





## **CVS caremark** Arkansas Formulary Exception/Prior Approval Request Form

This fax machine is located in a secure location as required by HIPAA regulations. Complete/review information, sign and date. Fax signed forms to **CVS/Caremark at 1-855-245-2134 for prior approval, step therapy, and quantity limit requests**. Please contact CVS/Caremark at **1-855-582-2022** with questions regarding the prior approval, step therapy, and quantity limit review process.

For Non-Formulary Exception requests, fax the form to 501-378-6980. For Non-Formulary request questions, contact 501-378-3392.

Patient Information				Prescriber Information			
Patient Name:			Prescriber Nam	Prescriber Name:			
Patient ID#:							
Address:			Address:	Address:			
City:	State:		City:			State:	
Home Phone:	ZIP:		Office Phone:		Office Fax:	ZIP:	
Gender: M or F	DOB:	DOB:		Contact Person at Doctor's Office:			
		Medicati	on Requesting and D	iagnosis			
Medication: Stre		Strength:		Directions for use (Frequency):			
Expected Length of Therapy:		Qty:	Day Supply:		If this is a continuation of therapy, how long has the patient been on the medication?		
Diagnosis:			Diagnosis (ICD)	Diagnosis (ICD) Code(s):			

## PLEASE ATTACH ALL RELEVANT CLINICAL DOCUMENTATION TO SUPPORT USE OF THIS MEDICATION WITH REQUEST

Expedited/Urgent Review Requested: By checking this box and signing below, I certify that applying the standard review time frame may seriously jeopardize the life or health of the patient or the patient's ability to regain maximum function.

Please list all medications the patient has tried specific to the diagnosis and specify below:

Medication Name	Trial period	Reason for Failure
0	0	0
0	0	0
0	0	0
0	0	0
0	0	0

Does the patient have a clinical condition for which other formulary alternatives are not recommended or are contraindicated due to comorbidities or drug interactions based on published clinical literature? *If so, please provide documentation including medication names and clinical reasons.* 

Is the request for a patient with one or more chronic conditions (e.g., psychiatric condition, epilepsy, dementia) who is stable on the current drug(s) and who might be at high risk for a significant adverse event with a medication change? *If yes, specify anticipated significant adverse event:* 

PRESCRIPTION BENEFIT PLAN MAY REQUEST ADDITIONAL INFORMATION OR CLARIFICATION, IF NEEDED, TO EVALUATE REQUESTS. I attest that the medication requested is medically necessary for this patient. I further attest that the information provided is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark, the health plan sponsor, or, if applicable, a state or federal regulatory agency. I understand that any person who knowingly makes or causes to be made a false record or statement that is material to a claim ultimately paid by the United States government or any state government may be subject to civil penalties and treble damages under both the federal and state False Claims Acts. *See, e.g.*, 31 U.S.C. §§ 3729-3733.

Prescriber Signature:

Date:

**Confidentiality Notice**: The documents accompanying this transmission contain confidential health information that is legally privileged. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution of these documents is strictly prohibited. If you have received this information in error, please notify the sender immediately (via return fax) and arrange for the return or destruction of these documents.

106-37207B 043019

Plan member privacy is important to us. Our employees are trained regarding the appropriate way to handle members' private health information. This document contains references to brand-name prescription drugs that are trademarks or registered trademarks of pharmaceutical manufacturers not affiliated with CVS Caremark<sup>®</sup>.