ClinGen at ACMG 2019
Visit us at Booth #529

ClinGen Reception
Wed April 3, 7:30-9:00pm, Washington State Convention Center, Room 606-609

Presentations
Platform Presentations, Washington State Convention Center
- **Wed April 3, 4:30-4:45pm - Room 6C** The Gene Curation Coalition: A global effort to harmonize gene-level resources (M. DiStefano)
- **Fri April 5, 4:45-5:00pm - Room 6E** Myeloid Malignancy Variant Curation Expert Panel: an ASH-sponsored ClinGen Expert Panel to optimize and validate ACMG/AMP variant interpretation guidelines for genes associated with inherited myeloid neoplasms (X. Luo)

Learning Lounge
- **Thurs April 4, 12:45-1:15pm - Learning Lounge 2** Collaboratively Transforming Access to Genetic Testing

Workshop
- **Fri April 5, 7:15-8:30am - Room 3AB** ClinGen Expert Panel Information Workshop (registration required)

Poster Presentations
(Exhibit Halls 4ABCD)
- **Thurs April 4, 10:00-11:30am**
  - 319 The Use and Potential Impact of Diversity Measures in Medical Genetics Practice: Results of a Survey of Clinical Genomics Professionals (A. Popejoy)
  - 467 Structured Narrative Triplets of Functional Assays to Support the Determination of Damaging Effect on Protein Function (A. Lopez Pineda)
  - 681 Contrasting the GAA Variant Spectrum in Newborn Screening and Pompe Diagnostic Testing Populations (B. Seifert)
  - 709 Toward Complete Interpretation of Known RYR1 Variants (J. Johnston)

- **Fri April 5, 10:30am-12:00pm**
  - 178 Clinical Genome Resource (ClinGen) Somatic Cancer Working Group Minimum Variant Level Data (MVLD) v2.0: Extensions Incorporating AMP Somatic Interpretation Guidelines (D. Ritter)
  - 294 ClinGen’s Variant Prioritization Tool (VPT): A Tool for Prioritizing Clinical Variants at Scale (M. Wright)
  - 682 Standards for the Classification and Reporting of Constitutional Copy Number Variants: A ClinGen/ACMG Joint Consensus Recommendation (E. Riggs)
  - 784 Opinions of ClinGen Members on the Use of ClinGen Clinical Validity Classifications in Genetic Testing (J. Goldstein)
  - 808 ClinGen Recurrent Copy Number Variant Annotation File to Aid Clinical Interpretation During Genomic Analysis (J. Herriges)
  - 834 ClinVar Makes Updates Easy: A New Online Form to Update a Single SCV (M. Landrum)