Your Child's

Doctor Discussion Guide



If you are considering CRYSVITA® (burosumab-twza) for your child, you may have questions. Take this guide to your next appointment to help start the conversation. Together, you and their doctor can decide if CRYSVITA is right for your child.



What is CRYSVITA?

CRYSVITA is a prescription medicine used to treat adults and children 6 months of age and older with X-linked hypophosphatemia (XLH).

Important Safety Information

You should not take CRYSVITA if:

- You take an oral phosphate supplement and/or a specific form of vitamin D supplement (such as calcitriol, paricalcitol, doxercalciferol, calcifediol).
- Your phosphorus levels from a blood sample are within or above the normal range for age.
- · You have kidney problems.

Please see Important Safety Information continued throughout this guide. Please see the full <u>Prescribing Information for CRYSVITA here</u>.



About CRYSVITA

What is CRYSVITA, and how is it different from other treatments?
Can you tell me about how CRYSVITA was studied in children? What does it mean for my child?
What are the possible side effects?
How is CRYSVITA given, and how often will my child need treatment?

How long does it take before CRYSVITA starts to work?

Is CRYSVITA right for my child?



Accessing CRYSVITA treatment

Will insurance cover CRYSVITA? What do I need to do to start the process for my child?

Are there any financial support programs available for CRYSVITA?



Resources and social support

What resources are available to help support people who are taking CRYSVITA?

Important Safety Information (cont'd)

What is the most important information you should know about CRYSVITA?

Some patients developed allergic reactions (e.g., rash and hives)
while taking CRYSVITA. Your doctor will monitor you for symptoms of
an allergic reaction while you are taking CRYSVITA. Your treatment
may need to be discontinued for serious allergic reactions.

Please see Important Safety Information continued throughout this guide. Please see the full <u>Prescribing Information for CRYSVITA here</u>.



Important Safety Information (cont'd)

What is the most important information you should know about CRYSVITA? (cont'd)

- High levels of phosphorus in the blood have been reported in some patients taking CRYSVITA.
 This may be related to a risk of high calcium levels in the kidneys. Your doctor will collect blood samples to monitor your levels. If you are already taking CRYSVITA, dose interruption and/or dose reduction may be required based on your serum phosphorus levels.
- Administration of CRYSVITA may result in reactions at the injection site, such as hives, reddening
 of the skin, rash, swelling, bruising, pain, severe itching of the skin, and collection of blood
 outside of a blood vessel (i.e., hematoma). Call your doctor if you develop an injection site
 reaction. CRYSVITA may be discontinued if severe injection site reactions occur.

What are the possible side effects of CRYSVITA?

Adverse reactions that were seen in children with XLH are:

Fever - Tooth abscess - Constipation
 Injection site reaction - Dental cavities - Muscle pain

Cough
 Diarrhea
 Pain in arms and legs
 Diarrhea
 Decreased vitamin D levels
 Dizziness
 Nausea

- Headache

• Adverse reactions that were seen in adults with XLH are:

Back pain
 Becreased vitamin D levels
 Muscle spasms
 Phosphorus levels
 Tooth infection
 Constipation
 Muscle spasms
 Phosphorus levels
 increased in the blood

- Restless legs syndrome

 Narrowing of the spaces within the spine is common in adults with XLH, and pressure on the spinal cord has been reported in adults taking CRYSVITA. It is not known if taking CRYSVITA worsens the narrowing of the spaces within the spine or the pressure on the spinal cord.

Before taking CRYSVITA, tell your doctor about all of your medications (including supplements) and medical conditions, including if you:

- Are taking oral phosphate and/or active vitamin D (such as calcitriol, paricalcitol, doxercalciferol, calcifediol).
- Are pregnant, think you may be pregnant, or plan to become pregnant. There is not enough experience to know if CRYSVITA may harm your unborn baby. Report pregnancies to the Kyowa Kirin, Inc. Adverse Event reporting line at 1-844-768-3544.
- Are breastfeeding or plan to breastfeed. There is not enough experience to know if CRYSVITA passes into your breast milk. Talk with your doctor about the best way to feed your baby while you receive CRYSVITA.

While taking CRYSVITA, tell your doctor if you experience:

- · An allergic reaction such as rash or hives
- · A rash, swelling, bruising, or other reaction at the injection site
- New or worsening restless legs syndrome

These are not all the possible side effects of CRYSVITA. Call your doctor for medical advice about side effects.

You may report side effects to the FDA at (800) FDA-1088 or www.fda.gov/medwatch. You may also report side effects to Kyowa Kirin, Inc. at 1-844-768-3544.

For important risk and use information, please see the full Prescribing Information here.



