

**Beneficiary Information**

1. Beneficiary Last Name: _____	2. First Name: _____
3. Beneficiary ID #: _____	4. Beneficiary Date of Birth: _____
5. Beneficiary Gender: _____	

**Prescriber Information**

6. Prescriber Name: _____	NPI #: _____
Mailing address: _____	City: _____ State: _____ ZIP: _____
7. Requester Contact Information: _____	
Name: _____	Phone #: _____ Fax #: _____

**Drug Information**

8. Drug Name: _____	9. Strength: _____	10. Quantity Per 30 Days: _____
11. Length of Therapy: Initial Request: <input type="checkbox"/> up to 30 days <input type="checkbox"/> 60 days <input type="checkbox"/> 90 days		
Reauthorization Request: <input type="checkbox"/> up to 30 days <input type="checkbox"/> 60 days <input type="checkbox"/> 90 days <input type="checkbox"/> 120 days <input type="checkbox"/> 180 days		

**Clinical Information**

**Initial Request for Hetlioz – Non-24 Sleep-Wake Disorder: (answer questions 1-5)**

1. Is the beneficiary at least 18 years old or older? Yes\_\_\_ No\_\_\_

2. Does the beneficiary have a documented diagnosis of Non-24 sleep-wake disorder? Yes\_\_\_ No\_\_\_

3. The diagnosis of Non-24 sleep-wake disorder is confirmed by meeting ONE of the following conditions:  
 Assessment of at least one physiologic circadian phase marker (e.g., measurement of urinary melatonin levels, dim light melatonin onset [as measured in blood or saliva], assessment of core body temperature.)  
 If the assessment of at least one physiologic circadian phase marker cannot be done, the diagnosis must be confirmed by actigraphy performed for >= 1 week plus evaluation of sleep logs recorded for >= 1 month.

4. Has the beneficiary had an insufficient response or intolerance to at least two (2) other medications, over-the-counter or prescription, for sleep? Yes\_\_\_ No\_\_\_

5. Is this medication being prescribed by, or is the physician consulting with, a physician who specialized in the treatment of sleep disorders? Yes\_\_\_ No\_\_\_

**Initial Request for Hetlioz – Nighttime Sleep Disturbances in Smith-Magenis Syndrome (SMS): (answer questions 6-10)**

6. Is the beneficiary at least 16 years of age or older? Yes\_\_\_ No\_\_\_

7. Does the beneficiary have a documented diagnosis of Nighttime sleep disturbances in Smith-Magenis Syndrome (SMS)? Yes\_\_\_ No\_\_\_

8. Has the beneficiary had an insufficient response or intolerance to at least two (2) other medications, over-the-counter or prescription, for sleep? Yes\_\_\_ No\_\_\_

10. Is this medication being prescribed by, or is the physician consulting with, a physician who specialized in the treatment of sleep disorders? Yes\_\_\_ No\_\_\_

**Initial Request for Hetlioz LQ – Nighttime Sleep Disturbances in Smith-Magenis Syndrome (SMS): (answer questions 11-14)**

11. Is the beneficiary between 3 years and 15 years of age? Yes\_\_\_ No\_\_\_

12. Does the beneficiary have a documented diagnosis of Nighttime sleep disturbances in Smith-Magenis Syndrome (SMS)? Yes\_\_\_ No\_\_\_

13. Has the beneficiary had an insufficient response or intolerance to at least two (2) other medications, over-the-counter or prescription, for sleep? Yes\_\_\_ No\_\_\_

14. Is this medication being prescribed by, or is the physician consulting with, a physician who specialized in the treatment of sleep disorders? Yes\_\_\_ No\_\_\_

**Re-authorization for Treatment: (answer questions 15 & 16 below)**

15. Has the beneficiary used Hetlioz or Hetlioz LQ continuously without gaps in treatment for the initial approval period of three (3) months? Yes\_\_\_ No\_\_\_

16. As the provider, have you included an objective evaluation of the beneficiary's sleep quality, including documentation of an improvement in overall sleep quality while taking Hetlioz/Hetlioz LQ? Yes\_\_\_ No\_\_\_

**\*\*Documentation of the beneficiary's overall sleep quality improvement must accompany this reauthorization for Hetlioz and Hetlioz LQ.\*\***

Signature of Prescriber: \_\_\_\_\_ Date: \_\_\_\_\_

**\*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.

**Fax this form to: 1-877-234-4274, or call Pharmacy Prior Authorization: 1-866-885-1406**