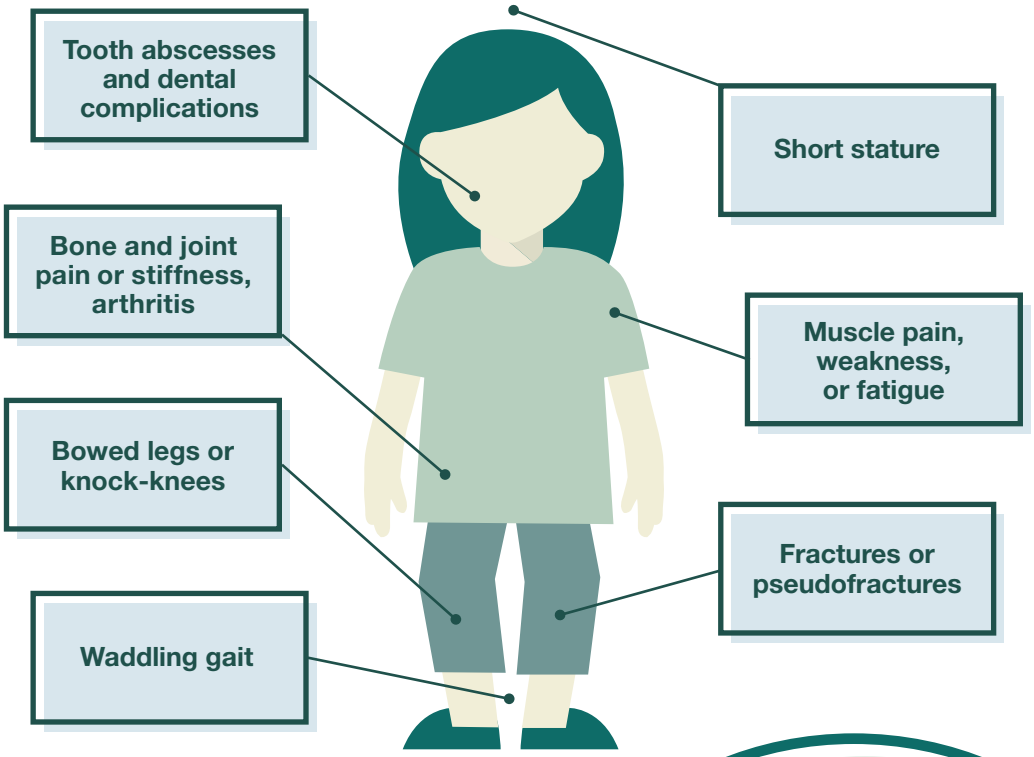
DO YOU HAVE ANY OF THESE SYMPTOMS? CONSIDER TESTING FOR HYPOPHOSPHATEMIA.



WHAT IS HYPOPHOSPHATEMIA?

Hypophosphatemia is a condition in which the levels of phosphorus in the blood are too low. Some forms of the condition are caused by genetic mutations. Symptoms can vary widely even among people with the same condition. The most common genetic form of hypophosphatemia is called X-linked hypophosphatemia (XLH).

A genetic test can help identify hypophosphatemia as the cause of these symptoms and can help your doctor create a care plan based on your or your child's specific needs.

SPEAK WITH YOUR DOCTOR ABOUT GENETIC TESTING FOR HYPOPHOSPHATEMIA

STEPS FOR GETTING TESTED

1

ASK YOUR DOCTOR IF YOU OR YOUR CHILD ARE CANDIDATES FOR TESTING

Your doctor will review your family's health history and your or your child's symptoms. If you or your child meet the program eligibility criteria, the genetic test for hypophosphatemia is provided at no cost to you.

2

PROVIDE A BLOOD OR SALIVA SAMPLE

Your doctor will collect a sample and send it in for testing. Results are confidential and usually available to you and your doctor within 3 weeks after processing begins.

3

DISCUSS THE RESULTS WITH YOUR DOCTOR

It is important to understand how the diagnosis may impact you or your child and your family members. Your doctor can also explain options for managing the condition.

Speak with your doctor about genetic testing for hypophosphatemia. Your doctor can learn more about this sponsored program at invitae.com/hypophosphatemia-test.

© 2019 Ultragenyx Pharmaceutical Inc. All rights reserved.

MRCP-KRN23-00548 04/2019