

Beneficiary Full Name: \_\_\_\_\_

Sponsor's SSN: \_\_\_\_\_ - \_\_\_\_\_ - \_\_\_\_\_

Date of Birth: \_\_\_\_\_

Beneficiary State of Residence: \_\_\_\_\_

Dear Provider,

Please complete the letter of attestation below and return as indicated on the additional information request letter.

**SECTION I**

The beneficiary needs a home sleep study because:  
(Please check one)

- The beneficiary has a high pretest probability of obstructive sleep apnea syndrome (OSAS) as evidenced by clinical features, signs and symptoms.  
**Please complete Section II and III.**
- A diagnosis of OSAS has been established, therapy has been initiated and response to treatment is to be evaluated.  
**Please complete Section III.**

**SECTION II**

Complete this section if the home sleep study is requested because the beneficiary has a high pretest probability of OSAS.  
**Please also complete Section III.**

1. Does the beneficiary have a high pretest probability of OSAS due to the following conditions? (Check all that apply.)
  - Age  Gender  High body mass index (BMI)
  - Loud snoring  Awakening with gasping or snoring
  - Excessive daytime sleepiness
  - Observed cessation of breathing during sleep
  - Other (Please explain condition in Section IV.)
  - None of the above
2. Has the ordering provider determined the home sleep study is an appropriate alternative to in-laboratory poly somnography (PSG)?  
 Yes  No
3. Does the beneficiary have any of the following co-morbidities? (Check all that apply.)
  - Moderate to severe pulmonary disease
  - Neuromuscular disease  Congestive heart failure
  - Other specific significant co-morbidity (Please explain in Section IV.)
  - None of the above

4. Are any of the following sleep disorders suspected? (Check all that apply.)
  - Central sleep apnea  Periodic limb movement disorder
  - Insomnia  Parasomnia  Circadian rhythm disorder
  - Narcolepsy  Other sleep disorder
  - None of the above
5. Is the sleep complaint of short duration?  
 Yes  No
6. Does the beneficiary experience functional disability during the day due to the sleep-related disorder?  
 Yes  No

**SECTION III**

1. Has the sleep study been ordered by an authorized provider acting within the scope of his/her license?  
 Yes  No
2. What type of monitor is to be used?  
 Type II  Type III  Type IV
3. Has the monitor been validated in a typical home environment?  
 Yes  No
4. Is the monitor an FDA-approved portable monitoring device?  
 Yes  No
5. Will test results be reviewed and interpreted by a physician who is board-eligible/board-certified in sleep medicine?  
 Yes  No

**SECTION IV**

Please explain:

\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

I attest the information provided is true and accurate to the best of my knowledge. I understand Health Net Federal Services, LLC or designee may perform a routine audit and request the medical documentation to verify the accuracy of the information reported on this form.

Additional information: \_\_\_\_\_

Provider's printed name and title: \_\_\_\_\_

TIN: \_\_\_\_\_ Signature: \_\_\_\_\_ Date: \_\_\_\_\_

This document may contain information covered under the Privacy Act (5 USC §552a) and/or the Health Insurance Portability and Accountability Act (P.L. 104-191) and its various implementing regulations and must be protected in accordance with those provisions. If you have received this correspondence in error, please notify 1-844-866-WEST (9378) at once and destroy the documents and any copies you have made.

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