



**Saturday, May 13, 2017**  
**Loudermilk Conference Center**  
**40 Courtland Street NE**  
**Atlanta, Georgia 30303**

Time	Session
8:30 am	<b>Registration</b>
9:00 am	<b>Our Journey: Driving Change Along the Rare Disease Road</b> Lisa Raman, Moderator, Klippel Feil Syndrome Alliance
10:00 am	<b>Turn Hope into Action: Plotting Your Path to Drive Change</b> Nicole Boice, Global Genes Kevin Rohrbach, Global Genes
11:00 am	<b>Discover the Keys to Your Story to Make the Most Impact</b> Alex Van Rees, SmithSolve
12:00 pm	<b>Lunch</b>
1:00 pm	<b>Messages from Members of Congress</b>
1:15 pm	<b>What is on the Docket to Accelerate Change in the Rare Disease Community</b> Julia Jenkins, EveryLife Foundation for Rare Diseases Lisa Schill, EveryLife Foundation for Rare Diseases
2:05 pm	<b>Break</b>
2:20 pm	<b>Public Policy is as Easy as a Phone Call</b> Julia Jenkins, EveryLife Foundation for Rare Diseases Lisa Schill, EveryLife Foundation for Rare Diseases
3:15 pm	<b>Workshop: Your Path Forward</b> Nicole Boice, Global Genes Kevin Rohrbach, Global Genes
4:15 pm	<b>Program Conclusion/Postcard Project</b> Lisa Raman, Moderator, Klippel Feil Syndrome Alliance