

#### Genomics Research to Elucidate the Genetics of Rare Diseases

### **GREGOR at ASHG 2025**

**Join us at our Ancillary Session:** "Expanded multi-omic data from the rare disease-focused GREGOR Consortium"

October 16th at 11:45 am - 1:15 pm in Room 102AB/Level 1

#### Agenda:

11:45 Gathering and Social time

**11:50 Introductory Remarks,** Dr. Ben Heavner, Senior Research Scientist, University of Washington GREGOR Data Coordinating Center

11:55 "Al-ready cohort allele frequency data from GREGoR using GA4GH Genomic Knowledge Standards", Dr. Alex Wagner (VRS)

12:20 "Genomics at scale with the NHGRI AnVIL", Dr. Mike Schatz (AnVIL)

12:45 "The Latest Data Release from the GREGOR Consortium", Dr. Ben

Heavner, Senior Research Scientist, University of Washington GREGoR Data

**Coordinating Center** 

1:05 Q & A, Social time

# Learn more & register here



#### Don't miss these presentations from our members:

Featured Symposium: "From Data to Diagnosis: Advancing Rare Disease Research through Collaborative Genomics"

October 15th at 8:30 - 10:00 am ET in Room 210C/Level 2

Dr. Marsha Wheeler's Platform talk: "Sharing valuable rare disease and multiomic data through the GREGOR Consortium"

October 15th at 2:15 - 2:30 pm ET in Room 210C/Level 2

See all GREGoR abstracts on our GREGoR at ASHG webpage -

## **Connect with GREGoR online**

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## **GREGoR Publications**

The Marker Paper, GREGoR: Accelerating Genomics for Rare Diseases, has been accepted for publication in Nature!

Since 2021 (the GREGOR Consortium's inception), GREGOR publications have been cited in 2,784 papers - scan the QR code to see our full list of publications.



# The GREGOR Data Variant Browser Data Output Output

#### **Growth of the GREGoR Dataset**

#### participants 10000 short-read DNA (WGS & WES) short-read RNA 8000 long-read DNA (Nanopore) - Iong-read DNA (PacBio) short-read ATAC-seq 6000 4000 2000 U02 U03 U05 U06 U07 U08 U09 U10 U11

## Multi-omic Coverage in GREGoR

