



Federal Employee Program. **WELIREG** PRIOR APPROVAL REQUEST

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

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

Patient Information (required)				Provider Information (required)		
Date:				Provider Name:		
Patient Name:				Specialty:		NPI:
Date of Birth:		Sex: <input type="checkbox"/> Male <input type="checkbox"/> Female		Office Phone:		Office Fax:
Street Address:				Office Street Address:		
City:		State:	Zip:	City:		State: Zip:
Patient ID: <b>R</b>				Physician Signature:		
<b>PHYSICIAN COMPLETES</b>						

**Welireg (belzutifan)**

**\*\*Check [www.fepblue.org/formulary](http://www.fepblue.org/formulary) to confirm which medication is part of the patient's benefit**

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

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

1. Will the patient need more than 120 milligrams per day? ☐ Yes\* ☐ No

**\*If YES**, please specify the requested milligrams per day? \_\_\_\_\_ mg per day

2. Is the patient's hemoglobin greater than or equal to 8 grams per deciliter (g/dL)? ☐ Yes ☐ No

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

☐ **NO** – this is **INITIATION**, please answer the following questions:

a. Does the prescriber agree to monitor for anemia and hypoxia before initiation of treatment and periodically throughout treatment? ☐ Yes ☐ No

b. **MALE Patient:** Does the patient have a female partner of reproductive potential? ☐ Yes\* ☐ No

**\*If YES**, will pregnancy be excluded before the start of treatment? ☐ Yes\* ☐ No

**\*If YES**, will the patient be advised to use effective non-hormonal contraception during treatment with Welireg and for 1 week after the last dose? ☐ Yes ☐ No

c. **FEMALE Patient:** Is the patient of reproductive potential? ☐ Yes\* ☐ No

**\*If YES**, has the patient had a negative pregnancy test? ☐ Yes\* ☐ No

**\*If YES**, will the patient be advised to use effective non-hormonal contraception during treatment with Welireg and for 1 week after the last dose? ☐ Yes ☐ No

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

a. Does the prescriber agree to monitor for anemia and hypoxia periodically throughout treatment? ☐ Yes ☐ No

b. Has the patient experienced disease progression or unacceptable toxicity while on the requested therapy? ☐ Yes ☐ No

c. **MALE Patient:** Does the patient have a female partner of reproductive potential? ☐ Yes\* ☐ No

**\*If YES**, will the patient be advised to use effective non-hormonal contraception during treatment with Welireg and for 1 week after the last dose? ☐ Yes ☐ No

d. **FEMALE Patient:** Is the patient of reproductive potential? ☐ Yes\* ☐ No

**\*If YES**, will the patient be advised to use effective non-hormonal contraception during treatment with Welireg and for 1 week after the last dose? ☐ Yes ☐ No

4. What is the patient's diagnosis?

☐ Advanced renal cell carcinoma (RCC)

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. Has the patient received previous treatment with a PD-1 inhibitor OR PD-L1 inhibitor? ☐ Yes ☐ No

ii. Has the patient received previous treatment with a VEGF tyrosine kinase inhibitor? ☐ Yes ☐ No

☐ Von Hippel-Lindau (VHL) disease

a. Does the patient require therapy for associated renal cell carcinoma (RCC), central nervous system (CNS) hemangioblastomas, or pancreatic neuroendocrine tumors (pNET)? ☐ Yes ☐ No

b. Does the patient require immediate surgery? ☐ Yes ☐ No

**PLEASE PROCEED TO PAGE 2 FOR ADDITIONAL DIAGNOSES**

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**BlueCross  
BlueShield**

Federal Employee Program.

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**PAGE 3 – PHYSICIAN COMPLETES**

**Patient Name:** \_\_\_\_\_ **DOB:** \_\_\_\_\_ **Patient ID: R** \_\_\_\_\_

☐ Pheochromocytoma or paraganglioma (PPGL)

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

☐ **NO** – this is **INITIATION**, please answer the following question:

i. Is the tumor locally advanced, unresectable, or metastatic? ☐ Yes ☐ No

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

i. Is the tumor advanced, unresectable, or metastatic? ☐ Yes ☐ No

☐ None of the above

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