



Fax completed prior authorization request form to 844-802-1412 or submit Electronic Prior Authorization through CoverMyMeds® or SureScripts.

Aetna Better Health®

All requested data must be provided. **Incomplete forms or forms without the chart notes will be returned**

Pharmacy Coverage Guidelines are available at <https://www.aetnabetterhealth.com/illinois-medicaid>

Opioids

Pharmacy Prior Authorization Request Form

Do not copy for future use. Forms are updated frequently.

REQUIRED: Office notes, labs and medical testing relevant to request showing medical justification are required to support diagnosis

Member Information									
Member Name (first & last):				Date of Birth:		Gender:		Height:	
						<input type="checkbox"/> Male	<input type="checkbox"/> Female		
Member ID:				City:		State:		Weight:	
Prescribing Provider Information									
Provider Name (first & last):				Specialty:		NPI#		DEA#	
Office Address:				City:		State:		Zip Code:	
Office Contact:				Office Phone			Office Fax:		
Dispensing Pharmacy Information									
Pharmacy Name:				Pharmacy Phone:			Pharmacy Fax:		
Requested Medication Information									
Preferred Short Acting Agents:		<input type="checkbox"/> Hydromorphone		<input type="checkbox"/> ascomp-codeine		<input type="checkbox"/> codeine sulfate		<input type="checkbox"/> morphine sulfate IR	
		<input type="checkbox"/> oxycodone		<input type="checkbox"/> Endocet		<input type="checkbox"/> tramadol		<input type="checkbox"/> Lorcet	
		<input type="checkbox"/> APAP-codeine		<input type="checkbox"/> hydrocodone-APAP					
Preferred Long Acting Agents:		<input type="checkbox"/> Morphine Sulfate ER 15mg							
Non-Preferred Short Acting Agent:		Specify drug:							
Non-Preferred Long Acting Agent:		Specify drug:							
Are there any contraindications to the preferred medications? (if yes, please specify):						<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> New request	<input type="checkbox"/> Continuation of therapy request
Directions for Use:				Strength:			Dosage Form:		
				Quantity:		Day Supply:		Duration of Therapy/Use:	
Medication request is NOT for an FDA- approved, or compendia-supported diagnosis (circle one): Yes No				Diagnosis:			ICD-10 Code:		
What medication(s) have been tried and failed for this diagnosis? Please specify:									
Turn-Around Time for Review									
<input type="checkbox"/> Standard – (24 hours)		<input type="checkbox"/> Urgent – If waiting 24 hours for standard decision could seriously harm life, health, or ability to regain maximum function, you can ask for an expedited decision. Signature: _____							
Clinical Information									
Pain is due to ONE of the following:		<input type="checkbox"/> Active Cancer		<input type="checkbox"/> Sickle Cell		<input type="checkbox"/> Palliative/End of life		<input type="checkbox"/> Hospice	
		<input type="checkbox"/> N/A							
Will member be on both opioid AND benzodiazepine at same		<input type="checkbox"/> Yes <input type="checkbox"/> No		Will Naloxone be provided/offered?			<input type="checkbox"/> Yes <input type="checkbox"/> No		<input type="checkbox"/> N/A

