

Opioid Analgesic Treatment Worksheet (Consolidated)

Aetna Better Health of Louisiana
 Fax: 1-844-699-2889
 For questions only, please call 1-855-242-0802.
www.aetnabetterhealth.com/louisiana/providers/pharmacy

Fee for Service (FFS)
Louisiana Legacy Medicaid
 Fax: 1-866-797-2329
 For questions, please call 1-866-730-4357.
www.lamedicaid.com

Healthy Blue
 Fax: 1-844-864-7865
 For questions, please call 1-844-521-6942
<https://providers.healthyblue.com>

LA Healthcare Connections
 Fax: 1-866-399-0929
 For questions, please call 1-888-929-3790.
www.louisianahealthconnect.com/for-members/pharmacy-services/

AmeriHealth Caritas Louisiana
 Fax: 1-855-452-9131
 For questions, please call 1-800-684-5502.
www.amerihealthcaritasla.com/pharmacy/index.aspx

UnitedHealthcare
 Fax: 1-866-940-7328
 For questions, please call 1-800-310-6826.
www.uhcommunityplan.com/health-professionals/la/pharmacy.html

Please fax the completed form to the appropriate plan using the designated fax number provided above.

Recipient Name:	FFS / MCO ID #:	Recipient DOB:	Medication Allergies:
Resident of long-term care facility: Yes / No If yes, name and phone number:		Recipient Weight (kg):	Recipient Height (ft/in):
Prescriber Name:	Prescriber Specialty:	Medicaid Provider ID # or NPI#:	
Prescriber Address:	Call-Back Phone#:		
Office Fax#:	Office Contact:	EPSDT Support Coordinator (Name/Address): <i>(optional)</i>	

DRUG INFORMATION (one drug per request)

Drug Name / Dosage Form: _____ Strength: _____ Quantity: _____

Requested medication is short-acting / long-acting. (CIRCLE ONE) Directions: _____

Diagnoses for which the opioid is prescribed (include primary and secondary diagnoses applicable to this request, ICD code and description):

Diagnosis: _____ Diagnosis: _____

Date of Diagnosis: _____ Date of Diagnosis: _____

This medication is being used for: _____ acute condition _____ chronic condition (check one only)

Is this medication being used for moderate to severe neuropathic pain or fibromyalgia? _____ Yes _____ No

Is this medication being used for postoperative pain? _____ Yes _____ No If yes, date of surgery _____

This medication is a PREFERRED / NON-PREFERRED Agent. (CIRCLE ONE)

If PREFERRED, CONTINUE to page 2.

If request is for a non-preferred agent, recipient must have had treatment failure with at least two (2) preferred agents.

Previously Tried Preferred Agents*	Reason for Discontinuation

**Refer to the appropriate MCO / FFS website at top of page for a list of preferred agents.*

OR if preferred agents have **not** been previously tried, provide explanation: _____

Does individual require an abuse deterrent agent based upon a history of substance abuse disorder OR individual's family member or household resident has active substance abuse disorder or a history of substance abuse disorder? _____ Yes _____ No

Does individual require Butrans (buprenorphine transdermal patch) or Belbuca (buprenorphine buccal film) due to concern for abuse or dependence with pure opioid agents? _____ Yes _____ No

Is this request for medication prescribed for treatment of pain related to cancer, palliative care, hospice, or end-of-life care?

Yes No

If NO, proceed to next section.

If YES, STOP HERE, sign below and fax form to the appropriate plan above.

Prescriber's signature: _____

(For FFS and Amerihealth Caritas, appropriate diagnosis code must be entered at POS.)

DOES QUANTITY REQUESTED EXCEED THE MAXIMUM QUANTITY LIMIT? YES / NO (CIRCLE ONE)

DOES DAILY MED EXCEED THE MAXIMUM MED ALLOWED PER DAY? YES / NO (CIRCLE ONE)

If answer is YES to either of these questions, continue to next section and complete the form in its entirety.

