

#### FLORIDA MEDICAID

### Prior Authorization Soma® (Carisoprodol)/Soma® Compound

Note: Maximum of 30 Days Approval (120 Tablets)/365 Days Note: Form must be completed in full. An incomplete form may be returned.

Beneficiary's Medicaid ID#										Date of Birth (MM/DD/YYYY)																				
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Ben	eficia	ary's	Full	Nan	ne					_				_				_												
Prescriber's Full Name																														
Prescriber's NPI															I															
Prescriber Phone Number										_							Pre	scrib	er Fa	ax Number										
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Pharmacy Name																														
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Pharmacy Phone Number  Pharmacy Fax Number													ı																	
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☐ Soma® (Carisoprodol)																										-				
☐ Soma® Compound										Directions									Quantity/30 Days											
Please indicate patient diagnosis: (Must provide supporting documentation.)																														
												_																		_
Plea	ase lis	st (2)	prefe	erred	skel	etal r	musc	le re	laxa	nts th	e pa	tient	rece	ived	l in	the	pas	st 365	i day	s. <i>(Pl</i>	ease	prov	ide s	шрр	orting	clini	cal d	ocun	nenta	tion
indi	cating	ther	ареи	ıtic o	utcor	ne oi	f trial	s and	d fail	ures.,	)																			
Drug Name:											Dates of Use:																			
Rea	son f	or Di	scon	tinuir	ng:																									_
Drug Name:Reason for Discontinuing:											Dates of Use:																			
Rea	son f	or Dis	scon	tinuir	ng: _																									_
Prescriber's Signature:																														
REQUIRED FOR REVIEW: All copies of medical records (e.g., diagnostic evaluations and recent chart notes), and the recopies of related labs. The provider must retain copies of all documentation for five years.												ne m	ost r	ecen	ıt															

Fax this form to 1-866-940-7328

Pharmacy PA Call Center: 1-800-310-6826

02.01.2025

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### FLORIDA MEDICAID

# PROTOCOL

## 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.

### **Approval Indications:**

- Beneficiary must have failed at least two preferred 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