

Kaiser Permanente Health Plan of Mid-Atlantic States, Inc. Antihyperglycemic – Incretin Mimetics Combination (Mounjaro) Prior Authorization (PA)

Pharmacy Benefits Prior Authorization Help Desk Length of Authorizations: Initial- 4 months; Continuation- 12 months

## **Instructions:**

This form is used by Kaiser Permanente and/or participating providers for coverage of Antihyperglycemic – Incretin Mimetics Combination (Mounjaro) for Commercial and FEHB (Federal) plans. Please complete all sections, incomplete forms will delay processing. Fax this form back to Kaiser Permanente within 24 hours (fax: 1-866-331-2104). If you have any questions or concerns, please call 1-866-331-2103. Requests will not be considered unless all sections are complete.

KP-MAS Formulary can be found at: <a href="Pharmacy">Pharmacy</a> | Community Provider Portal | Kaiser Permanente

1 – Patient Information				
Patient Name:	Kaiser Medical ID#:	Date of Birth:		
2 – Prescriber Information				
Prescriber Name:	Specialty:	NPI:		
Prescriber Address:				
Prescriber Phone #:	Prescriber Fax #:	·		
Do you have an approved provider referral number from Kaiser Permanente?  □ Yes – please provide your provider referral number here:				
3 – Pharmacy Information				
Pharmacy Name:	Pharmacy NPI:			
Pharmacy Phone #	Pharmacy Fax #:			
4 – Drug Therapy Requested				
Drug 1: Name/Strength/Formulation: _				
Sig:				
Drug 2: Name/Strength/Formulation:				
) Sig:				

## 5- Diagnosis/Clinical Criteria

	<b>.</b>		
1.	Is this request for initial or continuing therapy?  □ Initial therapy  □ Continuing therapy, state start date:		
2.	Indicate the patient's diagnosis for the requested medication:		
Clinical Criteria:  1. Diagnosis of type 2 diabetes mellitus,  □ No □ Yes			
2.	2. <b>AND</b> on maximally tolerated metformin dose (2000 mg) for at least 3 months or allergy/intolerance to metformin documented,  □ No □ Yes		
3.	<ul><li>AND patient is not using for chronic weight management (CWM),</li><li>□ No □ Yes</li></ul>		
4.	<b>AND</b> patient is not on any agent in any of the following drug classes: GLP-1 agonists or DPP-4 inhibitors, $\Box$ No $\Box$ Yes		
5.	AND A1c within 2% of goal (as per ADA guidelines) in the past 3 months, and previously failed adequate trial (≥3 months), experienced intolerance or contraindication to ALL of the following agents or associated combination products:  o Glipizide o Pioglitazone (if BMI <35) o Tradjenta o Jardiance o At least TWO of the following KP-preferred GLP-1 agonists: Victoza, Ozempic, OR Rybelsus		
6.	OR A1c ≥2% above goal (as per ADA guidelines) in the past 3 months and recurrent hypoglycemia on insulin therapy despite dose adjustments, and previously failed adequate trial (≥3 months), experienced intolerance or contraindication to insulin glargine or insulin glargine-yfgn, Tradjenta, Jardiance, and at least TWO of the KP-preferred GLP-1 agonists listed above  □ No □ Yes		
For continuation of therapy, please respond to <u>additional questions</u> below:			
1.	Has the patient failed adequate trial (≥3 months), or has intolerance or contraindication to BOTH Victoza and Ozempic?  □ No □ Yes		
1.	Is the patient using for CWM?  □ No □ Yes		
2.	Does the patient have documented A1c lowering of at least 0.5% from initial or A1c now at goal? $\hfill\Box$ No $\hfill\Box$ Yes		

## 6 - Prescriber Sign-Off

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Additional Information –		
1. Please submit chart notes/medical records for the patient that are applicable to this request.		
. If member has not tried preferred agent(s) please provide rationale/explanation and any additional supporting		
information that should be taken into consideration for the req	uested medication:	
I certify that the information provided is accurate. Supporting documentation is available for State audits.		
Prescriber Signature:	Date:	
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