

Duchenne Muscular Dystrophy: Vyondys 53 and Viltepso

| Member Information | |
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| 1. | Last Name: 2. First Name: |
| 3. | Last Name: 2. First Name: Trillium ID #: 4. Date of Birth: 5. Gender: |
| Pres | criber Information |
| 1. | Prescriber Name: 2. NPI #: |
| 3. | Requestor Name (Nurse/Office Staff): |
| 4. | Mailing Address: State: Zip: |
| 5. | Phone #: Ext Fax #: |
| Drug | g Information |
| 1. | Drug Name: 2. Strength: 3. Quantity per 30 Days: |
| | Length of Therapy (in Days): ☐ up to 30 Days ☐ 60 Days ☐ 90 Days ☐ 120 Days ☐ 180 Days |
| Clini | ical Information |
| Fo | r initial and re-authorization requests: (please answer questions 1-11) |
| 1. | What is the member's weight? |
| | 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 53 skipping? ☐ Yes ☐ No |
| 4. | Is Vyondys 53/Viltepso being prescribed by or in consultation with a neurologist? ☐ Yes ☐ No |
| 5. | Does the member have meaningful voluntary motor function? ☐ Yes ☐ No |
| 6. | Has the member been assessed for any physical therapy and/or occupational therapy needs? ☐ Yes ☐ No |
| 7. | Has the member'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 member'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)? \square Yes \square No |
| ۵ | Is there documentation of baseline movement/functional testing? Yes No |
| | Is the member taking any other RNA antisense agent or any other gene therapy? Yes No |
| | Is the member receiving a dose that does not exceed 30mg/kg once per week for (Vyondys 53) or 80mg/kg once |
| | per week (Viltepso)? ☐ Yes ☐ No |
| Fo | r reauthorization: (please answer questions 12 &13) |
| | Please attach documentation that shows the member has demonstrated a response to therapy compared to pretreatment baseline. |
| 13. | Has the member experienced any treatment-restricting adverse effects? ☐ Yes ☐ No |
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| Si | gnature 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.