

Kerendia

Patient Ir	nformation:			
Name:				
Member	ID:			
Address:				
City, Stat	e, Zip:			
Date of B	irth:			
	er Information:			
Name:				
NPI:				
Phone No				
Fax Num	ber			
Address:				
City, Stat	e, Zip:			
Requeste	ed Medication			
Rx Name	:			
Rx Streng	gth			
Rx Quant	tity:			
Rx Frequ	ency:			
Rx Route	of			
Administr	ation:			
Diagnosis	s and ICD Code:			
prescribed quantities of Upon rece SECTIO requests medicati	a medication for your an be provided. Plea ipt of the completed in A: Please not a Pharmacy prince that are not ions that are not in the provided in the provi	efit requires that we review certain requests for coverage with the presentation that requires Prior Authorization before benefit coverage or coverage complete the following questions then fax this form to the toll-free number of form, prescription benefit coverage will be determined based on the that supporting clinical documentation is required or authorization reviews can be subject to trial with a state of the trial with a listed within the criteria. The policies are subject to trial with a listed within the criteria. The policies are subject to trial with a listed within the criteria. The policies are subject to the listed within the criteria with a listed within the criteria.	erage of a umber liste the plar for ALI iddition change	additional ed below. n's rules. LPA al
1	Is the request an II	NITIAL or CONTINUATION of therapy?		
	[] Initial (If checked	. •		
2	[] Continuation (If o	checked, go to 2) ently receiving the requested medication?	Yes	No
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If you have any questions, call: 1-888-258-8250

Version 09.2025

	[If no, skip to question 7.]		
3	Has the patient been receiving medication samples of the requested medication? [If yes, skip to question 7.]		No
4	Does the patient have a previously approved prior authorization (PA) on file with the current plan? [NOTE: If the patient does NOT have a previously approved PA on file for the	Yes	No
	requested medication with the current plan, the renewal request will be considered under initial therapy.]		
	[If no, skip to question 7.]		
5	Has the patient been established on therapy for at least 3 months? [If no, skip to question 7.]	Yes	No
6	Has documentation been submitted to confirm that the patient has had a significant response to therapy, as determined by the provider? ACTION REQUIRED: Submit supporting documentation. [No further questions.]	Yes	No
7		V	NIa
7	Is the patient greater than or equal to 18 years of age? [If no, no further questions.]	Yes	No
8	Does the patient have a diagnosis of type 2 diabetes?	Yes	No
	[If no, no further questions.]		
9	Have non-diabetic kidney disease processes been evaluated and ruled out? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[If no, no further questions.]		
10	Does the patient have chronic heart failure with reduced ejection fraction and persistent symptoms (New York Heart Association [NYHA] Class II - IV)? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[If yes, no further questions.]		
11	Has the patient required dialysis for acute renal failure within the last 90 days?	Yes	No

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	ACTION REQUIRED: Submit supporting documentation.		
	[If yes, no further questions.]		
12	Has the patient experienced a stroke, transient ischemic attack (TIA), acute coronary syndrome or required hospitalization for worsening heart failure within the last 30 days? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[If yes, no further questions.]		
13	Does the patient have hepatic insufficiency classified as Child-Pugh Class C?	Yes	No
	[If yes, no further questions.]		
14	Has the patient currently been receiving a maximally tolerated dose of a sodium-glucose co-transporter 2 (SGLT2) inhibitor for at least 3 months? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[If yes, skip to question 16.]		
15	Does the patient have a contraindication to the use of sodium-glucose cotransporter 2 (SGLT2) inhibitors? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[If no, no further questions.]		
16	Has the patient currently been receiving a maximally tolerated labeled dosage of an angiotensin converting enzyme (ACE) inhibitor or angiotensin receptor blocker (ARB) for 4 weeks or is there a documented contraindication? ACTION REQUIRED: Submit supporting documentation.	Yes	No
	[If no, no further questions.]		
17	Prior to initiation, does the patient have ALL of the following: A) Estimated glomerular filtration rate greater than or equal to 25 mL/min/1.73 m2, B) Urine albumin-to-creatinine ratio greater than or equal to 30 mg/g, C) Serum potassium level between 3.5 to 5.0 mEq/L? ACTION REQUIRED: Submit supporting documentation.	Yes	No

Please document the diagnoses, symptoms, and/or any other information important to this review:							



SECTION B: Physician Signature

PHYSICIAN SIGNATURE

DATE

FAX COMPLETED FORM TO: 1-833-896-0656

Disclaimer: An authorization is not a guarantee of payment. Member must be eligible at the time services are rendered. Services must be a covered Health Plan Benefit and medically necessary with prior authorization as per Plan policy and procedures.

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