

BlueShield. LENVIMA
Federal Employee Program. 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 physician 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:	NPI:		
Date of Birth:		Sex:  Male  Female		Office Phone:	Office Fax:	Office Fax:	
Street Address:				Office Street Address:			
City:		State:	Zip:	City:	State:	Zip:	
Patient ID: R				Physician Signature:			
<u> </u>			PHYSICIAN	COMPLETES			
				(lenvatinib)			
	**Check	www.fenblue.org/fo		t (lenvaumo) t which medication is part o	of the natient's benefit		
	Check			_	_		
		NOTE: Form	must be complet	ed in its <b>entirety</b> for pr	ocessing		
1. Has the patien	nt been on Lenvin	na continuously f	or the last <b>6 mo</b>	nths, excluding sample	s? Please select answer	below:	
$\Box$ <b>YES</b> – this	is a PA renewal f	for CONTINUA	TION of therap	y, please answer the qu	estions on PAGE 2		
$\square$ <b>NO</b> – this is	s INITIATION	of therapy, please	e answer the que	stions below:			
2. Is this request	for brand or gene	eric? 🗆 Brand	Generic				
3. What is the page	atient's diagnosis	?					
•	Endometrial Carc						
a. Will L	envima be used ir	combination wi	th pembrolizum	ab (Keytruda)? □Yes	□No		
	patient's advanced not microsatellite				IMR), as determined by	an FDA-approved	
c. Has th	e patient experien	ced disease prog	ression followin	g prior systemic therap	y? □Yes □No		
d. Is the	patient a candidate	e for curative sur	gery or radiation	n? □Yes □No			
☐ Differentia	ted Thyroid Canc	er (DTC)					
	differentiated thyr		y recurrent?	Yes □No*			
* <b>I</b> f 1	<b>VO</b> , is the differen	tiated thyroid ca	ncer metastatic?	□Yes □No			
b. Has th	e patient had dise	ase progression a	after radioactive	iodine therapy (radioac	tive iodine-refractory)?	□Yes □No	
☐ Renal Cell	Carcinoma (cance	er) (RCC)					
a. Does t	he patient have ad	lvanced renal cel	l carcinoma (kid	ney cancer)? □Yes	□No		
b. Will L	envima be used in	n combination wi	th pembrolizum	ab (Keytruda) as a first	-line treatment? □Yes	□No	
	•	1 0	•	nti-angiogenic therapy? erolimus (Afinitor)?			
_	le Hepatocellular						
	envima be used as	,	,	lNo			
	envima be used a						
		•			eairment)? □Yes □No	)	
_	nosis ( <i>please spec</i>			- •			

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Date:

**Patient Information** (required)

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**Provider Information** (required)

Patient Name:			Specialty:	NPI	[:		
Date of Birth:		Sex:  Male  Female		Office Phone:	Offi	Office Fax:	
Street Address:				Office Street Address:			
City:		State:	Zip:	City:	State:	Zip:	
Patient ID: R	, ,	1 1 1	<del> </del>	Physician Signature:			
		]	PHYSICIAN	COMPLETES			
$\square$ <b>NO</b> – this i $\square$ <b>YES</b> – this	*Check on the state of the stat	www.fepblue.org/for  NOTE: Form r  na continuously for of therapy, please for CONTINUA	Lenvimal mulary to confirm must be complete for the last 6 most answer the que	IERAPY (PA RE (lenvatinib) which medication is part of the season of t	the patient's benefi scessing ? Please select a		
□ Advanced I a. Will L □ Differentia □ Renal Cell a. Does th □ Unresectab	ted Thyroid Cand Carcinoma (cand he patient have ad	einoma (EC) n combination with ter (DTC) ter) (RCC) dvanced renal cell c Carcinoma (HCC)	l carcinoma (kid	ab (Keytruda)? □Yes Iney cancer)? □Yes □			
4. Has the patier	nt experienced dis	sease progression	while on Lenvi	ma? □Yes □No			
cardiac dysfur severe QT pro	nction, hepatotox olongation (grade	icity, nephrotic sy	yndrome, renal i		ointestinal perfo	hypertension, severe oration/fistula formation, or arterial thromboembolic	

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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 <b>securely</b> online. <b>Online</b> submissions may receive <b>instant</b> 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 <b>Caremark.com/ePA.</b>
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.

better...

faster... Introducing ePA! Online Prior Authorizations in minutes through Caremark.com/ePA. Sign up today!

CVS/caremark<sup>\*</sup>

