

LOUISIANA MEDICAID NUSINERSEN (SPINRAZA®) CLINICAL AUTHORIZATION FORM

SECTION I — SUBMISSION					
Submitted to: Phone: Fax: Date: Aetna Better Health® of Louisiana 1-855-242-0802 1-844-699-2889	Date:				
SECTION II — PRESCRIBER INFORMATION					
Last Name, First Name MI: NPI# or Plan Provider #: Specialty:	Specialty:				
Address: City: State: Zip Cod	le:				
Phone: Fax: Office Contact Name: Contact Phone:					
SECTION III – PATIENT INFORMATION					
Last Name, First Name MI: DOB: Phone: Male Female	ale				
	nown				
Address: City: State: ZIP (Code:				
Plan Name (if different from Section I): Member or Medicaid ID #: Plan Provider ID:	Plan Provider ID:				
Is the patient currently a hospital inpatient getting ready for discharge? Yes No Date of Discharge: Per No Date of Discharge:					
SECTION IV — PRESCRIPTION DRUG INFORMATION					
Requested Drug Name: Nusinersen (Spinraza®)					
Strength: Dosage Form: Route of Admin: Days' Supply: Dosage Interval/Directions for Use: Expected Therapy Duration/Start Date:					
To the best of your knowledge this medication is: New therapy/Initial request Continuation of therapy, date of initiation: Has this medication been prescribed by, or in consultation with, a physician who specializes in the treatment of spinal muscular atrophy? YesNo Has this recipient previously been treated with onasemnogene abeparvovec-xioi (Zolgensma®)? Onasemnogene abeparvovec-xioi treatment date and result Will the patient receive the drug in the physician's office? YesNo If no, list name and NPI of servicing provider/facility: If yes, please complete the following: HCPCS/CPT-4 Code:NDC#:Dose Per Administration:Other Codes: SECTION V — PATIENT CLINICAL INFORMATION Does the patient have a diagnosis of spinal muscular atrophy (SMA)?YesNo If yes, date diagnosed:					
Type I (infantile onset or Werdnig-Hoffman disease [ICD-10-CM G12.0], symptoms are present at birth or by 6 months of age, unable to sit without assistance)					

Type II (intermediate SMA [ICD-10-CM G12.1], symptoms develop but unable to stand or walk independently)	betweer	6 months and 12 mon	ths of age, able to sit unassisted		
Type III (mild SMA or Kugelberg-Welander disease (ICD-10-CM G1	-	-	early childhood and		
adolescence, able to stand and walk independently but may lose to the diagnosis been confirmed by genetic testing?Yes		rater in life)			
If yes, did the testing confirm 5q SMA homozygous gene mutation, homYesNo	nozygous	gene deletion, or comp	oound heterozygote?		
Does the patient require ventilator support for 16 or more hours per da lf yes, date of initiation:	ay?	YesNo			
Motor Milestone Test*	Score	Measurement Date	Specialty of Provider Administering Test		
For recipients ≤ 2 years of age: Hammersmith Infant Neurological Examination Section 2 (HINE-2)					
For ambulatory recipients ≥3 years of age: Hammersmith Functional Motor Scale Expanded (HFMSE)					
For non-ambulatory recipients >3 years of age: Revised Upper Limb Module (RULM)					
*Results of most recent motor milestone test MUST be included for bot	h initial a	nd continuation / reau	thorization requests.		
Name of Pertinent Laboratory Test(s)	Date of Test		Results		
SECTION VI — FOR CONTINUATION OF THERAPY / REAUTHORIZATION REQUESTS ONLY					
From baseline motor milestone score to most recent motor milestone shas the patient received a clinical benefit from Spinraza® therapy as eventor skills or ability to sit, crawl, stand or walk, or new motor milestones.	idenced l		ntenance of		
When considering all categories of motor milestones, are the number of greater than the number that shows worsening?YesNo	of categoi	ies that show improver	ment		
SECTION VII — ADDITIONAL CLINICAL INFORMATION					
PHARMACY INFORMATION (OPTIONAL) Pharmacy Name:		Phone:			
By signing this request, the prescriber attests that the information pro knowledge. Also, by signing and submitting this request form, the pre section of the criteria specific to this request, if applicable.					
Signature of Prescriber:		Date:			