1 CC2D2A

This gene encodes a coiled-coil and calcium binding domain protein that appears to play a critical role in cilia formation. Mutations in this gene cause Meckel syndrome type 6, as well as Joubert syndrome type 9. Alternative splicing results in multiple transcript variants.

This gene is not differentially expressed.

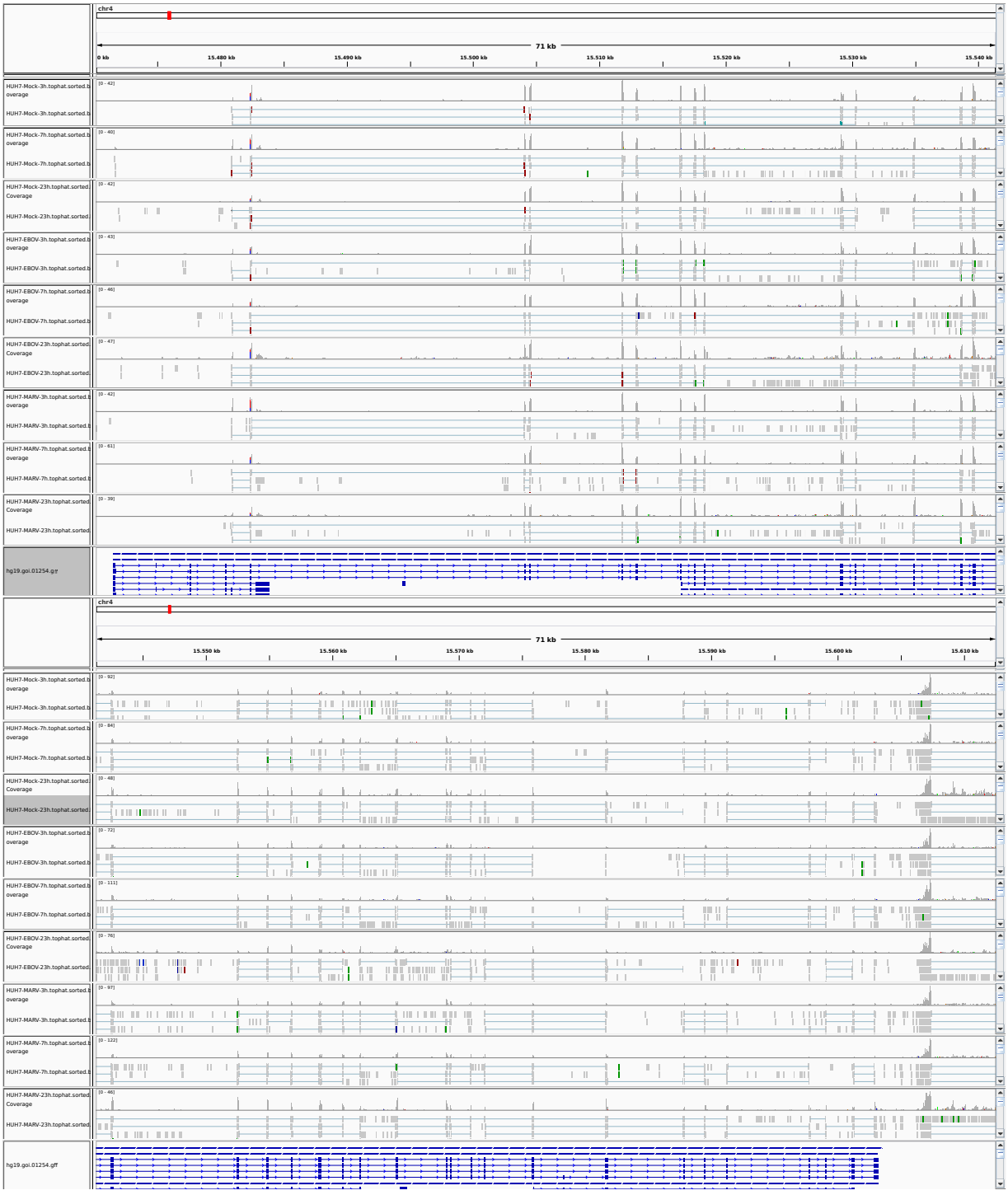

Figure 1: IGV Genome Browser screenshot of gene CC2D2A.

