

## **Beneficiary Information**

1. Beneficiary Last Name: 2. First Name:							
3. Beneficiary ID #:							
Prescriber Information							
6. Prescriber Name:				NPI #:			
Mailing address:			City:		State:	ZIP:	
7. Requester Contact Inf	ormation:						
Name:		Phone #:			Fax #:		
Drug Information							
8. Drug Name:		9. Strength:		1	LO. Quantity P	er 30 Days:	
11. Length of Therapy :	Initial Authorization:	up to 30 days	60 days	90 days	i		
	Reauthorization:	_up to 30 days	_60 days	_90 days _	120 days	180 days	
Clinical Information							
1. Is the beneficiary 7 years of age or older? Yes No							
2. Does the beneficiary have any current use of alcohol or sedative hypnotics? Yes No							
3. Does the beneficiary have succinic semialdehyde dehydrogenase deficiency? Yes No							
4. Has the beneficiary been evaluated for history of drug abuse? Yes No							
5. Will the prescriber monitor the beneficiary for signs of misuse or abuse of sodium oxybate (a.k.a. gamma-hydroxybutyrate							
[GHB]) including, but not limited to, the following: Use of increasingly large doses, increased frequency of use, drug-seeking behavior, feigned cataplexy, etc.? Yes No							
6. Does the beneficiary have a diagnosis of cataplexy associated with narcolepsy? Yes No							
7. Does the beneficiary have a diagnosis of excessive daytime sleepiness due to narcolepsy with daily periods of irrepressible need							
to sleep or daytime lapses into sleep occurring for <a>2</a> a months? Yes No							
8. Does the beneficiary have hypersomnolence secondary to another sleep disorder, neurologic disorder, medical condition, or by							
medicine, or substance use has been ruled out? YesNo							
For continuation of therapy, please answer questions 1-10							
9. For a diagnosis of Excessive Daytime Sleepiness, has the beneficiary responded to therapy with a reduction in excessive							
daytime sleepiness from pre-treatment baseline measured by a validated scale (e.g., Epworth Sleepiness Scale, Stanford							
Sleepiness Scale, Karolinska Sleepiness Scale, Cleveland Adolescent Sleepiness Questionnaire, or a Visual Analog Scale)? Yes No							
10. For a diagnosis of cataplexy, has the beneficiary had a reduced frequency of cataplexy attacks from pre-treatment baseline?							
YesNo							

Date: \_\_\_\_\_

Signature of Prescriber: \_\_\_\_\_\_ \*Prescriber signature mandatory

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.