

**Blue Cross Blue Shield/Blue Care Network of Michigan  
Medication Authorization Request Form**



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This form is to be used by participating physicians to obtain coverage for **drugs covered under the medical benefit**. For commercial members only, please complete this form and submit via fax to 1-877-325-5979. If you have any questions regarding this process, please contact BCBSM Provider Relations and Servicing or the Medical Drug Helpdesk at 1-800-437-3803 for assistance.

PATIENT INFORMATION	PHYSICIAN INFORMATION
<b>Name</b>	<b>Name</b>
<b>ID Number</b>	<b>Specialty</b>
<b>D.O.B.</b> ___/___/___ MM/DD/YYYY <input type="checkbox"/> Male <input type="checkbox"/> Female	<b>Address</b>
<b>Diagnosis</b>	<b>City /State/Zip</b>
<b>Drug Name</b> <b>Crysvita</b>	<b>Phone:</b>
<b>Dose and Quantity</b>	<b>Fax:</b>
<b>Directions</b>	<b>NPI</b>
<b>Date of Service(s)</b>	<b>Contact Person</b>
	<b>Contact Person Phone / Ext.</b>

**STEP 1: DISEASE STATE INFORMATION**

**Required Demographic Information:**

*Patient Weight:* \_\_\_\_\_ *kg*  
*Patient Height:* \_\_\_\_\_ *ft*    \_\_\_\_\_ *inches*

Will the provider be administering the medication to the FEP member within the health plan's geographic service area?  
 Yes    No    *If No, a prior authorization is not required through this process.*

**Prior authorizations are required for FEP members that will be serviced by a provider within the health plan's geographic service area. If you are not a provider in the geographic service area, please contact the health plan for questions regarding the FEP member's benefit requirements.**

Is this member's FEP coverage primary or secondary coverage?  
 If primary, continue with question set.  
 If secondary, **an authorization is not needed through this process. Please contact the member's primary coverage for determination of benefit and additional information.**

**Site of Care:**

- A. At what location will the member be receiving the requested medication?
- Physician's office, home infusion, non-hospital affiliated ambulatory infusion center.
  - Outpatient hospital infusion center. Please provide the name of the infusion center and rationale why the patient must receive this medication in a hospital outpatient setting. \_\_\_\_\_
  - Other. Please specify. \_\_\_\_\_

**Criteria Questions:**

1. Has the patient been on Crysvita continuously for the last **6 months, excluding samples**? ***Please select answer below:***

**NO** – this is **INITIATION** of therapy, please answer the following questions:

a. What is the patient's diagnosis?

FGF23-Related Hypophosphatemia in Tumor-Induced Osteomalacia (TIO)

i. Is the diagnosis associated with phosphaturic mesenchymal tumors that cannot be curatively resected or localized?  Yes    No

Rickets

i. Does the patient have a diagnosis of X-linked dominant hypophosphatemia rickets or X-linked vitamin D-resistant rickets?  
 Yes    No

ii. Has the diagnosis been confirmed by genetic testing of PHEX (phosphate regulating gene with homology to endopeptidases located on the X chromosome) mutation in the patient?  Yes    No

Other diagnosis (***please specify***): \_\_\_\_\_

b. Is the patient currently taking any oral phosphate or active vitamin D analog supplementation?  Yes\*  No

\*If YES, will the patient discontinue the oral phosphate or vitamin D analog supplementation at least one week prior to starting therapy with Crysivita?  Yes  No

c. Is the fasting serum phosphorus within or above the normal range for age?  Yes  No

YES – this is a PA renewal for CONTINUATION of therapy, please answer the following question:

a. What is the patient's diagnosis?

FGF23-Related Hypophosphatemia in Tumor-Induced Osteomalacia (TIO)

X-Linked Dominant Hypophosphatemic Rickets

X-Linked Hypophosphatemia (XLH)

X-Linked Vitamin D-Resistant Rickets

Other diagnosis (please specify): \_\_\_\_\_

2. Does the prescriber agree to measure serum phosphorus throughout therapy and withhold Crysivita when serum phosphorus is above the reference range for age?  Yes  No
3. Does the patient have severe renal impairment or end stage renal disease (ESRD), defined as eGFR less than 30 mL/min/1.73 m<sup>2</sup>?  Yes  No
4. Will Crysivita be administered by a healthcare provider?  Yes  No

Chart notes are required for the processing of all requests. Please add any other supporting medical information necessary for our review (required)

Coverage will not be provided if the prescribing physician's signature and date are not reflected on this document.

Request for expedited review: I certify that applying the standard review time frame may seriously jeopardize the life or health of the member or the member's ability to regain maximum function

Physician's Name	Physician Signature	Date
<b>Step 2:</b> Checklist	<input type="checkbox"/> Form Completely Filled Out <input type="checkbox"/> Provide chart notes	<input type="checkbox"/> Attach test results
<b>Step 3:</b> Submit	By Fax: BCBSM Specialty Pharmacy Mailbox 1-877-325-5979	By Mail: BCBSM Specialty Pharmacy Program P.O. Box 312320, Detroit, MI 48231-2320

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