

## Duchenne Muscular Dystrophy: Exondys 51

### Member Information

1. Last Name: \_\_\_\_\_ 2. First Name: \_\_\_\_\_  
3. Trillium ID #: \_\_\_\_\_ 4. Date of Birth: \_\_\_\_\_ 5. Gender: \_\_\_\_\_

### Prescriber Information

1. Prescriber Name: \_\_\_\_\_ 2. NPI #: \_\_\_\_\_  
3. Requestor Name (Nurse/Office Staff): \_\_\_\_\_  
4. Mailing Address: \_\_\_\_\_ City: \_\_\_\_\_ State: \_\_\_\_\_ Zip: \_\_\_\_\_  
5. Phone #: \_\_\_\_\_ Ext. \_\_\_\_\_ Fax #: \_\_\_\_\_

### Drug Information

1. Drug Name: **Exondys 51** 2. Strength: \_\_\_\_\_ 3. Quantity per 30 Days: \_\_\_\_\_  
4. Length of Therapy (in Days):  up to 30 Days  60 Days  90 Days  120 Days  180 Days

### Clinical Information

#### For initial authorization requests:

1. What is the member's weight? \_\_\_\_\_
2. Does the member have a diagnosis of Duchenne Muscular Dystrophy?  **Yes**  **No**
3. Are medical records attached to this request that confirm the mutation of the Duchenne Muscular Dystrophy gene is amenable to exon 51 skipping?  **Yes**  **No**
4. Is Exondys 51 being prescribed by or in consultation with a neurologist?  **Yes**  **No**
5. Is the member taking any other RNA antisense agent or any other gene therapy?  **Yes**  **No**
6. Is the member receiving a dose that does not exceed 30mg/kg once per week?  **Yes**  **No**

#### For reauthorization:

7. Please attach documentation that shows the member:  
 Has shown an improvement in dystrophin levels **OR**  
 Is not ventilator dependent **OR**  
 Has some functional use of upper extremities **OR**  
 Has an ability to walk with or without assistive devices

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.