



**MEDICARE ADVANTAGE
PRIOR AUTHORIZATION REQUEST FORM**

High Risk Medication - Lorazepam

Phone: 215-991-4300

Fax back to: 866-371-3239

Jefferson Health Plans manages the pharmacy drug benefit for your patient. Certain requests for coverage require review with the prescribing physician. Please answer the following questions and fax this form to the number listed above.

PLEASE NOTE: Any information (patient, prescriber, drug, labs) left blank, illegible, or not attached WILL delay the review process.

Member Name:	Prescriber Name:	
Member Number:	Fax:	Phone:
Date of Birth:	Office Contact:	
Line of Business: <input type="checkbox"/> Medicare Advantage	NPI:	State Lic ID:
Address:	Address:	
City, State ZIP:	City, State ZIP:	
Primary Phone:	Specialty/facility name (if applicable):	

☐ **REQUEST FOR EXPEDITED REVIEW:** By checking this box and signing below, I certify that applying the 72 hour standard review timeframe may seriously jeopardize the life or health of the enrollee or the enrollee's ability to regain maximum function.

Drug Name:	
Strength:	
Directions / SIG:	

Please attach any pertinent medical history including labs and information for this member that may support approval.
Please answer the following questions and sign.

Q1. Is this an initial request for Lorazepam? If YES go to 3, If NO go to 2.

☐ Yes

☐ No

Q2. Has the prescriber provided an explanation that the benefit continues to outweigh the potential risk of the high-risk medication with concurrent opioid therapy?

☐ Yes

☐ No

Q3. Is the patient 65 years of age or older? If YES, go to 4.

☐ Yes

☐ No

Q4. Are chart notes attached documenting an explanation of the risk versus benefit profile which shows the benefit outweighs the potential risk for the use of the high-risk medication with concurrent opioid therapy? If YES, go to 5.

☐ Yes

☐ No

Q5. Has the prescriber provided an attestation of intent to monitor and address treatment-related adverse events? If YES, go to 6. [see Q12-Q14 for attestation]



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Member Name:	Prescriber Name:
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Q6. Are chart notes attached documenting the patient has anxiety disorder? If YES, go to 7. If NO, go to 9.	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Q7. Is the requested drug is being used concurrently with a selective serotonin reuptake inhibitor (SSRI) or serotonin-norepinephrine reuptake inhibitor (SNRI) until the SSRI/SNRI becomes effective for the symptoms of anxiety? If YES, go to 10. If NO, go to 8.	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Q8. Is documentation attached showing inadequate response, intolerance, or contraindication to AT LEAST ONE agent from EACH of the following classes: a) selective serotonin reuptake inhibitors (SSRIs), b) serotonin-norepinephrine reuptake inhibitors (SNRIs)? If YES, go to 10.	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Q9. Are chart notes attached documenting the patient has insomnia? If YES, go to 10. If NO, go to 11.	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Q10. Is documentation attached showing inadequate response, intolerance, or contraindication to two non-high risk alternatives, such as trazodone, mirtazapine, doxepin 3 mg or 6 mg, ramelteon. If YES, go to 12.	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Q11. Are chart notes attached documenting the patient is taking the requested medication for an FDA approved indication?	
<input type="checkbox"/> Yes <input type="checkbox"/> No	
Q12. I have educated the member in regards to the risks of concurrent utilization of benzodiazepine and opioid therapy, and the member accepts these risks.	
<input type="checkbox"/> Yes <input type="checkbox"/> No	



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Member Name:	Prescriber Name:
<p>Q13. If applicable, I have consulted other prescribers involved in concurrent therapy and all prescribers involved agree to pursue concurrent opioid and benzodiazepine therapy for this member.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> NA</p>	
<p>Q14. I acknowledge, as the prescriber initiating or maintaining concurrent benzodiazepine and opioid therapy, the risk of adverse event(s) associated with concurrent utilization.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q15. Requested Duration:</p> <p><input type="checkbox"/> 12 Months <input type="checkbox"/> Other</p>	
<p>Q16. Additional Information:</p>	

Prescriber Signature

Date

v2026