Xultophy (insulin degludec; liraglutide) Prior Authorization Request Form

Caterpillar Prescription Drug Benefit Phone: 877-228-7909 Fax: 800-424-7640

MEMBER'S LAST NAME:		MEMBER'S FIRST NAME:	
	g., chart notes or lab data, to su	ly and legibly. Attach any addit apport the authorization reques	
			URGENT
MEMBER INFORMATION			
LAST NAME:		FIRST NAME:	
PHONE NUMBER:		DATE OF BIRTH:	
STREET ADDRESS:			
CITY:		STATE: ZIP CODE:	
PATIENT INSURANCE ID NU	IMBER:		
IF YOU ARE NOT THE PATIENT OR THE PRESC FOLLOWING LINK: PRIMETHERAPEUTICS.COI	CRIBER, YOU WILL NEED TO SUBMIT A PHI DISCIMINOPP PRESENTATIVE (IF APPLICABLE)	HT (LB/KG): ALLERG LOSURE AUTHORIZATION FORM WITH THIS REQ	UEST WHICH CAN BE FOUND AT THE
PRESCRIBER INFORMATION			
LAST NAME:	N	FIRST NAME:	
PRESCRIBER SPECIALTY:		EMAIL ADDRESS:	
NPI NUMBER:		DEA NUMBER:	
PHONE NUMBER:		FAX NUMBER:	
STREET ADDRESS:			
CITY:		STATE: ZIP CODE:	
REQUESTOR (if different than prescriber):		OFFICE CONTACT PERSON:	
MEDICATION OR MEDICAL	DISPENSING INFORMATION		
MEDICATION NAME:			
DOSE/STRENGTH:	FREQUENCY:	LENGTH OF THERAPY/REFILLS:	QUANTITY:
■ NEW THERAPY	RENEWAL	IF RENEWAL: DATE THERAPY	'INITIATED:
DURATION OF THERAPY (SP	ECIFIC DATES):		



Continued on next page.

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MEMBER'S LAST NAME:	MEMBER'S FIRST	NAME:
1. HAS THE PATIENT TRIED ANY OTHER	R MEDICATIONS FOR THIS CONDITION?	YES (if yes, complete below) NO
MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE):	DURATION OF THERAPY (SPECIFY DATES):	RESPONSE/REASON FOR FAILURE/ALLERGY:
2. LIST DIAGNOSES:		ICD-10:
☐ Type 2 diabetes ☐ Other DiagnosisICD-10 Co	ode(s):	
3. REQUIRED CLINICAL INFORMATION PRIOR AUTHORIZATION.	: PLEASE PROVIDE ALL RELEVANT CLINIC	AL INFORMATION TO SUPPORT A
Clinical Information: Is the patient already taking the reque	ested medication? 🗆 Yes 🗆 No	
Has the patient received prior treatme	ent with Soliqua (insulin glargine/lixiser	natide)? □ Yes □ No
Was the patient's most recent HgbA1c product) 7% or greater? ☐ Yes ☐ No Copy of HgbA1c level required	c (within the past 6 months or prior to s	tarting a GLP-1/insulin combination
Has the patient tried or is the patient	currently receiving treatment with met	formin? Yes No
Is this patient's estimated GFR less that	an or equal to 45 mL/min/1.73 m2? \Box Y	es □ No
Does the patient have advanced liver	disease with at least one of the following	ng? □Yes □No
If yes, please select:		
Has treatment with metformin been a	voided due to lactic acidosis or elevate	d liver enzymes? □ Yes □ No
Is the patient currently taking any of to • Kazano (alogliptin/metformin) • Glyxambi (empagliflozin/linagliptin) • Kombiglyze XR (saxagliptin/metform • Nesina (alogliptin) • Onglyza (saxagliptin)		
Januvia (sitagliptin) Oseni (alogliptin/nioglitazone)		



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MEMBER'S FIRST NAME:

WEIGHDER S EAST MAINE:
• Janumet/Janumet XR (sitagliptin/metformin)
• Tradjenta (linagliptin)
• Jentadueto/Jentadueto XR (linagliptin/metformin)
• QTERN (dapagloflozin/saxagliptin)
• Seglujan (ertugliflozin/sitagliptin)
If yes, will concomitant therapy with those agents be discontinued? ☐ Yes ☐ No
if yes, will conconitant therapy with those agents be discontinued: Tes No
Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the
physician feels is important to this review?
Please note: Not all drugs/diagnoses are covered on all plans. This request may be denied unless all required
information is received.
ATTESTATION: I attest the information provided is true and accurate to the best of my knowledge. I understand that
the Health Plan, insurer, Medical Group or its designees may perform a routine audit and request the medical
information necessary to verify the accuracy of the information reported on this form.
Drossvihov Signatura av Flastvania I.D. Varification
Prescriber Signature or Electronic I.D. Verification: Date:
CONFIDENTIALITY NOTICE: The documents accompanying this transmission contain confidential health information that is legally privileged. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents
of these documents is strictly prohibited. If you have received this information in error, please notify the sender immediately (via return FAX)

FAX THIS FORM TO: 800-424-7640

MAIL REQUESTS TO: Prime Therapeutics Management Prior Authorization Program
Attn: CP - 4201

P.O. Box 64811 St. Paul, MN 55164-0811



and arrange for the return or destruction of these documents.

MEMBER'S LAST NAME: