AGENDA

1. Discuss Key Questions Related to Data Collection:
   - What benefits could result from insurer collection of disaggregated demographic data for their policyholders and providers to better understand and then counteract health care disparities?
   - Are there any risks or other concerns that arise with the potential collection of such data?
   - Do any regulatory barriers exist to the collection of such data?

Consumer Remarks: Katie Keith (Out2Enroll), Karen Siegel (Health Equity Solutions) and Silvia Yee (Disability Rights Education and Defense Fund—DREDF)

Industry Remarks: Michelle Jester (AHIP) and Lauren Choi (Blue Cross Blue Shield Association—BCBSA)

Q&A
   - What should the role of insurance regulators be in thinking about this type of data collection? Is there a deliverable the NAIC should work towards?

Consumer Remarks: Katie Keith (Out2Enroll), Karen Siegel (Health Equity Solutions) and Silvia Yee (Disability Rights Education and Defense Fund—DREDF)

Industry Remarks: Michelle Jester (AHIP) and Lauren Choi (Blue Cross Blue Shield Association—BCBSA)

Q&A

2. Discuss Next Steps—Commissioner Jessica K. Altman (PA)

3. Discuss Any Other Matters Brought Before the Workstream—Commissioner Jessica K. Altman (PA)

4. Adjournment