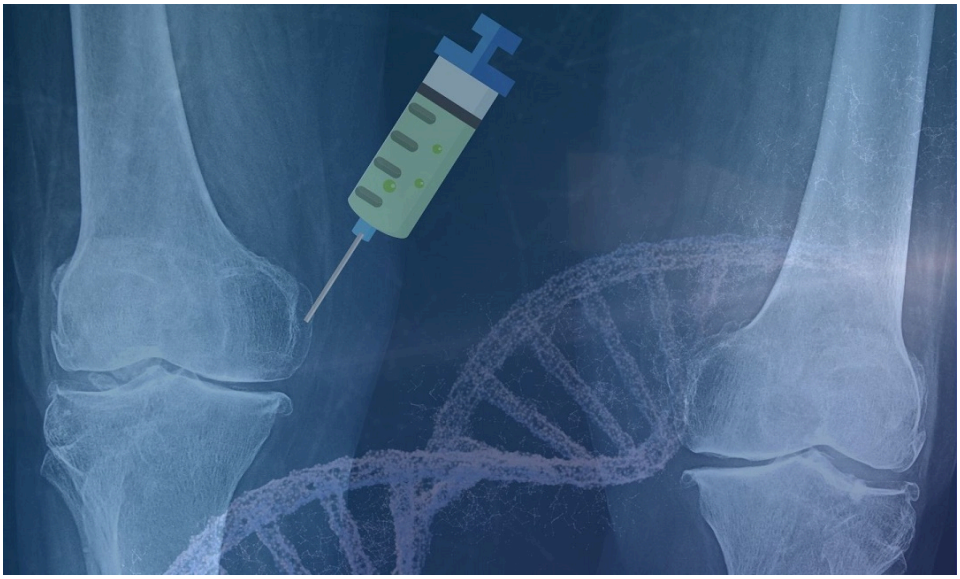
RQMO
REGROUPEMENT QUÉBÉCOIS DES MALADIES ORPHELINES

WEBINAR INVITATION

Are you concerned about X-LINKED HYPOPHOSPHATEMIA?



The Quebec Coalition of Orphan Diseases (Regroupement québécois des maladies orphelines) invites you to the first Canadian webinar about X-linked hypophosphatemia (XLH)!

WHEN?
TUESDAY, JANUARY 28, 2020 FROM 7:30 P.M. TO 8:30 P.M.

WHAT?
You will learn about:

- **XLH, its diagnosis and treatments**
- **The genetic cause of XLH and its hereditary transmission**
- **Resources available in Quebec and Canada**
- **And, how to stay in touch with your community.**

WHO?

Dr. Francis Glorieux (OC, MD, PhD):

Director Emeritus of Research at Shriners Hospital for Children Canada and Professor Emeritus, Pediatrics, Surgery and Human Genetics. He is the founder of the Genetics Unit of Shriners Hospital for Children. He has been recognized for his discoveries in the field of genetic bone diseases and for his contribution to improving orthopedic treatment and rehabilitation measures.

Gail Ouellette (PhD., M. Sc)

Geneticist and genetic counselor. Founder and president of the Quebec Coalition of Orphan Diseases (Regroupement québécois des maladies orphelines)

WHERE? Register for this webinar before January 17, 2020, by sending an email to: inscription@rqmo.org

The information necessary for connecting to the webinar will be provided following your registration.

✦ *A **webinar** is a conference that can be seen and listened to on the web with a computer, tablet or smartphone. At a minimum, speakers must be enabled to listen. Questions may be asked using the keyboard.*

The Quebec Coalition of Orphan Diseases/Regroupement québécois des maladies orphelines

IRARE CENTRE (Information and support centre for rare diseases)

https://rqmo.org/rare-disease-information-and-resource-centre/