

## BlueShield. METHOTREXATE INJECTIONS Federal Employee Program. 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 patient 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:		Office Phone:		Office Fax:	
Street Address:				Office Street Address:			
City:		State:	Zip:	City:	Sta	te:	Zip:
Patient ID: <b>R</b>	1 1			Physician Signar	ture:		
PHYSICIAN COMPLETES							
Methotrexate Injections							
NOTE: Form must be completed in its entirety for processing							
Please select m	edication:	□Otrexup		Rasuvo		□RediTrex	
**Check www.fepbl	ue.org/formulary to	confirm which medic	ation is part of the	patient's benefit			
Is this request for	brand or generic	? □Brand □Ge	eneric				
1. Has the patien	t been on this me	dication continuou	ısly for the last	6 months, exclud	ding samples? Plea	ise select answ	er below:
□ NO – this is <b>INITATION</b> of therapy, please answer the following questions: a. What is the patient's diagnosis?							
□Polyarticular Juvenile Idiopathic Arthritis (pJIA)							
i. Does the patient have a capturing discription on how they had an inches active resonance or intelegence to NSAIDs and							
<ul><li>ii. Does the patient have a contraindication or have they had an inadequate response or intolerance to NSAIDs, and oral methotrexate? □Yes □No</li></ul>							noaids, and
□Psoriasis (PsO) i. Does the patient have active psoriasis? □Yes □No							
ii. Does the patient have a contraindication or have they had either an inadequate response or intolerance to NSAIDs, topical corticosteroids and oral methotrexate? □Yes □No							
□Rheumatoid Arthritis (RA)  i. Does the patient have severely active rheumatoid arthritis? □Yes □No							
<ul> <li>ii. Does the patient have a contraindication or have they had an inadequate response intolerance to NSAIDs, and oral methotrexate? □Yes □No</li> </ul>							
□Other diagnosis (please specify):							
b. Is there a documented reason for requiring a special injection device? \( \text{YES}, \text{ select reason below} \) \( \text{No} \) \( \text{Lack of dexterity} \) \( \text{Visual acuity issues} \) \( \text{Other (please specify):} \)							
□YES – this is a PA renewal for CONTINUATION of therapy, please answer the following questions:  a. What is the patient's diagnosis?							
	•	•	is (JIA) □Pso	oriasis (PsO)	□ Rheumato:	id Arthritis (R	(A)
□Polyarticular Juvenile Idiopathic Arthritis (JIA) □Psoriasis (PsO) □Rheumatoid Arthritis (RA) □Other diagnosis ( <i>please specify</i> ):							·
b. Has the patient's condition improved or stabilized with therapy? □Yes □No							
2. <b>FEMALE Pa</b>	tient: Is the patien	nt of reproductive	potential? <b>\P</b> Y	es* □No			
*If YES, has the patient had a negative pregnancy test prior to initiating therapy? $\square \text{Yes*}$ (*If YES, answer question below) $\square \text{No}$							
a. <b>Otrexup or Reditrex Request:</b> Will the patient be advised to use effective contraception during treatment with this medication and for six months after the final dose? □Yes □No							
Rasuvo Request: Will the patient be advised to use effective contraception during treatment with Rasuvo and for at least one ovulatory cycle after the final dose? □Yes □No						o and for at least	
one of the state o							

PLEASE PROCEED TO PAGE 2 FOR ADDITIONAL QUESTIONS

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PAGE 2 - PHYSICIAN COMPLETES							
Patient Name:	DOB:	Patient ID: R					
MALE Patient: Does the patient have a female partner of reproductive potential? □Yes* □No *If YES, will the patient be advised to use effective contraception during treatment with this medication and for three months after the final dose? □Yes □No							
3. Does the prescriber agree to comply with regular monitoring of blood counts, renal function, and hepatic function testing? □Yes □No							
4. Will this medication be used in *If YES, specify the medica	combination with another form or tion:	brand of methotrexate? □Yes*	□No				

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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.

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