

Member Name: {{MEMFIRST}} {{MEMLAST}} DOB: {{MEMBERDOB}} PA Number: {{PANUMBER}}



{{PANUMCODE}}

Xywav

Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-866-249-6155.** If you have questions regarding the prior authorization, please contact CVS Caremark at 1-866-814-5506. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name: {{MEMFIRST}} {{MEMLAST}} **Date:** {{TODAY}}
Patient's ID: {{MEMBERID}} **Patient's Date of Birth:** {{MEMBERDOB}}
Physician's Name: {{PHYFIRST}} {{PHYLAST}}
Specialty: _____, **NPI#:** _____
Physician Office Telephone: {{PHYSICIANPHONE}} **Physician Office Fax:** {{PHYSICIANFAX}}
Request Initiated For: {{DRUGNAME}}

- What is the diagnosis?
 Cataplexy with narcolepsy
 Excessive daytime sleepiness with narcolepsy
 Other _____
- What is the ICD-10 code? _____
- Coverage for the requested drug is provided when the patient has tried and had a treatment failure with the formulary medication. The formulary alternative for the requested drug is Xyrem. Can the patient's treatment be switched to the formulary alternative? ***If Yes, please call 1-866-814-5506 to have the updated form faxed to your office OR you may complete the PA electronically (ePA). You may sign up online via CoverMyMeds at: www.covermymeds.com/epa/caremark/ or call 1-866-452-5017.***
 Yes
 No - Continue request non-formulary medication
- Has the patient tried and had a documented inadequate response or intolerable adverse reaction to the formulary alternative? Note: Formulary medications should be prescribed first unless the patient is unable to use or receive treatment with the alternative. Yes No

Formulary alternative(s): Xyrem

If Yes, indicate the formulary alternative the patient has tried and the reason for treatment failure and skip to #6.

Drug name: _____ Reason for treatment failure: _____

- Does the patient have a documented contraindication to the formulary alternative: Xyrem? Yes No
If No, complete this form in its entirety and State Step Therapy section.

If Yes, indicate the formulary alternative the patient is unable to take and describe the contraindication(s):

Drug name: _____ Contraindication: _____

Send completed form to: Case Review Unit, CVS Caremark Prior Authorization. Fax: 1-866-249-6155

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6. Has chart note(s) or other documentation supporting the inadequate response, intolerable adverse reaction or contraindication to the necessary number of formulary alternatives been submitted? ***ACTION REQUIRED: Submit chart note(s) or other documentation indicating prior treatment failure, severity of the adverse event (if any), and dosage and duration of the prior treatment, or contraindication to formulary alternatives.***
 Yes No *If No, complete this form in its entirety and State Step Therapy section.*
7. Has diagnosis of narcolepsy been confirmed by a sleep lab evaluation? Yes No
8. Is the patient currently receiving treatment with the requested medication?
 Yes No *If No, skip to diagnosis section if applicable.*
9. *If the diagnosis is cataplexy with narcolepsy*, has the patient demonstrated a beneficial response to treatment as defined by a decrease in cataplexy episodes from baseline? Yes No *No further questions*
10. *If the diagnosis is excessive daytime sleepiness*, has the patient demonstrated a beneficial response to treatment as defined by a decrease in daytime sleepiness with narcolepsy from baseline? Yes No *No further questions*

Complete the following section based on the patient's diagnosis, if applicable.

Section A: Excessive Daytime Sleepiness with Narcolepsy

11. Has the patient experienced an inadequate response, intolerance, or contraindication to at least one central nervous system (CNS) stimulant (i.e. amphetamine, dextroamphetamine, methylphenidate)?
If patient is 7 years old to 17 years old, no further questions Yes No
12. Has the patient experienced an inadequate response or intolerance to armodafinil or modafinil?
If Yes, no further questions Yes No
13. Does the patient have a contraindication to armodafinil and modafinil? Yes No

State Step Therapy

1. Is the requested drug being used for an FDA-approved indication or an indication supported in the compendia of current literature (examples: AHFS, Lexicomp, Clinical Pharmacology, Micromedex, current accepted guidelines)? Yes No
2. Does the prescribed quantity fall within the manufacturer's published dosing guidelines or within dosing guidelines found in the compendia of current literature (examples: package insert, AHFS, Lexicomp, Clinical Pharmacology, Micromedex, current accepted guidelines)? Yes No
3. Does the patient reside in Maryland? Yes No *If No, skip to #7*
4. Is the alternate drug (Xyrem) FDA-approved for the medical condition being treated?
 Yes No *If No, no further questions.*
5. Has the prescriber provided proof, documented in the patient's chart notes, indicating that the requested drug was ordered for the patient in the past 180 days? Yes No *If No, skip to #7*
6. Has the prescriber provided proof, documented in the patient chart notes, that in their opinion the requested drug is effective for the patient's condition? Yes No *No further questions*
7. Are any of the following conditions met for the alternate drug (Xyrem)? *If Yes, indicate below and no further questions.*
 - The alternate drug is contraindicated
 - The alternate drug is likely to cause an adverse reaction, physical or mental harm
 - The alternate drug is expected to be ineffective
 - The alternate drug was previously tried or a drug in the same class or with the same action was previously tried and was stopped due to ineffectiveness or an adverse event
 - The alternate drug is not in the patient's best interest
 - The alternate drug was tried while covered by the current or the previous health benefit plan
 - None of the above, *continue to #8*

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8. Is the patient stable or currently receiving a positive therapeutic outcome with the requested drug and a change in the prescription drug is expected to be ineffective or cause harm to the patient? Yes No

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

X _____

Prescriber or Authorized Signature

Date (mm/dd/yy)

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