

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: ___up to 30 days ___60 days ___90 days ___120 days ___180 days

Clinical Information

For initial authorization requests (please answer questions 1-11):

1. What is the beneficiary's weight? _____
2. Does the beneficiary 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 53 skipping? Yes ___ No ___
4. Is Vyondys 53/Viltepso being prescribed by or in consultation with a neurologist? Yes ___ No ___
5. Does the beneficiary have meaningful voluntary motor function? Yes ___ No ___
6. Has the beneficiary been assessed for any physical therapy and/or occupational therapy needs? Yes ___ No ___
7. Has the beneficiary's serum cystatin C, urine dipstick, and urine protein-to-creatinine ratio been measured prior to the start of therapy? Yes ___ No ___
8. Does the prescriber attest that the beneficiary's serum cystatin C, urine dipstick, and urine protein-to-creatinine ratio will be measured during treatment (monthly urine dipstick with serum cystatin C and urine protein-to-creatinine ratio every 3 months)? Yes ___ No ___
9. Is there documentation of baseline movement/functional testing? Yes ___ No ___
10. Is the beneficiary taking any other RNA antisense agent or any other gene therapy? Yes ___ No ___
11. Is the beneficiary receiving a dose that does not exceed 30mg/kg once per week (Vyondys 53) or 80mg/kg once per week (Viltepso)? Yes ___ No ___

For reauthorization (please answer questions 1-13):

12. Please attach documentation that shows the beneficiary has demonstrated a response to therapy compared to pretreatment baseline.
13. Has the beneficiary experienced any treatment-restricting adverse effects? Yes ___ No ___

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.