## Clinical Criteria Worksheet: Duchenne Muscular Dystrophy (DMD) Drug

## **Claim Submission**

**Enrollee Information** 

- Claim processing may be delayed if the information submitted in this worksheet is illegible.
- If the worksheet is left blank or information is missing the claim will be rejected for not enough documentation and reimbursement will be delayed.
- A claim should not be submitted until the drug has been administered to the patient.
- The manufacturer invoice showing the acquisition cost of the drug administered, including all discounts, rebates, and incentives must be submitted with the claim. The invoice must be dated within 6 months prior to the date of service and/or should include the expiration date of the drug, or it will be rejected for not enough documentation.

Enrollee Last Name:												_	Enrollee First Name:													
												ľ														
Date of Birth (MM/DD/YYYY):												Enrollee Medicaid ID (2 letters, 5 numbers, 1 letter):														
		/			/																					
Add	ress	:			_					_												_				
City	City, Town or Post Office:																	State:			ZIP Code:					
Pre	escr	ibe	r In	for	ma	tior	1																			
Pres	scrib	er La	ast N	lame	:								Prescriber First Name:													
Nat	iona	l Pro	vide	r Ide	entifi	er(N	NPI)	Nun	nber	:	•	_		•	•			<u> </u>			•	•				
Pre	ferre	d Co	ntac	t (Te	elepl	none	Nur	nbe	r)	_																
			]_				]_																			

Enrollee Last Name:	Enrollee First Name:													
Clinical Criteria – Drug Information														
Drug Administration:														
Provide the date of drug administration (MM/DD/YYYY):  / / / / / / / / / / / / / / / / / / /														
Provide the expiration date of the drug if the invoice administration (MM/DD/YYYY):  / / / / / / / / / / / / / / / / / / /	: date	e is g	reate	er tha	an 6	5 ma	onth	s fro	m th	e dat	te of	drug		
Drug Name and Strength:														
☐ casimersen (AMONDYS 45™)														
☐ eteplirsen (EXONDYS 51™)														
□ viltolarsen (VILTEPSO®)														
☐ golodirsen (VYONDYS 53™)														
☐ other DMD drug (unclassified code J3490)														
Strength:	Directions:													
Quantity:	_													
New Treatment: Yes No														
If <b>No</b> , date therapy initiated:														
Clinical Criteria – Diagnosis														
1. Duchenne Muscular Dystrophy  Other:														
<ol> <li>Is the patient currently being treated with another</li> </ol>														
☐ Yes ☐ No		1-1	. 5 -	_	. ,									
If this is a continuation of therapy for the patient and	you i	nave	alrea	dy re	ecei	ved	pay	ment	for	previ	ous			

If this is a continuation of therapy for the patient and you have already received payment for previous administration for this medication, provide attestation signature on page 3. Additional information on page 3 is not necessary.

Enr	Enrollee Last Name:													Enrollee First Name:											
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For	For diagnosis of Duchenne Muscular Dystrophy:																								
	Does the patient have documented genetic testing confirming the mutation of the DMD gene is amendable to exon 45, 51, or 53 skipping?  Yes No																								
4.	If <b>Yes</b> , please provide the date of the lab test result:																								
	Does the pateint have documented stable dose of corticosteroids prior to starting DMD therapy?  Yes No  If <b>Yes</b> , please provide therapy length:  Months:																								
	If <b>No</b> , please provide rationale for not utilizing a corticosteroid:																								
	Rationale:																								
	Does the patient have documented kidney function testing prior to starting therapy? ( <i>skip the question if the administered drug is eteplirsen</i> )   Yes   No												n if												
	If <b>Yes</b> , please provide the date of the testing (MM/DD/YYYY):  / / / / / / / / / / / / / / / / / / /																								
		tion																							
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Pre	Prescriber Signature (Required)																	ate (ſ	MM/	DD/\	(YYY)	)			