BlueCross BlueShield Federal Employee Program.

FARXIGA / JARDIANCE
PRIOR APPROVAL REQUEST

Additional information is required to process your claim for prescription drugs. Please complete the patient portion, and have the prescribing physician complete the provider portion and submit this completed form.

Send completed form to:
Service Benefit Plan
Prior Approval
P.O. Box 52080 MC 139
Phoenix, AZ 85072-2080
Attn. Clinical Services
Fax: 1-877-378-4727

Patient Information (required)			**Provider Information (required)**		
Date:			Provider Name:		
Patient Name:			Specialty:	NPI:	
Date of Birth:	Sex: ❑Male ❑Female		Office Phone:	Office Fax:	
Street Address:			Office Street Address:		
City:	State:	Zip:	City:	State:	Zip:
Patient ID: **R**			Physician Signature:		
PHYSICIAN COMPLETES					

NOTE: Form must be completed in its **entirety** for processing

Please select medication: ❑**Farxiga (dapagliflozin)** ❑**Jardiance (empagliflozin)**

****Check www.fepblue.org/formulary to confirm which medication is part of the patient's benefit**

Is this request for brand or generic? ❑Brand ❑Generic

1. What is the patient's diagnosis?

❑Chronic kidney disease (CKD)

a. Has the patient been on this medication continuously for the last **6 months** excluding samples? ***Please select answer below:***

❑**NO** – this is **INITIATION** of therapy, please answer the following questions:

i. Does the patient have a diagnosis of polycystic kidney disease (PKD)? ❑Yes ❑No

ii. Is the patient currently using or does the patient have a recent history of immunosuppressive therapy for the treatment of kidney disease (e.g., tacrolimus, sirolimus, cyclosporine, mycophenolate, etc)? ❑Yes ❑No

❑**YES** – this is a PA renewal for **CONTINUATION** of therapy, please answer the following question:

i. Has the patient had a reduced decline in renal function? ❑Yes ❑No

❑Heart failure (HF)

a. Has the patient been on this medication continuously for the last **6 months** excluding samples? ❑Yes***** ❑No

****If YES***, has the patient's symptoms improved or stabilized with therapy? ❑Yes ❑No

❑Type 2 diabetes mellitus (DM)

a. Has the patient been on this medication continuously for the last **6 months** excluding samples? ❑Yes ❑No*****

****If NO***, please answer the following questions:

i. Does the patient have an intolerance or contraindication or have they had an inadequate treatment response to metformin? ❑Yes ❑No*****

****If NO***, has the patient had an intolerance or contraindication or have they had an inadequate treatment response to one of the following metformin combination medications: Glucovance, Janumet/Janumet XR, Jentadueto/Jentadueto XR, Kazano, Kombiglyze XR, or Trijardy XR? ❑Yes ❑No

ii. **Standard Option:** Is this medication being requested as a change from one of the following to allow the member access to their copay benefit: Invokana, Invokamet, Invokamet XR, Steglatro, Steglujan, or Segluromet? ❑Yes***** ❑No

****If YES***, select drug: ❑Invokamet/Invokamet XR ❑Invokana ❑Segluromet ❑Steglatro ❑Steglujan

❑Other ***(please specify)***: _____

2. Will this medication be used in combination with other SGLT2 inhibitors? ❑Yes***** ❑No

****If YES***, please specify the medication: _____

****SGLT2 inhibitors: Brenzavvy (bexagliflozin), Farxiga (dapagliflozin), Glyxambi (empagliflozin/linagliptin), Inpefa (sotagliflozin), Invokamet/Invokamet XR (canagliflozin/ metformin), Invokana (canagliflozin), Jardiance (empagliflozin), Qtern (dapagliflozin/ saxagliptin), Qternmet XR (dapaglifozin/saxagliptin/metformin), Segluromet (ertugliflozin/metformin), Steglatro (ertugliflozin), Steglujan (ertugliflozin/ sitagliptin), Synjardy/Synjardy XR (empagliflozin/metformin), Trijardy XR (empagliflozin /linagliptin/ metformin), Xigduo XR (dapagliflozin/metformin)***

The information provided on this form will be used to determine the provision of healthcare benefits under a U.S. federal government program, and any falsification of records may subject the provider to prosecution, either civilly or criminally, under the False Claim Acts, the False Statements Act, the mail or wire fraud statutes, or other federal or state laws prohibiting such falsification. **Prescriber Certification:** I certify all information provided on this form to be true and correct to the best of my knowledge and belief. I understand that the insurer may request a medical record if the information provided herein is not sufficient to make a benefit determination or requires clarification and I agree to provide any such information to the insurer. Farxiga/Jardiance - FEP MD Fax Form Revised 4/26/2024

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Message:

Attached is a Prior Authorization request form.

For your convenience, there are 3 ways to complete a Prior Authorization request:

Electronically Online (ePA) Results in 2-3 **minutes** **FASTEST AND EASIEST**	Now you can get responses to drug Prior Authorization requests **securely** online. **Online** submissions may receive **instant** responses and do not require faxing or phone calls. Requests can be made 24 hours a day, 7 days a week. For more information on electronic prior authorization (ePA) and to register, go to **Caremark.com/ePA.**
Phone (4-5 **minutes** for response)	The FEP Clinical Call Center can be reached at **(877)-727-3784** between the hours of 7AM-9PM Eastern Time. A live representative will assist with the Prior Authorization, asking for the same information contained on the attached form. Please review the form and have your answers ready for faster service. The process over the phone takes on average between 4 and 5 minutes.
Fax (3-5 **days** for response)	Fax the attached form to **(877)-378-4727**. Requests sent via fax will be processed and responded to within 5 business days. The form must be filled out completely, if there is any missing information the Prior Authorization request cannot be processed. ***Please only fax the completed form once as duplicate submissions may delay processing times.***