

Division: Pharmacy Policy	Subject: Prior Authorization Criteria
Original Development Date: Original Effective Date: Revision Date:	July 19, 2018 March 13, 2020, June 29, 2020, July 1, 2024

# Crysvita<sup>®</sup> (burosumab)

#### LENGTH OF AUTHORIZATION: Initial therapy: 6 months

Continuation of therapy: 1 year

# **INITIAL REVIEW CRITERIA**:

- Patient is six months of age or older.
- Patient has not received oral phosphate and/or active vitamin D analogs within a week prior to starting therapy.
- Prescribed by, or in consultation with a geneticist, nephrologist or endocrinologist.
- Diagnosis of X-linked hypophosphatemia (XLH) confirmed by identifying at least one of the following:
  - Serum fibroblast growth factor-23 (FGF23) level >30pg/mL in children; OR
  - Phosphate regulating gene with homology to endopeptidases located on the X chromosome (PHEXgene) mutations in the patient.
- Low-serum phosphate concentration (baseline labs required, drawn within 30 days of prior authorization submission) AND
- Reduced tubular resorption of phosphate corrected for glomerular filtration rate.

#### OR

• Patient is 2 years of age or older for the treatment of FGF23-related hypophosphatemia in tumor induced osteomalacia (TIO) associated with phosphaturic mesenchymal tumors that cannot be curatively resected or localized.

# **CONTINUATION OF THERAPY REVIEW CRITERIA:**

- Patient met initial review criteria; AND
- Patient has not experienced any treatment-restricting adverse effects; AND
- Dosing is appropriate as per labeling or is supported by compendia; AND
- Disease response as indicated by at least one of the following:
  - Increased serum phosphorus levels (official labs required).
  - A reduction in serum total alkaline phosphatase activity.
  - Improvement in symptoms (e.g. skeletal pain, linear growth).
  - Improvement in radiographic imaging of rickets/osteomalacia.

# **DOSING AND ADMINISTRTATION:**

- Refer to product labeling https://www.accessdata.fda.gov/scripts/cder/daf/
- Available as 10 mg/mL, 20 mg/mL, or 30 mg/mL in a single-dose vials for injection

