

**Pharmacy Prior Approval Request for
Zolgensma**

Beneficiary Information

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

Prescriber Information

6. Prescribing Provider NPI #: _____		
7. Requester Contact Information		
Name: _____	Phone #: _____	Ext. _____

Drug Information

8. Drug Name: _____	9. Strength: _____
10. Quantity Per 30 Days: _____	11. Length of Therapy (in days): <u>4 weeks</u>

Clinical Information

1. Is the member less than 2 years of age? Yes ____ No ____	
2. Does the member have a diagnosis of spinal muscular atrophy (SMA), with bi-allelic mutations in the survival motor neuron 1 (SMN1) gene? Yes ____ No ____ (Please attach additional documentation)	
3. Does genetic testing confirm the presence of one of the following: Yes ____ No ____ (Please attach additional documentation and choose one or more of the following)	
<input type="checkbox"/> Homozygous deletions of SMN1 gene (e.g., absence of the SMN1 gene)	
<input type="checkbox"/> Homozygous mutation in the SMN1 gene (e.g., biallelic mutations of exon 7);	
<input type="checkbox"/> Compound heterozygous mutation in the SMN1 gene [e.g., deletion of SMN1 exon 7 (allele 1) and mutation of SMN1 (allele 2)]	
4. Is this medication being prescribed by or in consultation with a neurologist? Yes ____ No ____	
5. Does the member have advanced SMA (e.g., complete paralysis of limbs, permanent ventilator dependence, tracheostomy, non-invasive ventilation beyond the use for sleep)? Yes ____ No ____ (please attach documentation)	
6. Has the member been previously treated with Zolgensma? Yes ____ No ____	
7. Have documents been included for one of the following baseline scores:	
<input type="checkbox"/> Children's Hospital of Philadelphia Infant Test of Neuromuscular Disorder (CHOP-INTEND) score	
<input type="checkbox"/> Hammersmith Infant Neurological Examination (HINE) Section 2 motor milestone score	
8. Have documents been included for both of the following:	
- Baseline laboratory tests demonstrating Anti-AAV9 antibody titers ≤ 1:50 as determined by ELISA binding immunoassay	
- Baseline liver function test, platelet counts, and troponin-L	
9. Is Zolgensma be prescribed concurrently with Spinraza? Yes ____ No ____	
10. Does the member have an active viral infection? Yes ____ No ____	
11. Does the Total dose exceed 1.1 x 10 ¹⁴ vector genomes (vg) per kilogram (kg) body weight? Yes ____ No ____	
12. Is Zolgensma being given in conjunction with pre and post infusion parenteral corticosteroids? Yes ____ No ____	

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 all forms and lab work to: (866) 399-0929

Pharmacy PA Call Center: (833) 585-4309