Request is for: Initiation of therapy Continuation of Therapy If continuation, is dose currently being tapered? Yes No

If no, explain: _____

Recipient's current CUMULATIVE MED PER DAY: _____ (include MED of medication being requested)

Note: The Louisiana Prescription Monitoring Program (PMP) provides the cumulative MED for all of the recipient's controlled medications. Information is current through the previous day (the day before the PMP is accessed).

For **quantity limit override OR MED override**, explain in detail the need for requested quantity/MED:

List **treatments that have been tried or are currently being given** for this condition, both pharmacological and non-pharmacological:

Pharmacological Treatments (both opioid and nonopioid)

Drug / Strength	Directions	Start Date / End Date (or Current)	Reason for Discontinuation (if applicable)

Non-pharmacological Treatments

Treatment	Start Date / End Date (or Current)

PRESCRIBER ATTESTATION

Please indicate YES/True or NO/False for each of the following attestations. Explanation is required for each 'No/False' answer in order for the request to be considered for approval. For short-acting opioids, complete A – G; for long-acting opioids, complete A – L.

	YES (True)	NO (False)	THE PRESCRIBER ATTESTS TO THE FOLLOWING:
	SHORT AND LONG-ACTING OPIOIDS		
			B. The patient has been screened for substance abuse / opioid dependence and documentation is attached . <i>(Not required for recipients in long-term care facility.)</i>
			C. The PMP will be accessed each time a controlled prescription is written for this patient.
			D. A treatment plan which includes current and previous goals of therapy for both pain and function has been developed for this patient.
			E. Criteria for failure of the opioid trial and for stopping or continuing the opioid has been established and explained to the patient.
			F. Benefits and potential harms of opioid use have been discussed with this patient. In addition, if the patient has concurrent comorbidities or is taking medications that could potentially cause drug-drug interactions, an assessment of increased risk for respiratory depression has been completed and discussed with the patient. The risk of combining opioids with other central nervous system depressants, such as benzodiazepines, alcohol, or illicit drugs such as heroin, has also been specifically addressed. The level of risk for opioid abuse/overdose with the dose/duration prescribed to the patient has also been discussed.
			G. An Opioid Treatment Agreement signed by both the patient and prescriber is on file. <i>(Not required for recipients in long-term care facility.)</i>
LONG-ACTING OPIOIDS			H. The patient requires continuous around the clock analgesic therapy for which alternative treatment options have been inadequate or have not been tolerated.
			I. Patient previously utilized at least two weeks of short-acting opioids for this condition. Please enter drug(s), dose, duration and date of trial in <i>Pharmacological Treatment Section</i> on page 1.
			J. Medication has not been prescribed to treat acute pain, mild pain, or pain that is not expected to persist for an extended period of time.
			K. Medication has not been prescribed for use as an as-needed (PRN) analgesic.
			L. Prescribing information for requested product has been thoroughly reviewed by prescriber.
IF NO FOR ANY OF THE ABOVE (A-L), PLEASE EXPLAIN:			

Opioid overdose reversal medications are a covered benefit. Prior authorization is not required for some products. CDC guidelines recommend offering naloxone to patients at increased risk of overdose, defined as: history of overdose or substance use disorder, doses \geq 50 MED /day, or concurrent use with benzodiazepines. **Please refer to the appropriate MCO / FFS website (top of page 1) for a list of preferred agents.**

I certify that the benefits of opioid treatment for this patient outweigh the risks of treatment and that the information provided herein is true and accurate to the best of my knowledge and may be subject to a routine audit requesting the medical information necessary to verify the accuracy of the information provided.

Please note: An approval is not a guarantee of payment. All edits will apply when medication is processed at point-of-sale (POS). Payment on a claim will only be made when the claim is billed correctly and all conditions for payment are met.

Prescriber's Signature: _____ Date: _____

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