time?						
Is request for opioid naïve member?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Is member opioid tolerant?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Was non-pharmacologic therapy tried PRIOR to prescribing opioids (PT, exercise, CBT OR weight loss)?				<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Was non-opioid therapy tried PRIOR to prescribing opioids? (topical diclofenac NSAIDs, TCAs, and SNRIs OR anticonvulsants)				<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Signed treatment plan addresses the following (check that apply):	<input type="checkbox"/> Realistic goals for pain AND function	<input type="checkbox"/> When treatment will be stopped	<input type="checkbox"/> Consequences of lost medication	<input type="checkbox"/> Consequences of obtaining controlled substances from other prescribers		<input type="checkbox"/> Member using ONE pharmacy
Was member advised of harm AND benefits before treatment AND periodically during treatment (increased risks of respiratory depression, combination use with BNZ, risks to others in household, cognitive limitations AND side effects)?					<input type="checkbox"/> Yes	<input type="checkbox"/> No
Will treatment be prescribed at lowest effective dose?					<input type="checkbox"/> Yes	<input type="checkbox"/> No
Will treatment be reviewed within 1-4 weeks of starting opioid therapy for CHRONIC pain AND with any DOSE-ESCALATION AND RE-EVALUATED every 3 months?					<input type="checkbox"/> Yes	<input type="checkbox"/> No
Was there a review of the state's PMP Drug Monitoring Program for controlled substances?					<input type="checkbox"/> Yes	<input type="checkbox"/> No
Was UDS reviewed prior to starting treatment?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Were results of UDS consistent with prescribed controlled substances?		<input type="checkbox"/> Yes	<input type="checkbox"/> No
Is there evidence of substance use disorder?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was evidence-based treatment like MAT arranged?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Is request for female of reproductive age?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was counseling provided about opioid use during pregnancy AND neonatal abstinence syndrome?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Additional Clinical Information						
<input type="checkbox"/> Long Acting Opioids						
Will member exceed 90 MME per day limit?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was documentation submitted to support exceeding recommended limit?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Was pain specialist consulted?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A	Is request for chronic pain?		<input type="checkbox"/> Yes <input type="checkbox"/> No
Was treatment started with an IR opioid for at least 2 weeks prior to requesting ER/LA opioid?					<input type="checkbox"/> Yes	<input type="checkbox"/> No
Is request for buprenorphine weekly patch ?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Is there need for opioid with lower risk for abuse AND a noted concern that member OR member's household is at risk for abuse AND diversion?		<input type="checkbox"/> Yes	<input type="checkbox"/> No
Is request for non-preferred agent ?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there inadequate response OR intolerance to MSER for at least 2 weeks?		<input type="checkbox"/> Yes	<input type="checkbox"/> No
Is request for abuse-deterrent product ?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there trial AND failure with buprenorphine patch for at least 2 weeks?		<input type="checkbox"/> Yes	<input type="checkbox"/> No
			Is there a NEED for abuse deterrent agent AND a noted concern that member OR household is at risk for abuse AND diversion?		<input type="checkbox"/> Yes	<input type="checkbox"/> No
Is request for methadone ?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Is female member pregnant?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
<input type="checkbox"/> Short Acting Opioids						
Will member exceed 90 MME per day limit?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there documentation to support medical necessity of exceeding recommended MME, or day supply limit?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Is request for non-preferred short-acting agent?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was there inadequate response OR intolerance to 2 preferred short-acting opioids?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	
Was documentation submitted supporting continued use of a SHORT ACTING AGENT beyond 30 days AND when used in combination with LONG-ACTING agent?				<input type="checkbox"/> Yes	<input type="checkbox"/> No	
<input type="checkbox"/> Acute Pain Pediatric Members less than 18 years of Age						
Is request for ACUTE pain (post-dental procedure)?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was a pain assessment completed?		<input type="checkbox"/> Yes	<input type="checkbox"/> No
Were member AND their parent(s)/guardian(s) screened for previous AND current opioid use?					<input type="checkbox"/> Yes	<input type="checkbox"/> No

Has provider checked state's PMP Drug Monitoring Program for controlled substances?					<input type="checkbox"/> Yes	<input type="checkbox"/> No
Was concomitant use with BNZ appropriately addressed, if present?				<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Was COMBO therapy of APAP and NSAIDs tried AND failed OR there is C/I present for use of both?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Will OPIOID THERAPY be used in COMBO with APAP and NSAIDs unless there is C/I present for use with both?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Is medication prescribed codeine or tramadol with age being <12 years? (NOTE: use of these medications is C/I in ages younger than 12 AND not recommended for ages 12-17.)				<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Will prescription be limited to 8 – 12 tablets?					<input type="checkbox"/> Yes	<input type="checkbox"/> No
Will IR opioids be prescribed, limited to lowest effective dose AND no quantity greater than expected pain duration that is severe enough to require opioids will be given (NOTE: 3 days or fewer is recommended by CDC)?					<input type="checkbox"/> Yes	<input type="checkbox"/> No
<input type="checkbox"/> RENEWAL ONLY						
Was there sustained improvement in Pain OR Function?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was tapering plan initiated to D/C treatment of current medication?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Was UDS performed in past year?	<input type="checkbox"/> Yes	<input type="checkbox"/> No				
State's PMP was reviewed AND verified (check that apply):	<input type="checkbox"/> Prescriptions from other providers		<input type="checkbox"/> Benzodiazepine use	<input type="checkbox"/> ER / LA use for acute pain	<input type="checkbox"/> UDS is consistent with prescribed controlled substances	
Is dose ≥50 MME per day?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Did provider offer Naloxone to member?		<input type="checkbox"/> Yes	<input type="checkbox"/> No
						<input type="checkbox"/> N/A
Is dose ≥90 MME per day?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Did provider refer member to Pain Specialist?		<input type="checkbox"/> Yes	<input type="checkbox"/> No
						<input type="checkbox"/> N/A
Is there continued concomitant use of opioid with BNZ?	<input type="checkbox"/> Yes	<input type="checkbox"/> No	Was member counseled on FDA BBW on concomitant use AND provider to prescribe at LOWEST effective dose AND duration?		<input type="checkbox"/> Yes	<input type="checkbox"/> No
						<input type="checkbox"/> N/A
Additional information the prescribing provider feels is important to this review. Please specify below or submit medical records						

Signature affirms that information given on this form is true and accurate and reflects office notes.

Prescribing Provider's Signature: _____ **Date:** _____

Please note: Incomplete forms or forms without the chart notes will be returned

Office notes, labs, and medical testing relevant to the request that show medical justification are required.

Standard turnaround time is 24 hours. You can call 866-329-4701 to check the status of a request.