Trillium Health Resources Pharmacy Prior Approval Request for



Zolgensma

Member Information					
1.	Last Name:	2. First Name:			
3.	Trillium ID #:	2. First Name:4. Date of Birth:		5. Gender:	
	criber Information				
1.	Prescriber Name:	2. NPI #			
3.	Requestor Name (Nurse/Office Staff): _				
4.	Requestor Name (Nurse/Office Staff): _ Mailing Address: Phone #:		City:	State:	Zip:
3.	Phone #:	Ext	Fax #:		
Drug Information					
1. [1. Drug Name: Zolgensma 2. Strength: 3. Quantity Per 30 Days:				
4. Length of Therapy: ⊠ 1 Dose					
Clinical Information					
1. 2.	Is the member less than 2 years of age? \square Yes \square No Does the member have a diagnosis of spinal muscular atrophy (SMA), with bi-allelic mutations in the survival motor neuron 1 (SMN1) gene? \square Yes \square 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) □ Homozygous deletions of SMN1 gene (e.g., absence of the SMN1 gene)				
	☐ Homozygous mutation in the SMN1 gene (e.g., biallelic mutations of exon 7);				
	☐ Compound heterozygous mutation in the SMN1 gene [e.g., deletion of SMN1 exon 7 (allele 1) and mutation of SMN1 (allele 2)]				
4. 5.	Is this medication being prescribed by cooper the member have advanced SMA	(e.g., complete para	ysis of limbs, permane	nt ventilator deper	
tracheostomy, non-invasive ventilation beyond the use for sleep)? ☐ Yes ☐ No (please a 6. Has the member been previously treated with Zolgensma? ☐ Yes ☐ No				case attach abcam	Citation
7.	Have documents been included for one				
☐ Children's Hospital of Philadelphia Infant Test of Neuromuscular Disorder (CHOP-INTEND) score					
	☐ Hammersmith Infant Neurological Examination (HINE) Section 2 motor milestone score ☐ Newborn Screening results indicating baby has SMA				
_	-		□V□N.		
8.	Have documents been included for both ☐ Baseline laboratory tests demonstration immunoassay	ating Anti-AAV9 antil	oody titers ≤ 1:50 as de	etermined by ELISA	binding
_	☐ Baseline liver function test, platelet	•			
	Is Zolgensma being prescribed concurre	•			
10. Does the member have an active viral infection? Yes No					
 11. Does the Total dose exceed 1.1 x 1014 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 					
		· ,			
Çi.	gnature of Prescriber		Dato:		
31	gnature 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.