

## Methadone – Virginia Prior Authorization Request Form

Please complete this <u>entire</u> form and fax it to: 866-940-7328. If you have questions, please call 800-310-6826. This form may contain multiple pages. Please complete all pages to avoid a delay in our decision.

Allow at least 24 hours for review.

| Member Information  |                  |                           |                           | Prescriber Information |               |                      |
|---|------------------|---------------------------|---------------------------|------------------------|---------------|----------------------|
| Member Name:  |                  |                           | Provider Name             | Provider Name:         |               |                      |
| Member ID:  |                  |                           | NPI#:                     | I#: Specialty:         |               |                      |
| Date Of Birth:  |                  |                           | Office Phone:             | Office Phone:          |               |                      |
| Street Address:   |                  |                           | Office Fax:               | Office Fax:            |               |                      |
| City:   | State: ZIP Code: |                           | Office Street Ad          | Office Street Address: |               |                      |
| Phone:  | Aller            | gies:                     | City:                     | State                  | :             | ZIP Code:            |
| Is the requested medi   | cation:   No     | ew or 🗆 Continua          | ition of Therapy? If c    | ontinuation, li        | st start da   | ate:                 |
| Is this patient current   | y hospitaliz     | ed? □ Yes □ N             | o If recently dischar     | ged, list disch        | arge date:    | :                    |
| Is this member pregna   | ant? □ Yes       | □ No If yes, wh           | at is this member's d     | ue date?               |               | <del></del>          |
|   |                  | Medi                      | cation Informatio         | n                      |               |                      |
| Medication:   |                  |                           |                           |                        | Strength:     |                      |
| Directions for use:   |                  |                           |                           | Quantity:              |               |                      |
| Medication Administered   | i: □ Self-Adı    | ministered   Pr           | nysician's Office □ O     | ther:                  |               |                      |
|   |                  |                           | nical Information         |                        |               |                      |
| What is the nationt's   | diagnasia fe     |                           |                           |                        |               |                      |
| What is the patient's diagnosis for the medication being requested? |                  |                           |                           |                        |               |                      |
| ICD-10 Code(s):   |                  |                           |                           |                        |               |                      |
| Are there any supporting  | laboratory o     | or test results relate    | ed to the patient's diagr | nosis? (Please s       | pecify or pro | ovide documentation) |
|   |                  |                           |                           |                        |               |                      |
|   | Pr               | evious Medica             | ation Trials / Cont       | raindication           | ıs            |                      |
| Pleas   |                  |                           | ww.uhcprovider.com f      |                        |               | natives              |
| What medication(s) does   |                  |                           |                           |                        |               | <del></del>          |
| length of trial, and reason for discontinuation of each medication) |                  |                           |                           |                        |               |                      |
|   |                  |                           |                           |                        |               |                      |
|   |                  |                           |                           |                        |               |                      |
|   |                  |                           |                           |                        |               |                      |
| What medication(s) does   | s the patient    | have a <u>contraindic</u> | ation or intolerance to?  | ? (Please specify      | ALL medic     | cation(s) with the   |
| associated contraindication   | n to or specifi  | ic issues resulting in    | intolerance to each med   | ication)               |               | . ,                  |
|   |                  |                           |                           |                        |               |                      |
|   |                  |                           |                           |                        |               |                      |
|   |                  |                           |                           |                        |               |                      |
|   | Addi             | tional information        | n that may be importa     | ant for this rev       | riew          |                      |
|   |                  |                           |                           |                        |               |                      |
|   |                  |                           |                           |                        |               |                      |
|   |                  |                           |                           |                        |               |                      |



