsychotics	Rykindo (risperidone ER intramuscular injection) will be added to the pharmacy benefit with prior authorization and quantity limit of 2 injections per 28 days.Note: The MassHealth Pediatric Behavioral Health Medication Initiative may apply to members <18 years of age due to polypharmacy, age, and/or drug restrictions.
Asthma and Allergy Monoclonal antibodies	The Xolair criteria was updated to include off-label diagnosis of systemic mastocytosis.



Benzodiazepines & other Antianxiety Agents	The MassHealth Concomitant Opioid Benzodiazepine Initiative was updated to require a prior authorization of any benzodiazepine and opioid agent used concomitantly greater than 15 days within the past 45-day period.
Opioids and Analgesics	
Beta thalassemia, MDS, and SCD Agents	Reblozyl criteria was updated to include diagnosis of myelodysplastic syndromes associated anemia.
Breast Cancer Agents	Enhertu (fam-trastuzumab deruxtecan-nxki injection) will no longer be available on the pharmacy benefit and will be restricted to medical benefit only with prior authorization. Trodelvy criteria was updated to be more in line with the FDA-labeled indications.
Cardiovascular Antihypertensives and Miscellaneous Cardiovascular Medications	 Filspari (<i>sparsentan tablet</i>) will be added to the pharmacy benefit with prior authorization and quantity limit of 30 tablets per 30 days. Prior authorization was <u>removed</u> from diltiazem CD 360 mg capsule. Entresto will have a quantity limit of 60 tablets per 30 days and <u>will remain</u> on the pharmacy benefit with a prior authorization.
Cerebral Stimulants & ADHD	 The following medications had quantity limits (QL) added and prior authorization will remain on the pharmacy benefit: Dyanavel XR (amphetamine extended-release 2.5mg/mL oral suspension) - QL ≤ 8 mL (20 mg) per day. Dyanavel XR (amphetamine extended-release chewable tablet) - QL 30 tablets per 30 days. Quillivant XR (methylphenidate extended-release oral suspension) - QL ≤ 12 mL (60 mg) per day. Relexxii (18mg, 27mg, 36mg, 54mg) - QL 60 tablets per 30 days Relexxii (45mg, 63mg and 72mg) - QL 30 tablets per 30 days Clonidine ER 0.1 mg will require a prior authorization through the pharmacy benefit if member is less than 3 years of age and the requested quantity exceeds 4 tablets per day.



CGRP Inhibitors	Zavzpret (<i>zavegepant nasal spray</i>) will be added to the pharmacy benefit with prior authorization and quantity limit of 12 units per 30 days.
Complement Inhibitors & Immunosuppresive Agents	 The following new drugs will be available on <u>only the medical benefit</u> with prior authorization: Rystiggo (rozanolixizumab-noli) Vyvgart Hytrulo (efgartigimod alfa and hyaluronidase-qvfc)
Corticosteroids – oral agents	Tarpeyo criteria was updated to ensure consistency with Filspari criteria for the same indication, immunoglobulin A nephropathy (IgAN).
Dermatological Agents (Topical Chemo-Genital Wart Therapy)	New drug, Ycanth (<i>cantharidin</i>), will be added to the medical benefit only with prior authorization.
Diabetic Testing Supplies	Freestyle Neo criteria was updated to include Freestyle Libre 3 as a compatible CGM device.
Glaucoma Agents	Iyuzeh (<i>latanoprost PF solution</i>) will be added to the pharmacy benefit with prior authorization.
Glycopyrrolate Agents	Dartisla ODT was added as a step-through option for Cuvposa for members ≥ 17 years of age.
Growth Hormone Agents	New drug, Ngenla (<i>somatrogon-ghla</i>), will be added to the pharmacy benefit with prior authorization. Skytrofa will be a preferred agent. Requests for Sogroya or Ngenla will require a step-through of Skytrofa, while requests for other non-preferred agents will continue to require a step-through of Genotropin.



Hepatitis Antiviral Agents FAQ and Guideline	Criteria was updated to remove the 12-week treatment recommendation for members if treatment-naïve with cirrhosis and HIV-coinfection.
Immunosuppressants	The following medications will be available on the medical benefit only without prior authorization:
	 Sandimmune IV (cyclosporine injection) Simulect (basiliximab)
	The criteria for Prograf granules and Sandimmune solution was updated to accept members < 13 years of age for medical necessity.
Kinase Inhibitors	Lytgobi (<i>futibatinib</i>) <u>will remain</u> on the pharmacy benefit with prior authorization and quantity limit of 5 tablets per day.
	Ayvakit criteria was updated to include the expanded indication of indolent systemic mastocytosis (ISM).
Melanoma Agents	Kimmtrak (<i>tebentafusp injection</i>) will be removed from the pharmacy benefit and will <u>remain</u> on the medical benefit only with prior authorization.
	The following medications will be added to the pharmacy benefit with prior authorization and quantity limit:
	 Tafinlar (<i>dabrafenib tablet for oral solution</i>) – QL ≤ 30 mL per day Mekinist (<i>trametinib solution</i>) – QL ≤ 40 mL per day
	The following medications were updated to include new indications, listed respectively:
	 Cotellic – histiocytic neoplasms Mekinist – low grade glioma Tafinlar – low grade glioma
Narcolonsy Agonts	Lumryz ER (<i>sodium oxybate extended-release suspension</i>) will be added to the pharmacy benefit with prior authorization.
Narcolepsy Agents	Note : This medication <u>will not</u> be part of the federal rebate program. Additional restrictions may apply.
Nonhormonal Agents for Menopausal Symptoms	Veozah (<i>fezolinetant tablet</i>) will be added to the pharmacy benefit with prior authorization and quantity limit of 30 tablets per 30 days.



