

Spinal Muscular Atrophy: Evrysdi

Member Information

1.	Last	Name:	2. First Name:				
3.	Trilli	ium ID #:	4. Date c	of Birth:	5. Gende	er:	
_							
		r Information					
		Prescriber Name: 2. NPI #: Requestor Name (Nurse/Office Staff):					
3.	Req	uestor Name (Nurse/Off	ce Staff):				
4. r	IVIAI	ling Address:		City:	State:	Zip:	
5.	Pho	ne #:	E;	Kt Fax #:			
Drug Information							
1.	Dru	Drug Name: Evrysdi 2. Strength: 3. Quantity per 30 Days:					
4.	Leng	Length of Therapy (in Days): 🛛 up to 30 Days 🖓 60 Days 🖓 90 Days 🖓 120 Days 🖓 180 Days 🖓 365 Days					
		□ Other					
Clinical Information							
Fo	For initial authorization requests, please answer questions 1-5						
1.	Is the member 2 months of age or older? Yes No						
		Does the member have a diagnosis of 5q-autosomal recessive spinal muscular atrophy (SMA)?					
		Does the member have SMA phenotype 1, 2, 3? Yes No					
4.	Will the member use Evrysdi concomitantly with nusinersen (Spinraza) or onasemnogene abeparvovec-xioi						
_		(Zolgensma)? Yes No					
5.	Is this medication being prescribed by or in consultation with a neurologist? Yes No						
Fo	For reauthorization, please answer questions 1-7						
6.	Has the member experienced any treatment related adverse effects or unacceptable toxicity?						
7.		Has the member had clinically meaningful response to treatment as demonstrated by at least 1 of the following:					
	Stability or improvement in net motor function/milestones, including but not limited to the following						
	validated scales: Hammersmith Infant Neurologic Exam (HINE), Hammersmith Functional Motor Scale						
	Expanded (HFMSE), Children's Hospital of Philadelphia Infant Test of Neuromuscular Disorders (CHOP						
	INTEND), Bayley Scales of Infant and Toddler development Third Ed. (BSID-III), 6-minute walk test (6MWT), upper limb module (ULM), etc.						
			in respiratory function tes	te le al forced vital	capacity(E)(C) at 1		
					, _	nfection in	
	Reduction in exacerbations necessitating hospitalization and/or antibiotic therapy for respiratory infection in the preceding year/timeframe						
	 Stable or increased member weight (for members without a gastrostomy tube) 						
		-					

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.