POLICY NAME Duvyzat (givinostat) POLICY # 3257P

Criteria

| Cove | erage Criteria |
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| | Diagnosis of Duchenne Muscular Dystrophy confirmed by one of the following: • Genetic testing documenting a mutation in the dystrophin (DMD) gene • Muscle biopsy documenting lack of muscle dystrophin |
| | Age 6 years or older |
| | Prescribed by or in consultation with a physician who specializes in the treatment of Duchenne Muscular Dystrophy (DMD) and/or neuromuscular disorders |
| | Patient is currently ambulatory (able to walk independently) |
| | Documented concurrent use (for at least the last 6 months) of prednisone unless member has experienced at least one of the following significant intolerable adverse effects (AE) • Cushingoid appearance • Central (truncal) obesity • Undesirable weight gain defined as a 10% of body weight gain increase over a 6-month period • Diabetes and/or hypertension that is difficult to manage • Severe behavioral AE that would require a prednisone dose reduction • Clinically significant growth stunting as evidenced by decline in mean height percentile from baseline, decrease in growth velocity or decrease in serum bone formation biomarkers |
| | If member is unable to tolerate prednisone, concurrent use of generic deflazacort is required |
| | Documentation of a baseline motor milestone score from one of the following assessments: • 4-stair climb (4SC) • North Star Ambulatory Assessment (NSAA) • 6-minute walk test (6MWT) • Time to stand test (TTSTAND) |
| | Review of clinical documentation and confirming that patient has met all of the above requirements for treatment completed by both a pharmacist and medical director |

| Exclu | sion | Criteri | a – A | ny of | the | followi | ing pi | revents | covera | ıge |
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Duvyzat will not be covered in combination with or in patients who have previously received any