

Date: _____ Date Medication Required: _____
 Ship to: Physician Patient's Home Other _____

Patient Information

Last Name:		First Name:		Middle:	DOB: ___/___/___	
Address:			City:		State:	Zip:
Daytime Phone:		Evening Phone:		Sex: <input type="checkbox"/> Male <input type="checkbox"/> Female		

Insurance Information (Attach copies of cards)

Primary Insurance:		Secondary Insurance:			
ID #	Group #	ID #	Group #		
City:		State:	City:		State:

Physician Information

Name:		Specialty:		NPI:	
Address:			City:		State: Zip:
Phone #:		Secure Fax #:		Office Contact:	

Primary Diagnosis

ICD-10 Code: _____
 Duchenne muscular dystrophy (DMD) Other: _____

Prescription Information

MEDICATION	STRENGTH	DIRECTIONS	QUANTITY	REFILLS
Viltepso (viltolarsen)				

Clinical Information

***** Please submit supporting clinical documentation *****

INITIAL THERAPY CONTINUATION OF THERAPY; Therapy start date: _____

- Has patient had a positive response to the prescribed therapy within the last 30 days?
 Yes ****Mark all that apply**** No Not applicable
 - Ambulatory function with a 6 minute walk test distance (6MWT) ≥ 201 m? Yes: _____ m No
 - Ambulatory function with a time-to-stand (TTSTAND) < 10 seconds? Yes: _____ seconds No
 - Stable cardiac function with left ventricular ejection fraction (LVEF) ≥ 40%? Yes: _____ % No
 - Stable pulmonary function with predicted forced vital capacity (FVC) ≥ 50%? Yes: _____ % No
 - Other: _____
- Is Viltepso prescribed concurrently with an oral corticosteroid? Yes No No, contraindicated/intolerant
- Is Viltepso prescribed concurrently with other exon-skipping therapies (e.g. Exondys 51, Vyondys 53)? Yes No
- Please document patient's weight: _____ kg

Complete this section ONLY if the patient is initiating therapy OR if the patient is new to this health plan:

- Is therapy prescribed by or in consultation with a neurologist? Yes No
- If DMD, is mutation amenable to exon 53 skipping confirmed with genetic testing? Yes, mutation: _____ - _____ No
- Has the patient had an inadequate response (evidence by significant decline in 6MWT, LVEF, or FVC) despite adherent use of an oral corticosteroid (e.g., prednisone, Emflaza™) for ≥ 6 months? Yes No No, contraindicated/intolerant

Complete this section ONLY for indications other than DMD:

- Has patient tried and failed, or is contraindicated to, accepted standards of care? Yes No
****If yes, submit documentation and answer the following:****
 - Please list all previous therapies: _____
 - Was patient adherent to previously tried therapies? Yes No No, patient intolerant to drug

Physician's Signature: _____ Date: _____ DAW

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Telephone: (800) 514-0083 option 2
Fax: (866) 374-1579

Viltolarsen (Viltepso)

Prior Authorization Form/Prescription

Date: _____ Date Medication Required: _____
Ship to: Physician Patient's Home Other _____

Patient Name: _____ DOB: _____

INFORMATION BELOW IS TO BE COMPLETE BY THE HEALTH PLAN/EPS PA STAFF

Authorization Information

Authorization number:	Decision Due Date:
J-Code:	Coverage: <input type="checkbox"/> State excludes <input type="checkbox"/> COB (secondary)
Line of Business: <input type="checkbox"/> Commercial <input type="checkbox"/> Health Insurance Marketplace <input type="checkbox"/> Medicaid <input type="checkbox"/> Medicare	Benefit: <input type="checkbox"/> Medical <input type="checkbox"/> Pharmacy
Criteria: <input type="checkbox"/> Centene Policy Date Policy last reviewed/approved by plan (we want to be sure we are using the version approved by your plan): _____ <input type="checkbox"/> State Specific (please include policy)	
Medicare only criteria for CY2019 and CY2020: <input type="checkbox"/> PART B use LCD or NCD <input type="checkbox"/> PART D use the Medicare Part D Viltepso specific criteria	