

1. Patient information





CONTAINS CONFIDENTIAL PATIENT INFORMATION

GLP-1 Agonist/Basal Insulin Combinations Prior Authorization of Benefits (PAB) Form

Complete form in its entirety and fax to:

Prior Authorization of Benefits Center at 1-844-512-9004 Provider Help Desk 1-800-454-3730

2. Physician information

Patient name:		Prescribing physician: _	Prescribing physician:					
Patient ID #: Patient DOB: Date of Rx: Patient phone #: Patient email address:		Physician address: Physician phone #: Physician fax #: Physician specialty: Physician DEA: Physician NPI #: Physician email address:						
				3. Medication	4. Strength	5. Directions	6. Quantity per 30 days	
						_	Specify:	
				7. Diagnosis:				
				8. Approval criteria: CHECK ALL BO) Note: Any areas not filled out are considered. Prior authorization is required for GL considered for patients when the following statements.	ed not applicable to you .P-1 agonist recepto	or/basal insulin combination		
				1) A diagnosis of Type 2 Diabetes Mellitus; and				
				2) Patient is 18 years of age or o	2) Patient is 18 years of age or older; and			
 The patient has not achieved maximally tolerated dose, unl contraindicated; and 	-							
 Documentation of an adequation agonist and one preferred longer 	•	•	ne preferred GLP-1 receptor					
5) Will not be used concurrently	5) Will not be used concurrently with prandial insulin; and							
6) Clinical rationale is provided a preferred long-acting insulin a		·	P-1 receptor agonist and a					
7) Medication will be discontinu dosage of: a) Soliqua below 15 units or b) Xultophy persistently belo	over 60 units, or	·	used if patients require a daily					
Most recent HgbA1C level:Date this level was obtained:								







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Prior Authorization of Benefits Center at 1-844-512-9004

Patient name:	Patient ID #:			
Metformin trial: Trial start date:	_Trial end date:	Trial dose:		
Reason for failure:				
Medical or contraindication reason to override	trial requirements:			
Preferred GLP-1 Receptor Agonist Trial: Drug r Trial start date: Trial end date:				
Preferred Long-Acting Insulin Trial: Drug name Trial start date:Trial end date:	e/dose:Reason for failur	re:		
Clinical rationale as to why patient cannot use a preferred GLP-1 receptor agonist and a preferred long-acting insulin agent concurrently:				
Is prandial insulin being used concurrently? Yes No				
Medication will be discontinued and alternation Soliqua – below 15 units or over 60 units		ill be used if patients require a daily dosage of:		
Xultophy - persistently below 16 units or	over 50 units Yes	□ No		
Attach lab results and other documentation a	s necessary.			
Patient name:	Patient	t ID #:		
9. Physician signature				
Prescriber or authorized signature Prior Authorization of Benefits is not the practice of medici	Date ine or the substitute for the indepe	endent medical judgment of a treating physician. Only a treating physician		
limitations, and exclusions. The submitting provider certific indicated and necessary to the health of the patient.	es that the information provided is	e plan for the detailed information regarding benefits, conditions, strue, accurate, and complete and the requested services are medically rization does not guarantee payment.		
The document(s) accompanying this transmission may cor	ntain confidential health information	ion that is legally privileged. This information is intended only for the use hibited from disclosing this information to any other party unless		
If you are not the intended recipient, you are hereby notif	, , , , ,	distribution, or action taken in reliance on the contents of these the sender immediately and arrange for the return or destruction of		