PYQUVI™ (deflazacort) oral suspension
Patient Enrollment Form

Fax: (877) 860-1978
Phone: (847) 393-4099

Patient Information *(Please Print)*

Last Name: First Name: SSN: Sex: M ◯ F ◯

Address: City: State: Zip:

Phone *(Day)*: Phone *(Evening)*: Cell: Preferred method of contact: Day # ◯ Evening # ◯ Cell # ◯

DOB: Age: Weight *(kgs.)*: Height *(Inches)*:

If Patient is a Minor, Guardian/Parent Name: Relationship to Patient:

Emergency Contact/Relation: Phone#:

Insurance Information

Please attach front and back of patient's insurance card, prescription card, and/or Medicaid card

Primary Insurance : Phone #:

Policy Holder: Policy # Group #:

Secondary Insurance: Phone #:

Policy Holder: Policy # Group #:

Physician Information

Prescriber: Institution:

NPI:

Address: City: State: Zip:

Name & Title of Office Contact: Office Contact Email:

Office Contact Phone #: Office Contact Fax:

Prior Authorization Office Contact: Prior Authorization Office Contact #:

Prescription Information

PYQUVI™ (deflazacort) 22.75 mg/ mL oral suspension

◯ SIG: Take 0.9 mg/kg orally once daily

◯ Take ____ milligram orally once daily

Other directions Quantity: ◯ 30 Days ◯ 90 Days Other Refills

Clinical Information

Please fax clinical documentation to pharmacy along with Enrollment Form

Primary Diagnosis: Date of Diagnosis:

Patient Age at Diagnosis:

Please check applicable ICD-10 code: Therapy Start Date:

◯ Duchenne Muscular Dystrophy (G71.01) ◯ Other

Allergies: ◯ No Known Drug Allergies

I certify I am prescribing PYQUVI™ for this patient for a medically necessary purpose. Date Written:

Dispense as Written: _____ Substitution Allowed: _____

(Stamped Signatures Are Not Valid) *(Stamped Signatures Are Not Valid)*

Please see Important Safety information on page 2. For more information, please see full Prescribing Information, including Instructions for Use or go to PyquviHCP.com

Pyquvi™ (deflazacort) Oral Suspension

INDICATION

PYQUVI™ is a corticosteroid indicated for the treatment of Duchenne muscular dystrophy (DMD) in patients 5 years of age and older.

IMPORTANT SAFETY INFORMATION

PYQUVI is contraindicated in patients with a hypersensitivity to deflazacort or any of the inactive ingredients.

WARNINGS AND PRECAUTIONS

- **Alterations in Endocrine Function:** Corticosteroids can cause serious and life-threatening endocrine alterations, especially with chronic use. Monitor for Cushing's syndrome, hyperglycemia, and adrenal insufficiency after withdrawal. Risk is higher in patients with hypopituitarism, primary adrenal insufficiency, congenital adrenal hyperplasia, altered thyroid function, or pheochromocytoma. Acute adrenal insufficiency or "withdrawal syndrome" may occur if discontinued abruptly and can be fatal; taper gradually. Dose may need to be increased during times of medical stress.

- **Immunosuppression and Increased Risk of Infection:** Increased risk of new, exacerbation, dissemination, or reactivation of latent infections, which can be severe or fatal. Signs of infection may be masked. Advise patients/caregivers to report recent infections or vaccinations. Patients without prior chickenpox or measles should avoid exposure and contact their healthcare provider immediately if exposed.

- **Alterations in Cardiovascular/Renal Function:** Monitor blood pressure; dietary salt restriction and potassium supplementation may be needed.

- **Gastrointestinal (GI) Perforation:** Risk is increased in patients with certain GI disorders (eg, active or latent peptic ulcers, diverticulitis, recent intestinal anastomoses, inflammatory bowel disease). Signs may be masked.

- **Behavioral and Mood Disturbances:** May include euphoria, insomnia, mood swings, personality changes, depression, and psychosis. Advise patients to seek medical attention if symptoms develop or worsen.

- **Effects on Bones:** Prolonged use increases risk of osteoporosis, and vertebral and long bone fractures. Monitor bone density with chronic use.

- **Ophthalmic Effects:** May cause cataracts, ocular infections, or glaucoma. If corticosteroids are used for >6 weeks, monitor intraocular pressure.

- **Vaccination:** Do not administer live or live-attenuated vaccines during immunosuppressive corticosteroid therapy. Administer such vaccines at least 4 to 6 weeks before starting PYQUVI.

- **Serious Skin Rashes:** Toxic epidermal necrolysis has been reported. Discontinue at the first sign of rash, unless the rash is clearly not drug related.

- **Effects on Growth and Development:** Long-term use of corticosteroids may slow growth and development in children.

- **Thromboembolic Events:** Observational studies have shown an increased risk. Use with caution in at-risk patients.

ADVERSE REACTIONS

The most common adverse reactions (≥10% and greater than placebo) are Cushingoid appearance, weight gain, increased appetite, upper respiratory tract infection, cough, pollakiuria, hirsutism, central obesity, and nasopharyngitis.

DRUG INTERACTIONS

Moderate or strong CYP3A4 inhibitors: Use one third of the recommended PYQUVI dose.

Moderate or strong CYP3A4 inducers: Avoid concomitant use as efficacy may be reduced.

To report SUSPECTED ADVERSE REACTIONS, contact Aucta Pharmaceuticals, Inc. at 1-800-655-9902, or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

Please see full Prescribing Information.

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