LOUISIANA UNIFORM PRESCRIPTION DRUG PRIOR AUTHORIZATION FORM

Submitted to:		Phone:			Fa	Fax:		Date:			
SECTION	I II - PRESCRIBI	ER INFORMATIO	ON								
Last Name, First Name MI:				NPI# or Plan Provider #:			Sp	Specialty:			
Address:				City:						State:	ZIP Code:
Phone:	Phone: Fax:			Office Contact Name:				Contact Phone:			
SECTION	N III - PATIENT	INFORMATION	N.								
Last Name, First Name MI:				DOB:		Phone:			☐ Male ☐ Other		Female Unknown
Address:			•	City:		1				State:	ZIP Code:
Plan Nam	e (if different fro	om Section I):	Membe	er or Medic	caid ID #:	Plan Provi	der ID:				1
Patient is Patient is Patient is	being discharge being discharge a long-term care	pital inpatient get od from a psychia od from a residen e resident? or contact inforn	tric facility tial substa _ Yes	/? ince use fa No l	icility? f yes, nam	Yes Yes	No No	Date Date	of Disch of Disch	narge: narge:	
SECTION	I IV - PRESCRIP	TION DRUG IN	FORMAT	ION							
Requeste	d Drug Name:										
Strength:	trength: Dosage Form: Route of Admin: Quantity: Da			ys' Supply: Dosage Interval/Directions for Us			ns for Use:	se: Expected Therapy Duration/Start Date:			
To the be	st of your knowl	edge this medica	tion is:								
For Provid	der Administere	d Drugs only:		Contin	uation of t	:herapy/Re	authoriz	ation re	quest		
HCPCS/CF	PT-4 Code:		_NDC#:_			Dose Per	Adminis	tration:			
Other Cod	des:										
Will patie	nt receive the di	rug in the physici	an's office	?Yes	No						
	– If r	no, list name and	NPI of ser	vicing pro	vider/facil	ity:					

SECTI	ON V - PA	ATIENT C	LINICAL INFO	ORMATION						
Primary diagnosis relevant to this request: ICD-10 Diagnosis Code: Date										
	dary diagr	ICD-10 Diagnosis Code: Date Diagnosed:								
For pain-related diagnoses, pain is: Acute Chronic										
For po	stoperativ	/e pain-rel	ated diagnoses	s: Date of Surgery						
Pertir	nent labor	atory valu	es and dates (a	attach or list below):						
Date Name of Test Value										
SECTI	ON VI - T	HIS SEC	TION FOR OP	IOID MEDICATIONS ONLY						
					es No (If yes, provide justification below.)					
	nulative da		Stea exceed th	ie max quantity mint anowed:	esNo (ii yes, provide justification below.)					
		-	MF exceed the	e daily max MMF allowed? Ye	sNo (If yes, provide justification below.)					
	.s carriara	ive daily iv	TIVIL CACCCO CIT	<u></u>	==					
	YES	NO		THE DESCRIPTION AT						
DS	(True)	(False)	THE PRESCRIBER ATTESTS TO THE FOLLOWING:							
Ιō			A Complete assessment for pain and function was performed for this patient.							
0.0			B The patient has been screened for substance abuse / opioid dependence. (Not required for recipients in long-							
(True) (False) A A complete assessment for pain and function was performed for this patient. B The patient has been screened for substance abuse / opioid dependence. (Not required for recipien term care facility.) 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 had developed for this patient. E Criteria for failure of the opioid trial and for stopping or continuing the opioid has been established a explained to the patient. F Benefits and potential harms of opioid use have been discussed with this patient. G An Opioid Treatment Agreement signed by both the patient and prescriber is on file. (Not required for stopping or continuing the opioid has been established and potential harms of opioid use have been discussed with 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.										
ND I		E Criteria for failure of the opioid trial and for stopping or continuing the opioid has been established and explained to the patient.								
₹				d potential harms of opioid use have be	en discussed with this patient.					
l g										
S		G An Opioid Treatment Agreement signed by both the patient and prescriber is on file. (Not required for recipients in long-term care facility.)								
					nalgesic therapy for which alternative treatment options					
S			have been in	adequate or have not been tolerated.						
			I. Patient prev	viously utilized at least two weeks of sho	ort-acting opioids for this condition. Please enter drug(s),					
OPI			dose, duration and date of trial in pharmacologic/non-pharmacologic treatment section below.							
LING			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.							
ĄĊ			K Medication has not been prescribed for use as an as-needed (PRN) analgesic.							
LONG-ACTING OPIOIDS			L. Prescribing information for requested product has been thoroughly reviewed by prescriber.							
IF NO	O FOR ANY	OF THE AB	OVE (A-L), PLEAS	SE EXPLAIN:						

SECTION VII - PHARMACOLOGIC & NON-PHARMACOLOGIC TREATMENT(S) USED FOR THIS DIAGNOSIS (BOTH PREVIOUS & CURRENT):

Drug name	Strength Frequency		Dates Started and Stopped or Approximate Duration	Describe Response, Reason					
			h	11.1.1.1					
Drug Allergies:			Height (if applicable):	Weight (if applicable):					
Is there clinical evidence or patient history that suggests the use of the plan's pre-requisite medication(s), e.g. step medications, vill be ineffective or cause an adverse reaction to the patient?YesNo (If yes, please explain in Section VIII below.) SECTION VIII - JUSTIFICATION (SEE INSTRUCTIONS)									
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By signing this request, the prescriber attests	that the info	rmation provi	ided herein is true and accurate	a to the best of					
by signing this request, the prescriber attests iis/her knowledge. Also, by signing and subm Attestation' section of the criteria specific to	itting this red	quest form, th	ne prescriber attests to stateme						
Signature of Prescriber:			Date:	·					