

Request for Prior Authorization for Preferred Sodium-Glucose Co-Transporter 2 (SGLT2) Inhibitors GLP-1 Receptor Agonists, and Dipeptidyl Peptidase IV (DPP-IV) Inhibitors

Website Form – <u>www.highmarkhealthoptions.com</u> Submit request via: Fax - 1-855-476-4158

All requests for Sodium-Glucose Co-Transporter 2 (SGLT2) Inhibitors, GLP-1 Receptor Agonists, and Dipeptidyl Peptidase IV (DPP-IV) Inhibitors require a prior authorization and will be screened for medical necessity and appropriateness using the criteria listed below.

Prior Authorization Criteria:

For all requests for Preferred Sodium-Glucose Co-Transporter 2 (SGLT2) Inhibitors and the diagnosis is Type 2 diabetes all of the following criteria must be met:

• Claims will pay at the point of sale when a diagnosis of Type 2 diabetes is entered

For all requests for Preferred Sodium-Glucose Co-Transporter 2 (SGLT2) Inhibitors and the diagnosis is Heart Failure (HF) all of the following criteria must be met:

- Documentation the member's HF is NYHA class II-IV
- Member must be 18 years of age or older
- Must be prescribed by or in consultation with a cardiologist
- Member must have a reduced left ventricular ejection fraction of 40% or less
- Member has tried and failed (which will be verified via pharmacy claims if available) had an intolerance, contraindication, or is currently taking at least **two** of the following:
 - Beta blocker
 - o ACEi/ARB/ARNI
 - o Diuretic
 - o Aldosterone antagonist (if an aldosterone antagonist is indicated)
- The requested dose and frequency is in accordance with FDA-approved labeling, nationally recognized compendia, and/or evidence-based practice guidelines
- **Initial Duration of Approval:** 12 months

For all requests for Preferred GLP-1 Receptor Agonists all of the following criteria must be met:

• Claims will pay at the point of sale when a diagnosis of Type 2 diabetes is entered.

For all requests for Preferred Dipeptidyl Peptidase IV (DPP-IV) Inhibitors all of the following criteria must be met:

- Claims will pay at the point of sale when both of the following is met:
 - o A diagnosis for Type 2 diabetes is entered at the point of sale.



- O The member has filled metformin or a metformin containing product in the past 90 days.
- o For any request that does not meet automatic point of sale approval all of the following criteria must be met:
 - Documentation the member has a diagnosis of Type 2 diabetes
 - Documentation the member has tried and failed or has an intolerance or contraindication to any version of metformin or a metformin combination product.
 - Initial Duration of Approval: 12 months

• Reauthorization criteria

- Members with historical pharmacy claims data meeting the following criteria will receive automatic reauthorization at the pharmacy point of service without the requirement for documentation of additional information. If pharmacy claims data cannot obtain the criteria below, documentation will be required to indicate the member meets the reauthorization criteria below. Claims will automatically adjudicate on-line, without a requirement to submit for reauthorization when the following criteria is met:
 - Documentation the member has at least 1 fill of the requested medication in the past 45 days
- Reauthorization Duration of Approval: 12 months

Coverage may be provided for any non-FDA labeled indication if it is determined that the use is a medically accepted indication supported by nationally recognized pharmacy compendia or peer-reviewed medical literature for treatment of the diagnosis(es) for which it is prescribed. These requests will be reviewed on a case by case basis to determine medical necessity.

Drugs are authorized in generic form unless the branded product is on the preferred drug list or the prescriber has indicated in writing that the branded product is medically necessary. If only the branded product is on the preferred drug list, the generic form will be considered nonpreferred and shall not require the prescriber to indicate in writing that the branded product is medically necessary.



Sodium-Glucose co-Transporter 2 (SLGT2) Inhibitors GLP-1 Receptor Agonists, and Dipeptidyl Peptidase IV (DPP-IV) Inhibitors PRIOR AUTHORIZATION FORM

Please complete and fax all requested information below including any progress notes, laboratory test results, or chart documentation as applicable to Highmark Health Options Pharmacy Services. **FAX:** (855) 476-4158

If needed, you may call to speak to a Pharmacy Services Representative

If needed, you may call to speak to a Pharmacy Services Representative.				
PHONE: (844) 325-6251 Monday through Friday 8:00am to 7:00pm PROVIDER INFORMATION				
Requesting Physician:	NPI:			
Physician Specialty: Office Address:	Office Contact:			
Office Address:	Office Phone:			
Office Fax: MEMBER INFORMATION				
Member Name:	INFORMATION			
Health Options ID:	DOB:			
	NFORMATION			
Medication:	Strength:			
Directions:	Quantity: Refills:			
Is the member currently receiving requested	Date Medication Initiated:			
medication? Yes No	Date Wedication inflated.			
Is this medication being used for a chronic or long-term condition for which the medication may be necessary				
for the life of the patient? Yes No				
BILLING INFORMATION				
This medication will be billed: at a retail pharmacy OR				
medically (if medically please provide a JCODE:				
Place of service: Hospital Provider's office Member's home Other				
PLACE OF SERVICE INFORMATION				
Name:	NPI:			
Address:	Phone:			
MEDICAL HISTORY				
Member's Diagnosis: Type 2 Diabetes Heart Failure Other ICD-10:				
If the diagnosis is for Type 2 diabetes and the requested medication is a DPP-IV inhibitor: has the member tried and failed metformin or a metformin containing product?				
If the diagnosis is HF please answer the following two questions: 1. Does the member have heart failure with a NYHA class II-IV and a reduced ejection fraction ≤ 40%? ☐ Yes ☐ No				



2. Please mark which of the	e following the member	er has tried and faile	ed or had a contraindication to:	
Beta blocker	C			
ACEi/ARB/ARNI				
Diuretic				
	ist (if an aldosterone a	ntagonist is indicate	ed)	
	ist (if all algosterone a	inagomet is marcat		
CURRENT or PREVIOUS THERAPY				
Medication Name	Strength/	Dates of Therapy Status (Discontinued & W		
	Frequency		Status (Discontinued & Why/Current)	
	DE A LITHO	DIZATION CDI	TEDIA	
REAUTHORIZATION CRITERIA				
Has the member been on a SGLT2-inhibitor, GLP-1 Receptor Agonists, and Dipeptidyl Peptidase IV (DPP-IV)				
Inhibitors within the last 45 days? Yes No				
SUPPORTING INFORMATION or CLINICAL RATIONALE				
Prescribing Physician	Signatura		Date	
Trescribing Thysician	Signature		Date	