

RAISE YOUR VOICE FOR RARE

Support Patients. Advance Research. Change Lives.



DLG4 SHINE—SHINE Syndrome—is **an ultra-rare genetic condition** affecting fewer than 300 people in the world. In addition to a lack of public awareness and most patients facing delayed diagnoses, there are currently no treatment options.

But with your help, we can change that!

**YOUR
SUPPORT
WILL HELP:**

- **Fund groundbreaking research**
- **Support families with critical resources**
- **Advocate for better care and policies**

DLG4 SHINE is an extremely rare neurodevelopmental disorder characterized predominantly by global developmental delay/intellectual disability of varying severity, autism spectrum disorder, attention deficit hyperactivity disorder, hypotonia, and epilepsy.

The DLG4 SHINE Foundation collaborates with researchers, healthcare professionals, and other organizations to accelerate medical advances focused on the treatment of this rare disease.

YOUR DONATION SUPPORTS:

Patient &
caregiver
support
services

Research
grants &
clinical trial
funding

Legislative
advocacy

Educational
outreach for
medical
professionals



venmo



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dlg4shine.org