Health Professionals’ Services Program  
Phone: 888-802-2843 Fax: 503-961-7142  

Medication Management Form  

(Due within 3 business days from date of prescription)  

Dear Provider,  

As part of a monitoring agreement, this patient must submit a log of all new and continuing prescriptions of medications with addictive potential and/or psychotropic medications including Buprenorphine and Methadone. For a definition of medications with addictive potential and/or psychotropic medications, please consult the HPSP letter of information. Opioid antagonists like Naltrexone and Vivitrol are often prescribed to assist in treatment and recovery. Please discuss with your patient if they are taking an opioid antagonist and add the medication to this form.  

In addition, several over the counter medications may have sedating or stimulating effects. This includes centrally acting antihistamines and decongestants such as diphenhydramine. Participants who have a non-negative test result due to the presence of one of these over the counter medications will be required to provide a letter from their treating physician indicating that the physician is aware that the patient is participating in the Health Professionals’ Services Program, and is also aware that the participant is taking an over the counter medication that has potentially sedating or stimulating effects, and agrees with the participant’s use of the medication. If you approve of your patient using over the counter medications that may have sedating or stimulating effects, please add them to the list below.  

We ask that you complete and sign this form and fax it to HPSP at: 503-961-7142 within three business days. The participant is responsible for submitting a copy of valid prescription(s) within 24 hours to HPSP. Thank you for your attention to this delicate matter. Please call us with any concerns.  

Thank you,  

Health Professionals’ Services Program
Medication Management Form

Name of Participant or Acct#: ___________________________ Participant DOB: ___________________________

Name of Provider (First Middle Last, Credentials): ___________________________

Name of Provider Practice and Specialty: ___________________________

Provider Office Address: ___________________________

Provider Phone: ___________________________

Provider Fax: ___________________________

Medications with Addictive Potential

<table>
<thead>
<tr>
<th>Date of Most Recent Rx</th>
<th>Initial Start Date of Rx</th>
<th>Medication Name (Generic)</th>
<th>Dosage, route, freq (ex: 25 mg PO BID)</th>
<th>#</th>
<th>Condition Prescribed for</th>
<th>Expected duration of treatment</th>
<th>Can patient continue to work while taking this medication?</th>
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Psychotropic Medications and/or sedating or stimulating OTC medications

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<tr>
<th>Date of Most Recent Rx</th>
<th>Initial Start Date of Rx</th>
<th>Medication Name (Generic)</th>
<th>Dosage, route, freq (ex: 25 mg PO BID)</th>
<th>#</th>
<th>Condition Prescribed for</th>
<th>Expected duration of treatment</th>
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If the patient should not work while taking a listed medication, WHEN can patient return to his/her professional duties?

______________________________

Comments:____________________________________________________________________________________

___________________________________________________________________________________________

Please verify:

☐ I have been informed this patient is in recovery for chemical dependency or has a mental health diagnosis. I have discussed opioid antagonists with participant and noted above if participant is taking Naltrexone or Vivitrol and was prescribed any opioid medication.

☐ I have been given the HPSP letter of information regarding prescription of medications with addictive potential and/or psychotropic medications.

Practitioner’s Name (Please Print) ___________________________ Practitioner’s Signature (no stamps, please) ___________________________ Date ____________

This information has been disclosed to you from records whose confidentiality is protected by Federal Law. Federal Regulation (42 CFR, Part 2) prohibits you from making any further disclosure of it without the specific written consent of the person to whom it pertains, or as otherwise permitted by such regulations. A general authorization for the release of medical or other information is NOT sufficient for this purpose. The Federal rules restrict any use of the information to criminally investigate or prosecute the patient.

Rev. 4/6/2022