HEALTHY U MEDICAID

PRIOR AUTHORIZATION REQUEST FORM SUNOSI

For authorization, please answer each question and fax this form PLUS chart notes back to the Healthy U Prior Authorization Department at 385-425-4052.

Failure to submit clinical documentation to support this request will result in a dismissal of the request.

If you have prior authorization questions, please call for assistance: 385-425-5094

Disclaimer: Prior Authorization request forms are subject to change in accordance with Federal and State notice requirements.

Date:	Member Name:	ID#:
Bate.		
DOB:	Gender:	Physician:
DOD.	Gender.	
Office Phone:	Office Fax:	Office Contact:
office filone.	Office Fax:	office contact.

Height/Weight:

Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.

Product being requested:
Sunosi[®] (solfiamfetol)

Dosing/Frequency:_

If the request is for reauthorization, proceed to reauthorization section.					
	Questions	Yes	No	Comments/Notes	
EXCESSIVE SOMNOLENCE ASSOCIATED WITH NARCOLEPSY					
1. Is the member 18 years	of age or older?				
2. Does the member have	a baseline ESS score of 15 or higher?			Please provide documentation	
3. Does the member have polysomnography and N	a diagnosis of narcolepsy confirmed by /SLT?			Please provide documentation	
 Is Sunosi[®] prescribed by disorder specialist or ne 	, or in consultation with, a sleep urologist?				
following categories for • Central nervous syst	t least one agent from each of the at least 3 months each: em stimulant (e.g. methylphenidate) ting agent (e.g. modafinil)			Please provide documentation	
6. Is the member's blood p	ressure adequately controlled?			Please provide documentation	
7. Will the member be mo exacerbations?	nitored for psychologic disorders or				
EXCESSIVE SOMNOLENCE ASSOCIATED WITH SLEEP APNEA					
1. Is the member 18 years	of age or older?				
2. Does the member have	a baseline ESS score of 15 or higher?			Please provide documentation	
	a diagnosis of obstructive sleep apnea order specialist with either CST?			Please provide documentation	

4. Is Sunosi [®] prescribed by, or in consultation with, a sleep			
disorder specialist or pulmonologist?			
5. Is the member being treated with non-pharmacologic primary treatment modalities (CPAP or similar)?			Please provide documentation
6. Is the member at least 90% compliant on non-pharmacologic			
primary treatment modalities with at least 5 hours of use per			
night for at least 3 months prior to initiation of Sunosi [®] ?			
7. Will the member continue to use CPAP therapy for at least 6			
hours per night with at least 90% compliance during Sunosi [®] therapy?			
8. Has the member tried modafinil or armodafinil for at least 3			Please provide documentation
months while using CPAP?			
9. Is the member's blood pressure adequately controlled?			Please provide documentation
10. Will the member be monitored for psychologic disorders or exacerbations?			
REAUTHORIZATIO	N		
1. Is the request for reauthorization of therapy?			
 Does documentation show the member had an improvement in 			Please provide documentation
ESS score from baseline?			riease provide documentation
 At least 5 point improvement for initial renewal 			
 Maintenance of ESS score improvement for ongoing 			
renewals			
3. For OSA, has the member continued to use non-pharmacologic	Π		Please provide documentation
primary treatment modalities with at least 90% compliance for		_	
at least 6 hours per night?			
What medications and/or treatment modalities have been tried in	the pas	st for this	condition? Please document
name of treatment, reason for failure, treatment dates, etc.	•		
Additional information:			
Physician Signature:			

** Failure to submit clinical documentation to support this request will result in a dismissal of the request.**

Policy: PHARM-HU-107 Origination Date: 01/01/2022 Reviewed/Revised Date: 01/18/2023 Next Review Date: 01/18/2024 Current Effective Date: 02/01/2023

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