

**State of Oklahoma  
Oklahoma Health Care Authority  
Spinraza™ (Nusinersen) Prior Authorization Form**

**Member Name:** \_\_\_\_\_ **Date of Birth:** \_\_\_\_\_ **Member ID#:** \_\_\_\_\_

**Drug Information**

Physician billing (HCPCS code: \_\_\_\_\_)  Pharmacy billing (NDC: \_\_\_\_\_)  
**Start Date (or date of next dose):** \_\_\_\_\_ **Dose:** \_\_\_\_\_ **Regimen:** \_\_\_\_\_

**Billing Provider Information**

**NPI:** \_\_\_\_\_ **Provider Name:** \_\_\_\_\_  
**Provider Phone:** \_\_\_\_\_ **Provider Fax:** \_\_\_\_\_  
**Name of outpatient hospital facility where Spinraza™ will be delivered to and administered at:** \_\_\_\_\_

**Prescriber Information**

**Prescriber NPI:** \_\_\_\_\_ **Prescriber Name:** \_\_\_\_\_  
**Prescriber Phone:** \_\_\_\_\_ **Prescriber Fax:** \_\_\_\_\_ **Specialty:** \_\_\_\_\_

**Criteria**

**For Initial Authorization (Initial approval will be for the duration of 6 months):**

- Has the member previously been treated with Spinraza™ (nusinersen)? Yes \_\_\_ No \_\_\_  
A. If member has previously received nusinersen, please provide dates of previous doses: \_\_\_\_\_
- What is the member's diagnosis?  
 Spinal Muscular Atrophy (SMA)  
A. What type of SMA does the member have (0-4)? \_\_\_\_\_  
B. Does member currently have symptoms consistent with SMA? Yes \_\_\_ No \_\_\_  
C. Has the diagnosis been confirmed by molecular genetic testing? Yes \_\_\_ No \_\_\_  
D. Does member have biallelic pathogenic variants in the survival motor neuron gene 1 (SMN1)? Yes \_\_\_ No \_\_\_  
 Other: \_\_\_\_\_
- Is member currently dependent on permanent ventilation? Yes \_\_\_ No \_\_\_  
A. If member is currently dependent on permanent ventilation, please specify number of hours per day member requires ventilator support: \_\_\_\_\_
- Is Spinraza™ being prescribed by a neurologist, specialist with expertise in treatment of SMA, or an advanced care practitioner with a supervising physician who is a neurologist or specialist with expertise in treatment of SMA? Yes \_\_\_ No \_\_\_
- Has platelet count, coagulation laboratory testing, and quantitative spot urine protein testing been obtained? Yes \_\_\_ No \_\_\_  
A. If yes, are levels acceptable to the prescriber? Yes \_\_\_ No \_\_\_
- Does prescriber agree to do a platelet count, coagulation test, and quantitative spot urine protein testing prior to each dose? Yes \_\_\_ No \_\_\_
- Will Spinraza™ be administered in a healthcare facility by a specialist experienced in performing lumbar punctures? Yes \_\_\_ No \_\_\_
- Has a baseline assessment been performed and documented using at least one of the following exams as functionally appropriate: Hammersmith Infant Neurological Exam (HINE), Children's Hospital of Philadelphia Infant Test of Neuromuscular Disorders (CHOP-INTEND), Upper Limb Module (ULM) Test, or Hammersmith Functional Motor Scale Expanded (HFMSSE)? Yes \_\_\_ No \_\_\_  
A. If yes, please indicate the exam performed: \_\_\_\_\_  
B. Please provide member's baseline score to exam listed above: \_\_\_\_\_

Additional Information: \_\_\_\_\_

**For Continued Authorization:**

- Has the member previously been approved through the SoonerCare prior authorization process? Yes \_\_\_ No \_\_\_  
A. If no, please complete the initial authorization section above.
- Is member responding to the medication as demonstrated by a clinically significant improvement or maintenance of function from pre-treatment baseline status using the same exam as performed at baseline assessment? Yes \_\_\_ No \_\_\_
- Please indicate exam used to perform assessment: \_\_\_\_\_  
A. Please provide member's baseline score to exam listed above: \_\_\_\_\_  
B. Please provide member's current score to exam listed above: \_\_\_\_\_
- If member is currently dependent on permanent ventilation, please specify number of hours per day member requires ventilator support: \_\_\_\_\_

Additional Information: \_\_\_\_\_

**Prescriber Signature:** \_\_\_\_\_ **Date:** \_\_\_\_\_

*I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge.*

*Please do not send in chart notes. Specific information will be requested if necessary. Failure to complete this form in full will result in processing delays.*

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy  
Pharmacy Management Consultants  
Product Based Prior Authorization Unit  
Fax: 1-800-224-4014  
Phone: 1-800-522-0114 Option 4

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