

Prior Authorization Soma® (Carisoprodol)/Soma® Compound

(Maximum of 30 Days Approval (120 Tablets)/365 Days)

Note: Form must be completed in full. An incomplete form may be returned.																												
Recipient	t's Me	edica	id ID	#	1		1				Date	e of E	<u>3</u> irth	(<u>MM</u>	<u>/DD/</u>	<u> </u>	<u></u>			1	1							
													/			/												
Recipient	t's Fu	II Na	me		1		ı						_						1	ı		1	1		ı	1		
Prescribe	er's F	ull Na	ame																									
Prescribe	er Lice	ense	# (N	IE, C	S, Al	RNP,	PA)									,			1			1				•		
Prescriber Phone Number										Pres	scribe	er Fa	x Nui	mber														
						-																		-				
Pharmac	v Nar	ne					!										<u> </u>											
Pharmac	y Med	dicai	d Pro	vide	r#								1	1			1											
Pharmac	y Pho	one N	lumb	er													Pha	rmac	y Fa	x Nur	nber							
						-																		-				
	Son	n a [®]	(Car	icon	rodo	۱۱)												-	ı		<u> </u>					ı		
			•			,ı,		_					Direc	tion							-	Oua	ntity	/30 [)21/6			
Soma [®] Compound Directions Quantity/30 Days																												
Please in	dicat	e pat	tient	diagı	nosis	: (Mu	st pro	ovide	sup	porti	ng d	ocun	nenta	ation)													
Please list (2) skeletal muscle relaxants the patient received in the past 365 days? (Please provide supporting clinical documentation indicating																												
therapeu	tic ou	tcom	ne of	trials	and	failu	res)																					
Drug Name: Dates of Use:											_																	
Reason for Discontinuing:																												
Drug Name: Dates of Use:										_																		
Reason f	or Dis	scont	tinuir	ng: _																								
Prescriber's Signature: Date:									_																			
REQUIRED FOR REVIEW: Copies of medical records (i.e., diagnostic evaluations and recent chart notes), a copy of the original prescription, and the most recent copies of related labs. Supporting documentation includes chart notes, progress notes, and discharge summaries. The provider must retain copies of all documentation for five years.																												

Fax Information to:



Pharmacy Provider Services Fax: 855-825-2717 Phone: 1-800-617-5727

FLORIDA MEDICAID

PROTOCOL

Soma® (Carisoprodol/Soma® Compound)





Note: Form must be completed in full. An incomplete form may be returned.

Approval Indications

- Beneficiary must have failed at least two other skeletal muscle relaxants in the past 365 days.
- Approval limited to a one month supply (120 tablets) during a 365 day period.

Approval Period

Maximum of 30 days approval (120 tablets) / 365 days

Tapering Guidelines (Sample)

Short Taper	Long Taper									
Reduce Carisoprodol over 4 days:	Reduce Carisoprodol over 9 days:									
350mg TID X 1 day, then	350mg TID X 3 days, then									
350mg BID X 2 days, then	350mg BID X 3 days, then									
• 350mg QD X 1 day	• 350mg QD X 3 days									