DONATE

Let's help one another find our way forward.

Our foundation advocates for people with Galactosemia and their families. We connect families and support networking efforts between clinicians and researchers.

GET INVOLVED

Virtual: Back to School – Teacher Letter, IEP, 504 & More

UPCOMING EVENT – AUGUST 14, 2025

REGISTER

RECENTLY DIAGNOSED?

START HERE

WE NEED YOU!

Advocating for a Treatment

The Galactosemia Foundation is urging the U.S. Food and Drug Administration (FDA) to incorporate the experiences and perspectives of people living with galactosemia in the agency's review of govorestat (AT-007), the potential first-ever treatment for our rare genetic disease.

Understanding Galactosemia

About Galactosemia

Galactosemia is a disorder caused by a genetic mutation that affects how galactose is broken down in the metabolic pathway.

LEARN MORE

Recent Diagnosis

Whether your child was recently diagnosed or just trying to learn more about galactosemia, you've come to the right place.

LEARN MORE

Resources

Tools and resources for the care of your children and loved ones.

LEARN MORE

LIVING WITH GALACTOSEMIA

Newborns

Babies are first diagnosed with galactosemia during the newborn screening process, which is done in all 50 states.

NEWBORNS

Latest News

PRESS RELEASE – NOVEMBER 27

BLOG – FEBRUARY 11

ction Needed: NIH nding Cuts Could pact Galactosemia esearch

PRESS RELEASE – NOVEMBER 27

Govorestat, First and Only Potential Treatment for Classic Galactosemia, Does

U.S. FDA Regarding New Drug

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Not Receive FDA Approval

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Help our fight against Galactosemia today!

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Stay in the loop and get updates.

Sign up for the Foundation's mailing list

SIGN UP TODAY!

Galactosemia Foundation

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