

# WANTED

## Stories, Experiences, and Case Examples

***Heartland Regional Genetics Network (HRGN) is seeking the following types of cases for presentation and discussion during sessions of the 2022 Summer Project ECHO series: "The Genetic Testing Journey: Who, What, and How"***

- Patient story of not being able to afford confirmatory testing, but diagnosis and management achieved (e.g. neurofibromatosis, type 1; Noonan syndrome)
- Patient story of successful referral to genetics, completed genetics evaluation, testing, and management plan
- Patient story of challenges in accessing genetics services and/or genetic testing
- Genetic testing pre-authorization successes and failures
- Genetic testing pre-authorization trouble-shooting
- Patient experiences with sponsored testing
- Provider experiences with sponsored testing
- Other cases pertaining to genetic testing

***Please send deidentified cases to [Cara-Vaught@ouhsc.edu](mailto:Cara-Vaught@ouhsc.edu).  
To register for this series [click here](#) or scan the QR code.***

***Thank you for your interest in and contribution  
to the HRGN Project ECHO series!***

