

**Beneficiary Information**

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

**Prescriber Information**

6. Prescriber Name: \_\_\_\_\_ NPI #: \_\_\_\_\_  
Mailing address: \_\_\_\_\_ City: \_\_\_\_\_ State: \_\_\_\_\_ ZIP: \_\_\_\_\_  
7. Requester Contact Information: \_\_\_\_\_  
Name: \_\_\_\_\_ Phone #: \_\_\_\_\_ Fax #: \_\_\_\_\_

**Drug Information**

8. Drug Name: \_\_\_\_\_ 9. Strength: \_\_\_\_\_ 10. Quantity Per 30 Days: \_\_\_\_\_  
11. Length of Therapy: Initial Request:  up to 30 days  60 days  90 days  120 days  180 days  
Reauthorization Request:  up to 30 days  60 days  90 days  120 days  180 days  365 days

**Clinical Information**

**Initial Authorization Request:**

1. Is the beneficiary age 2 or older? Yes \_\_\_ No \_\_\_
  2. Does the beneficiary have a diagnosis of Duchenne Muscular Dystrophy confirmed by genetic testing (documentation required)? Yes \_\_\_ No \_\_\_
  3. Has the beneficiary tried prednisone? Yes \_\_\_ No \_\_\_
- Answer questions 3a and 3b when the response to question 3 is 'Yes'.**
- 3a. Has the beneficiary had an inadequate treatment response to prednisone? If yes, documentation required. Yes \_\_\_ No \_\_\_
  - 3b. Has the beneficiary experienced unmanageable and clinically significant side effects such as significant weight gain/obesity, persistent psychiatric/behavioral issues, diabetes, hypertension, or Cushingoid appearance? If yes, documentation required. Yes \_\_\_ No \_\_\_
4. A baseline motor milestone assessment is required. **Please select all that apply and submit documentation for each.**  
 6-minute walk test (6MWT)  
 North Star Ambulatory Assessment (NSAA)  
 Motor Function Measure (MFM)  
 Hammersmith Functional Motor Scale (HFMS)  
 Other. Please explain: \_\_\_\_\_  
 None of the above.
  5. Is the medication prescribed by or in consultation with a neurologist? Yes \_\_\_ No \_\_\_
  6. Will the provider ensure the Emflaza is not being given concurrently with live vaccinations? Yes \_\_\_ No \_\_\_
  7. Is Emflaza dosing for Duchenne Muscular Dystrophy in accordance with the USFDA approved labeling? Yes \_\_\_ No \_\_\_

**Reauthorization Request:**

Please check all of the applicable clinical benefits the beneficiary has received from Emflaza therapy. (Please submit documentation for each.)

8. A baseline motor milestone assessment is required. Please select all that apply and submit documentation:  
 Stabilization, maintenance or improvement of muscle strength  
 Stabilization, maintenance or improvement of pulmonary function  
 Improvement in motor milestone assessment scores from baseline testing  
 Motor function is superior relative to that projected for the natural course of Duchenne Muscular Dystrophy  
 Other – Please explain: \_\_\_\_\_  
 None of the above.

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.

Fax this form to: 1-877-234-4274, or call Pharmacy Prior Authorization: 1-866-885-1406