Opioid Dependence and Reversal Agents	 Opvee (nalmefene nasal spray) will be added to the pharmacy benefit with prior authorization and quantity limit of 2 inhalers per year. Brixadi criteria was updated to allow more options to the treatment failure requirement.
PARP Inhibitors	 The following medications will be added to the pharmacy benefit with prior authorization: Zejula tablet (<i>niraparib 100mg, 200mg, 300mg</i>) with quantity limit of 30 tablets per 30 days Talzenna capsule (<i>talazoparib 0.1mg, 0.35mg</i>) The following medications were updated to include new indications, listed respectively: Lynparza – metastatic castration-resistant prostate cancer (mCRPC) Talzenna – BRCA-mutated locally advanced or metastatic breast cancer and metastatic castration-resistant prostate cancer (mCRPC)
Pharmaceutical Compounding	Criteria was updated to specify that members who are dependent on feeding tubes should not use agents that contain sweeteners or flavorings. Topical compounds requiring a prior authorization will now include transdermal route of administration.
Polivy (polatuzumab)	Criteria was updated to include expanded indication of previously untreated diffuse large B-cell lymphoma (DLBCL).
Prostate Cancer Agents	Akeega (niraparib/abiraterone tablets) will be added to the pharmacy benefit with prior authorization and quantity limit of 60 tablets per 30 days. New indication for Xtandi was added for metastatic castration-resistant prostate cancer (mCRPC).
Pulmonary Hypertension (PH) Agents	New drug, Liqrev (sildenafil oral suspension), will be added to the pharmacy benefit with prior authorization.
Respiratory Agents - Inhaled	New drug, Airsupra (<i>albuterol/budesonide</i>), will be added to the pharmacy benefit with prior authorization. Prior authorization was <u>removed</u> from ProAir Respiclick (<i>albuterol inhalation powder</i>) from the pharmacy benefit.
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	Criteria for Pulmicort Respules for eosinophilic esophagitis (EoE) was
	updated to remove the step-through fluticasone proprionate requirement.
	Off-label indication of EoE was added for fluticasone proprionate.
Skyrizi	Skyrizi IV will be added to the medical benefit with prior authorization required.
	Skyrizi SC will <u>remain</u> on the pharmacy benefit with prior authorization required.
	The following medications will be added to the pharmacy benefit with prior authorization:
	 New adalimumab biosimilar agents and associated unbranded generics
	• Litfulo (<i>ritlecitinib</i>) capsule – QL of 30 capsules per 30 days
	Subcutaneous Entyvio has been added to the same criteria as the IV formulation.
Targeted Immunomodulators	Entyvio criteria for Crohn's disease and ulcerative colitis was updated to require a step-through an anti-TNF agent.
	The step-through requirement of an anti-TNF agent has been removed from the following:
	 Taltz criteria for psoriatic arthritis, non-radiographic axial spondyloarthritis, and ankylosing spondylitis Stelara for psoriatic arthritis
	Cibinqo (abrocitinib) criteria was updated to be in line with the current FDA labeling for use in members 12 years of age and older.
	The following medications will be added to the medical benefit only with prior authorization:
T-cell immunotherapies	• Epkinly (epcoritamab-bysp) injection
	Columvi (glofitamab-gxbm) injection
Vitamins	ADEK Gummies (<i>multivitamins/zinc gummy</i>) will be added to the pharmacy benefit with prior authorization.
Vesicular monoamine transporter 2 (VMAT2) Inhibitors	The following medications will be added to the pharmacy benefit with prior authorization and quantity limit:
	 Austedo XR (deutetrabenazine extended-release) (6mg, 12mg) – QL 90 tablets per 30 days.



	 Austedo XR (deutetrabenazine extended-release) (24mg) – QL 60 tablets per 30 days. A new indication for Ingrezza was added for Huntington's disease.
Wound Care	Vyjuvek gel (<i>beremagene geperpavec-svdt</i>) will be added to the medical benefit only with prior authorization.

Upcoming Important Updates for MassHealth Members

	Novo Nordisk announced to the United States Food and Drug Administration (FDA) on November 8, 2023, the planned discontinuation of Levemir (<i>insulin detemir</i>) FlexPen and vials. Supply disruptions of the FlexPen are anticipated to begin mid-January 2024, followed by discontinuation of the FlexPen formulation on April 1, 2024, and discontinuation of the vial formulation on December 31, 2024.
	There are currently no commercially available biosimilars or authorized generics for insulin detemir.
Discontinuation of Levemir (insulin detemir)	Alternative long-acting insulin products covered under the MassHealth UPPL (Unified Pharmacy Product List) include the following:
	Agents without Prior Authorization (PA) • Lantus (<i>insulin glargine</i>) • Toujeo (<i>insulin glargine</i>) • Tresiba (<i>insulin degludec</i>)
	Agents Requiring PA • Basaglar (insulin glargine) • Semglee (insulin glargine-yfgn)
	Novo Nordisk announced to the FDA on November 18, 2023 , the planned discontinuation of GlucaGen HypoKit (glucagon) in the United States. This discontinuation will be effective July 1, 2024 .
Discontinuation of GlucaGen HypoKit (<i>glucagon</i>)	Generic formulations of glucagon are commercially available. Additionally, there are multiple branded alternative formulations. The following glucagon products are available through the MassHealth UPPL (Unified Pharmacy Product List).
	Agents without PA • Baqsimi (glucagon nasal powder) • glucagon vial



 Gvoke (glucagon auto-injection, prefilled syringe, vial)
 Zegalogue (dasiglucagon) – PA required until March 4, 2024