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| Patient First n | ame:   | Patient Last name:                        | Patient DOB:                        |  |  |
|-----------------|--|---|-------------------------------------|--|--|
|                 | Cli  | nical and Drug Specific Informat          | ion                                 |  |  |
| □ Yes □ No      | Does the prescriber attest that a treatment plan has been established with the patient and includes <a href="ALL">ALL</a> of the following? <ul> <li>Signed agreement with the patient</li> <li>Established expected outcome and improvement in both pain relief and function or just pain relief as well as limitations (i.e., Function may improve yet pain persist OR pain may never be totally eliminated)</li> <li>Established goals for monitoring progress toward patient-centered functional goals; e.g., walking the dog or walking around the block, returning to part-time work, attending family sports or recreational activities, etc.</li> <li>Goals for pain and function, how opioid therapy will be evaluated for effectiveness and the potential need to discontinue if not effective</li> <li>Emphasize serious adverse effects of opioids (including fatal respiratory depression and opioid use disorder, OR alter the ability to safely operate a vehicle)</li> <li>Emphasize common side effects of opioids (constipation, dry mouth, nausea, vomiting, drowsiness, confusion, tolerance, physical dependence, withdrawal)</li> </ul> <li>Prescriber's Signature:</li> |   |                                     |  |  |
|                 |  | ALL REQUESTS                              |                                     |  |  |
| □ Yes □ No      | Is the patient an infan taper?   | t up to 1 year of age who was discharge   | ed from the hospital on a methadone |  |  |
| □ Yes □ No      | Does the prescriber attest that the patient has intractable pain associated with any of the following? (If yes, check which applies)  □ Active cancer □ Hospice care □ Palliative care (treatment of symptoms associated with life limiting illnesses)   |   |                                     |  |  |
| □ Yes □ No      | Does the patient have any of the following diagnoses? (If yes, check which applies)  □ Chronic severe pain □ Metastatic neoplasia □ Sickle cell  |   |                                     |  |  |
| □ Yes □ No      | Is the prescriber one of the following specialists? (If yes, check which applies)  □ Chronic pain specialist □ Oncologist □ Palliative care □ Sickle cell specialist   |   |                                     |  |  |
| □ Yes □ No      | Does the patient have a contraindication to ALL other preferred and non-preferred long acting opioids? (MedWatch form must be submitted)  If yes, list contraindications:  |   |                                     |  |  |
| □ Yes □ No      | Does the patient have  | a history of, or received treatment for,  | drug dependency or drug abuse?      |  |  |
| □ Yes □ No      | Has the prescriber checked the prescription monitoring program (PMP) website on the date of this request to determine whether the patient is receiving opioid dosages or dangerous combinations (e.g. opioids and benzodiazepines) that put the patient at risk for fatal overdose?  If yes, list date the website was accessed:   |   |                                     |  |  |
|                 | What is the date of the website?   | e patient's last opioid prescription with | in the last 12 months from the PMP  |  |  |



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| Patient First name:         |  | Patient Last name:  | Patient DOB:                         |  |
|-----------------------------|--|---|--------------------------------------|--|
|                             | What is the date of the the PMP website?   | e patient's last benzodiazepine prescrip  | ption within the last 12 months from |  |
| □ Yes □ No □ Not Applicable | Has the prescriber coupenzodiazepines and  | unseled the patient of the risks associa<br>opioids?  | ated with combined use of            |  |
|                             | What is the patient's t  | otal morphine milligram equivalents (N  | MME) from the PMP website?           |  |
| □ Yes □ No                  | (If yes, check which app<br>□ MME/Day of 51 to 90<br>prevention education  | <u>0,</u> the prescriber considered offering a pre  | escription for naloxone and overdose |  |
|                             |  | prescriber considered offering a prescripti<br>, and consulting with a pain specialist  | ion for naloxone, providing overdose |  |
| □ Yes □ No                  | of 10 substances inclu<br>amphetamines, and m  | ne drug screen (UDS) been done at lea<br>uding heroin, prescription opioids, coc<br>etabolites?<br>nt UDS must be submitted)  |                                      |  |
|                             | EXCI   | EEDING 90 MME CUMULATIVE THRES  | HOLD                                 |  |
| □ Yes □ No                  | Does the patient have  | chronic, moderate to severe pain OR s   | severe post-operative pain?          |  |
| □ Yes □ No                  | <ul> <li>That they will I</li> <li>Have reviewed</li> <li>Acknowledge overdose</li> </ul>  | ttest to ALL of the following? be managing the patient's opioid thera I the Virginia board of medicine (BOM) the warnings associated with high dos s medically necessary for this patient | Regulations for Opioid Prescribing   |  |
| □ Yes □ No                  | Has the prescriber prescribed naloxone?  |   |                                      |  |
| □ Yes □ No □ Not Applicable | If the patient is a woman between the ages of 18 to 45 years of age, has the provider counseled the patient on becoming pregnant while receiving opioids, including the risk of Neonatal Opioid Withdrawal Syndrome and offered access to contraceptive services when necessary? |   |                                      |  |
| □ Yes □ No                  | Has the prescriber ordered and reviewed a urine drug screen (UDS) or serum medication level at least every 3 months for the first year of treatment and at least every 6 months thereafter to ensuadherence?   |   |                                      |  |
|                             |  |   |                                      |  |

| Provider Signature:   | Date: |
|-----------------------|-------|
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