

Updated: 09/2024 DMMA Approved: 09/2024

Request for Prior Authorization for Carisoprodol Website Form – <u>www.highmarkhealthoptions.com</u> Submit request via: Fax - 1-855-476-4158

All requests for Carisoprodol require a Prior Authorization and will be screened for medical necessity and appropriateness using the criteria listed below.

## Carisoprodol Prior Authorization Criteria:

Coverage may be provided with a <u>diagnosis</u> of <u>acute</u> musculoskeletal pain and the following criteria is met:

- o Member is age appropriate according to FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature
- o Must have tried and failed two preferred skeletal muscle relaxant medications
- The requested dose and frequency is in accordance with FDA-approved labeling, nationally recognized compendia, and/or evidence-based practice guidelines and will not exceed a max daily dosing of four times per day
- o Carisoprodol should only be used for a maximum of 2 to 3 weeks due to lack of evidence of effectiveness with prolonged use
- o The member will not use carisoprodol in combination with a benzodiazepine
- The member has no contraindications to the medication including acute intermittent porphyria.

## • **Initial Duration of Approval:** up to 3 weeks.

Coverage may be provided for any non-FDA labeled indication if it is determined that the use is a medically accepted indication supported by nationally recognized pharmacy compendia or peer-reviewed medical literature for treatment of the diagnosis(es) for which it is prescribed. These requests will be reviewed on a case by case basis to determine medical necessity.

Drugs are authorized in generic form unless the branded product is on the preferred drug list or the prescriber has indicated in writing that the branded product is medically necessary. If only the branded product is on the preferred drug list, the generic form will be considered nonpreferred and shall not require the prescriber to indicate in writing that the branded product is medically necessary.



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## CARISOPRODOL PRIOR AUTHORIZATION FORM

Please complete and fax all requested information below including any progress notes, laboratory test results, or chart

	able to Highmark Health Options you may call to speak to a Pharn			
	<b>NE</b> : (844) 325-6251 Monday thro		ve.	
THO	PROVIDER INFORM			
Requesting Physician:		NPI:		
Physician Specialty:		Office Contact:	Office Contact:	
Office Address:		Office Phone:		
		Office Fax:		
	MEMBER INFORM	ATION		
Member Name:				
Member ID:		DOB:		
	REQUESTED DRUG INF			
Medication:		Strength:	<u>U</u>	
Frequency:		Duration:		
Is the member currently receiving requested medication? \( \subseteq \text{Yes} \)		Date Medication Init	Date Medication Initiated:	
No				
Is this medication being used for a		for which the medication m	nay be necessary for the	
life of the patient? Yes	No MEDICAL HIGH	ODY		
D' '	MEDICAL HIST			
Diagnosis:		ICD-10:	<del></del>	
Has the member tried and failed to	vo masfamad altalatal mayaala mala	xants? Yes No		
	*			
Will the member be using carisopa	rodol in combination with a benz	odiazepine? LYes LN	lo	
Does the member have any contra	indications to carisoprodol include	ling acute intermittent porp	ohyria? Yes No	
	<b>CURRENT or PREVIOU</b>	STHERAPY		
Drug Name	Strength/ Frequency	Dates of Therapy	Status (Discontinued & Why / Current)	
		A P POP GOVERNMANTA		
SUPPORTING INFORMATI	ON or CLINICAL RATIONA	LE FOR CONTINUATION	ON OF TREATMENT	
Duocavihina Dh	ysician Signature		Date	
Trescribing Fil	ysician signature		Date	



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