

Please note: All information below is required to process this request.

Fax to 1-844-403-1029

Mon-Sat: 7am to 7pm Central

Evrysdi® Prior Authorization Request Form (Page 1 of 3) DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

Member Information (required)			Provider Information (required)			
Member Name:		Provider Name:				
Insurance ID#:		NPI#:	NPI#:		Specialty:	
		Office Phone:				
Street Address:			Office Fax:			
State:	Zip:	Office Street Address:				
Phone:			State:	State: Zip:		
N	ledication Infor	mation (require	d)			
Medication Name:			<i>'</i>	Dosage Fo	orm:	
☐ Check if requesting brand			Directions for Use:			
r continuation of thera						
	Clinical Inform	nation (required)				
elow:						
□ Spinal muscular atrophy (SMA): Type □ Other diagnosis: ICD-10 Code(s):						
			lowing special	ists:		
ith expertise in the diag	nosis and treatment of S	MA				
gene deletion or mutati	on (e.g., homozygous de	letion of exon 7 at lo				
eterozygous mutation (e	e.g., deletion of SMN1 ex	on 7 [allele 1] and m	utation of SMN	1 [allele 2]) _		
		□ Yes □ No				
endent on use of non-in	vasive ventilation beyond	d use for naps and n	ghttime sleep?	□ Yes □	No	
er exams listed below	(based on patient's age a	and motor ability) bee	n conducted to	o establish ba	seline motor ability I	
eurologist?	e Evnanded (HEMSE)					
Infant Neurological Ex	am (HINE) (infant to earl	y childhood)				
		lar Disorders (CHOE	INTEND)			
spilai di Filliadelpilia ili	iani Tesi oi Neuromuscu					
n Measure 32 (MFM-32		iai Disorders (Crior				
n Measure 32 (MFM-32				nt of SMA (e.g	յ., Spinraza)?	
	er Information State: Normand r continuation of thera elow: phy (SMA): Type sted medication is prese ith expertise in the diage copies? or deletion of genes in gene deletion or mutation eterozygous mutation (eleterozygous mutation (eleterozygous mutation) eterozygous eter	State: Zip: Medication Information (required) rand r continuation of therapy Clinical Inform elow: phy (SMA): Type sted medication is prescribed by or in consultation the expertise in the diagnosis and treatment of S copies? or deletion of genes in chromosomes 5q result gene deletion or mutation (e.g., homozygous desterozygous mutation (e.g., deletion of SMN1 extendent on invasive ventilation or tracheostomy? endent on use of non-invasive ventilation beyond the exams listed below (based on patient's age at eurologist? Functional Motor Scale Expanded (HFMSE) Infant Neurological Exam (HINE) (infant to early lodule (ULM) Test (Non ambulatory)	Provider Name: Provider Name: NPI#:	Provider Information (required) Provider Name: NPI#: Office Phone: Office Fax: Office Street Address: City: State: Medication Information (required) Strength: rand Directions for Use: Phy (SMA): Type	Provider Name: NPI#: Specialty: Office Phone: Office Fax: Office Street Address: City: State: Medication Information (required) Strength: Dosage Form Clinical Information (required) Clinical Information (required) elow: phy (SMA): Type	



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9.	in we hi	patient has previously received gene therapy for the treatment of SMA (e.g., Zolgensma), provider to attests that there has been an adequate response to gene therapy (e.g., sustained decrease in at least one motor test score over a period of 6 months) or or or sening in clinical status since receiving gene therapy as demonstrated by a decline of minimally clinical important difference fro ghest score achieved on one of the following exams: HFMSE: decline of at least points on kicking and points on any other milestones (excluding voluntary grasp) HINE-2: decline of at least points CHOP INTEND: decline of at least points
		y limit requests:
		the quantity requested per DAY?the reason for exceeding the plan limitations?
		ation or loading dose purposes
	Red	ent is on a dose-alternating schedule (e.g., one tablet in the morning and two tablets at night, one to two tablets at bedtime) juested strength/dose is not commercially available er:
		prization:
If th	is is	a reauthorization request, answer the following:
1.	Ho	v many SMN2 copies?
2.	as	vide documentation of positive clinical response to therapy (e.g., chart notes, laboratory values) from pretreatment baseline status demonstrated by the most recent results (less than 1 month prior to reauthorization request) from one of the following exams: One of the following HINE-2 milestones
		One of the following HINE-2 milestones Improvement or maintenance of previous improvement of at least a 2-point (or maximal score) increase in ability to kick Improvement or maintenance of previous improvement of at least a 1-point increase in any other HINE-2 milestone (e.g., head control, rolling, sitting, crawling, etc.), excluding voluntary grasp Patient exhibited improvement, or maintenance of previous improvement in more HINE motor milestones than worsening,
		from pretreatment baseline (net positive improvement)
		□ Patient has achieved and maintained any new motor milestones when they would otherwise be unexpected to do so (e.g., sit unassisted, stand, walk)
		One of the following HFMSE milestones
		☐ Improvement or maintenance of a previous improvement of at least a 3-point increase in score from pretreatment baseline ☐ Patient has achieved and maintained any new motor milestone from pretreatment baseline when they would otherwise be unexpected to do so (e.g., sit unassisted, stand, walk)
		One of the following ULM test milestones
		☐ Improvement or maintenance of a previous improvement of at least a 2-point increase in score from pretreatment baseline ☐ Patient has achieved and maintained any new motor milestone from pretreatment baseline when they would otherwise be unexpected to do so (e.g., sit unassisted, stand, walk)
		One of the following CHOP-INTEND milestones
		□ Improvement or maintenance of a previous improvement of at least a 4-point increase in score from pretreatment baseline □ Patient has achieved and maintained any new motor milestone from pretreatment baseline when they would otherwise be unexpected to do so (e.g., sit unassisted, stand, walk)
		One of the following MFM-32 milestones
		□ Improvement or maintenance of a previous improvement of at least a 3-point increase in score from pretreatment baseline □ Patient has achieved and maintained any new motor milestone from pretreatment baseline when they would otherwise be unexpected to do so (e.g., sit unassisted, stand, walk)
3.	ls t	ne patient dependent on invasive ventilation or tracheostomy? □ Yes □ No



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4.	Is the patient dependent on the use of non-invasive ventilation beyond use for naps and nighttime sleep? Yes No
5.	Is the requested medication prescribed by or in consultation with a neurologist with expertise in the diagnosis and treatment of SMA? Yes D No
6.	Is the patient is receiving concomitant chronic survival motor neuron (SMN) modifying therapy for the treatment of SMA (e.g., Spinraza)? Ves No
7.	Has the patient previously received gene replacement therapy for the treatment of SMA (e.g., Zolgensma)? No
8.	Was there inadequate response to gene therapy (e.g., sustained decrease in at least one motor test score over a period of 6 months)? If so, submit medical records (e.g., chart notes) documenting the inadequate response to gene therapy.
	there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to review?
Pleas	se note: This request may be denied unless all required information is received.

For urgent or expedited requests please call 1-855-401-4262.

This form may be used for non-urgent requests and faxed to 1-844-403-1029.