## Immediate-Release Opioid Prior Authorization Form



Please Fax Form To: 866-930-0019 Date Of Request: \_\_\_\_\_

Patient Information						
Member Name:				CareSource ID:		
Member DOB:						
Pharmacy: Pharmacy Phone #:						
Prescriber Information						
Name:			NPI/DEA:		Specialty:	
Address:						
Office Contact:		Phone:			Fax:	
Diagnosis & Required Information						
Diagnosis Code (ICD-10):						
Prescriber attests to reviewing state prescription drug monitoring program (PDMP) prior to writing prescription. Date:						
Prescriber attests benefits and risks of opioid therapy have been discussed with patient.						
Prescriber attests to a documented patient-specific treatment plan (e.g., assessment of pain and function scores, a baseline urine drug test, plans for random urine drug screens, opioid contract, etc.)						
Prescriber attests to periodic assessment of patient's outcomes (e.g., adherence, progress notes documenting pain and function scores, random urine drug screens, no serious adverse outcomes) to ensure that continued therapy outweighs risk to patient safety.						
Prescriber attests to reassessment of patient's addiction risk or mental health concerns (e.g., using Screening, Brief Intervention, and Referral to Treatment [SBIRT] tools), including referral to an addiction medicine specialist when appropriate.						
If the patient is taking a benzodiazepine, prescriber affirms to assessment to ensure benefit outweighs the risk of benzodiazepine use along with the opioid analgesic.						
If patient's cumulative dose of opioids is above 80 MED per day, prescriber must be a pain specialist or must attest to consulting a pain specialist. If pain specialist is unavailable, provide documentation supporting so and rationale for higher dose.						
Prescriber attests member has tried and failed at least two preferred non-opioid analgesics (NSAIDs, APAP, anticonvulsants, antidepressants) at maximally tolerated doses unless all contraindicated. Please list drugs that have been tried and/or explanation of contraindication.						
Medication Name Date <u>Trial</u> Started Length			<u>R</u>	Reason For Discontinuation/ Contraindication		
Immediate-Release Opioid Requested						
Drug Name:				Strength:		
Quantity: SIG: Dosag				iorm:		
If member is currently treated on this medication, please list start date:						
Which limits you are requesting to exceed? (Circle all that apply)						
>14 Day Supply Within 45 > 7 Day Supply for this Fill Days				>90 Days of Therapy	> 60 MED Per Script (MED	= Morphine Equivalent Dose)
Reason for Request:						
Physician Signature:					Date:	IN-P- 0379 First Issue XX/XX/2018 OMPP CEECEE