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

MEMBER'S LAST NAME: _____

MEMBER'S FIRST NAME: _____

Instructions: Please fill out all applicable sections completely and legibly. Attach any additional documentation that is important for the review (e.g., chart notes or lab data, to support the authorization request). Information contained in this form is Protected Health Information under HIPAA.

URGENT

MEMBER INFORMATION		
LAST NAME:	FIRST NAME:	
PHONE NUMBER:	DATE OF BIRTH:	
STREET ADDRESS:		
CITY:	STATE: ZIP CODE:	
PATIENT INSURANCE ID NUMBER:		
MALE FEMALE HEIGHT (IN/CM): WEIGHT (LB/KG): ALLERGIES:		

IF YOU ARE NOT THE PATIENT OR THE PRESCRIBER, YOU WILL NEED TO SUBMIT A PHI DISCLOSURE AUTHORIZATION FORM WITH THIS REQUEST WHICH CAN BE FOUND AT THE FOLLOWING LINK: <u>PRIMETHERAPEUTICS.COM/NOPP</u>

PATIENT'S AUTHORIZED REPRESENTATIVE (IF APPLICABLE):

AUTHORIZED REPRESENTATIVE'S PHONE NUMBER:

PRESCRIBER INFORMATION		
LAST NAME:	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:	
DURATION OF THERAPY (SPE	RENEWAL CIFIC DATES):	IF RENEWAL: DATE THERAPY INITIATED:		

Continued on next page.



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MEMBER'S LAST NAME:	MEMBER'S FIRST I	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 II diabetes Type II diabetes with established cardiovascular disease Congestive heart failure Other DiagnosisICD-10 Code(s): 				
3. REQUIRED CLINICAL INFORMATION: PRIOR AUTHORIZATION.	PLEASE PROVIDE ALL RELEVANT CLINIC	AL INFORMATION TO SUPPORT A		
Clinical information: Has patient tried Jardiance(empagliflozin) AND Trajenta(linagliptin)/Jentadueto as single entities? □ Yes □ No Please provide documentation. Is the patient a Type II diabetic? □ Yes □ No If prescribing for Type II Diabetes, please answer the following: Is the patient's estimated glomerular filtration rate (eGFR) below 45 mL/min/1.73 m2? □ Yes □ No Please provide documentation.				
Is the patient's most recent (pre-Jardiance) HgbA1C obtained in the past 6 months or prior to starting TrijardyXR(empagliflozin/linagliptin/metformin) 7% or greater? u Yes u No <i>Please provide documentation.</i> Is the patient on dialysis? u Yes u No				
Is the patient currently on metformin? Yes No				
Did the patient have an inadequate response or intolerance to metform? Please provide documentation				
Does the patient have at least one of the following contraindications to metformin? Yes No (Please Check one) Estimated glomerular filtration rate (eGFR) less than or equal to 30 mL/min/1.73 m ² Advanced liver disease with cirrhosis, portal hypertension, ascites, and/or hepatic encephalopathy				
Is the patient's most recent hemoglobin A1c level within the past 6months or prior to starting TrijardyXR(empagliflozin/linagliptin/metformin) 7.0–10%, inclusive? □ Yes □ No Please provide documentation.				
Does the patient's body mass index(BMI) exceed 45 kg/m ² ? ☐ Yes ☐ No				
Is the patient's estimated glomerular filtration rate (eGFR) above 30 mL/min/1.73 m ² ? D Yes D No Please provide documentation.				



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Is the patient's medical history positive for a Please check at least one of the following: MI or Stroke	t least one of the following? □ Yes □ No			
 Imaging shows single-vessel or multi-ve Previous coronary revascularization pro 				
 Positive cardiac stress test Hospital admission for unstable angina 				
	efined as limb revascularization procedure, limb or foot amputation			
	g or non-invasive study showing evidence of more than 50% stenosis in			
For diagnosis of congestive heart failure, plea	ase answer the following:			
Does patient have an ejection fraction (EF) e	qualing 40% or less? Ves No Please provide documentation.			
Does patient have an ejection fraction (EF) g	reater than 40%? Please provide documentation.			
Has patient ever had NYHA class II, III or IV sy	ymptoms of heart failure? Yes No Please provide documentation.			
Does patient's body mass index (BMI) equal	less than 45 kg/m ² ? Yes No Please provide documentation.			
Does patient have a NT-proBNP greater than	300 pg/ml? Yes No Please provide documentation.			
For patients with A-fib, is the NT-proBNP gre	ater than 900 pg/ml? Yes No Please provide documentation.			
IF NT-proBNP not available, does patient have a BNP >100 pg/ml without kidney failure? Yes No Please submit chart documentation.				
If NT-proBNP not available and patient has k Please submit chart documentation.	idney failure, does patient have a BNP>200 pg/ml? □ Yes □ No			
If NT-proBNP not available and patient has A Please submit chart documentation	trial fibrillation(AF), does patient have a BNP >150 pg/ml? \Box Yes \Box No			
Please provide documentation from echocard	se such as one or more of the following:? ☐ Yes ☐ No diogram.			
□ LA width >4.0cm				
 LA length >5.0 cm LA area >20cm2 				
□ LA area >20cm2 □ LA volume >55ml				
□ LA volume index >34ml/m2				
	ophy defined by at least one of the following:? \Box Yes \Box No			
Please provide documentation from echocardiogram.				
□ Septal thickness or posterior wall thickness >1.1 cm				
 LV mass index(LVMI) >115g/m2 for main to E/e² (mean septal and lateral) >13 	ies and >95 g/m2 for temaies			
\Box e' (mean septal and lateral) <9cm/s				
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CAT0122

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Has patient been hospitalized in the past 12 months before starting Trijardy XR (empagliflozin/linagliptin/ metformin) ?
Solve Yes
No Please provide documentation.

Is patient on a stable dose of a diuretic?

Yes
No Please provide documentation.

Has patient had a myocardial infarction, coronary bypass graft surgery or other major cardiovascular surgery, stroke or TIA in the past 90 days of starting Jardiance?
□ Yes □ No Please provide documentation.

Has patient had a heart translplant?

Yes No

Does patient have acute decompensated heart failure?

Ves
No

Does patient have severe <u>pulmonary disease</u> including severe COPD, requiring home oxygen therapy for their COPD, chronic nebulizer therapy or chronic oral steroid therapy for treatment of their severe COPD?
Yes No Please submit chart documentation.

Does patient have severe <u>pulmonary disease</u> including primary pulmonary hypertension?
□ Yes □ No Please submit chart documentation.

Does patient have any other condition or diagnosis causing patient's heart failure symptoms such as patient has significant mitral valve regurgitation causing the heart failure, any dilated cardiomyopathy, infiltrative cardiomyopathy, drug induced cardiomyopathy, or viral myocarditis?
□ Yes □ No Please submit chart documentation.

Does patient have and eGFR less than 20ml/min/1.73m²?
□ Yes □ No

Does patient require dialysis?

Yes
No

Is patient's heart failure related to any of the following?
□ Yes □ No Please check at least one of the following:

- □ infiltrative disease
- accumulation disease
- muscular dystrophy
- □ hypertrophic obstructive cardiomyopathy
- □ known pericardial restriction
- valvular disease expected to lead to surgery
- $\hfill\square$ atrial fib/flutter with a resting heart rate greater than 110 bpm

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/diagnosis are covered on all plans. This request may be denied unless all required information is received.



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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.

Prescriber Signature or Electronic I.D. Verification:

Date:

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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