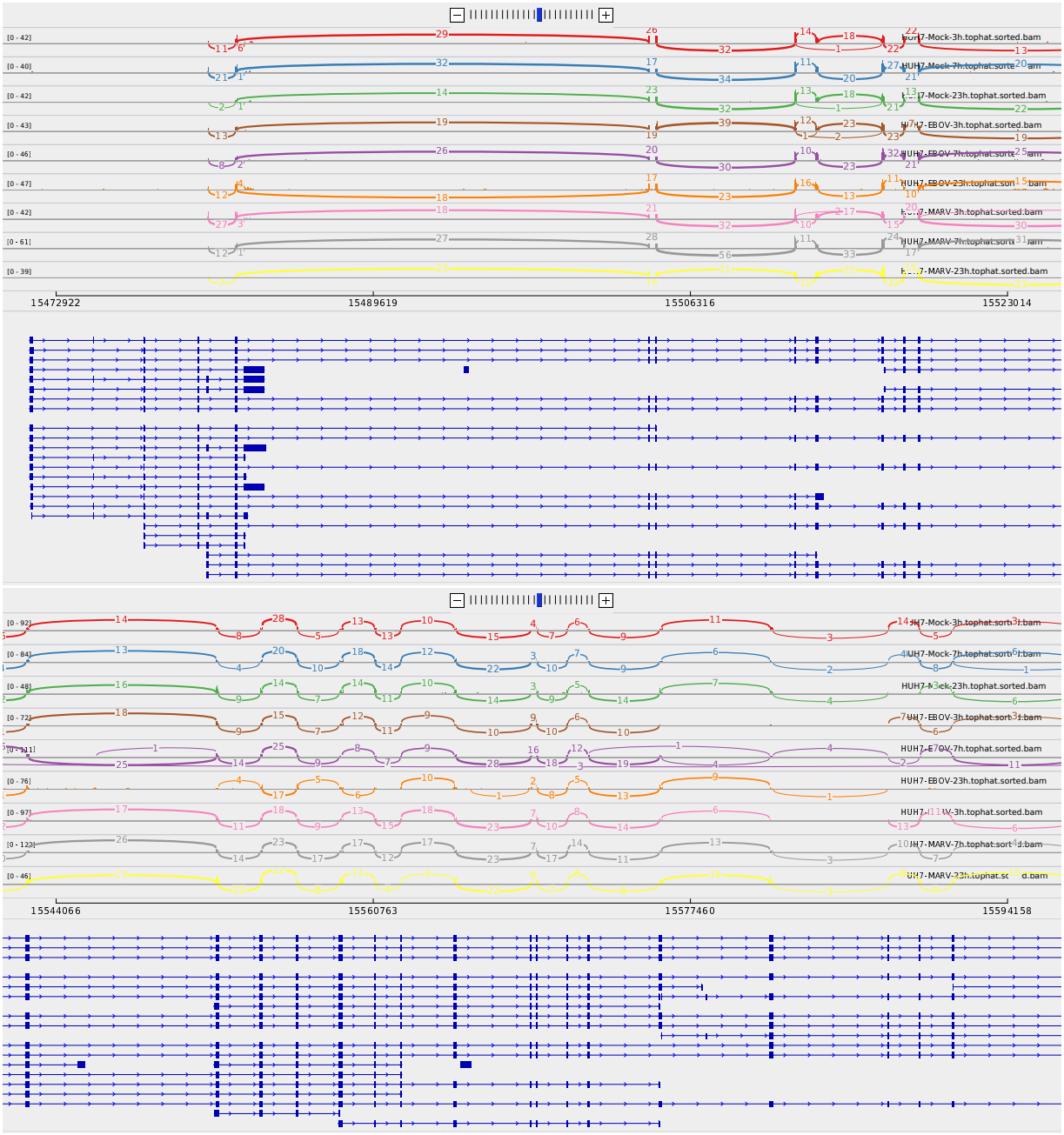

Figure 2: Sashimi plot of gene CC2D2A.

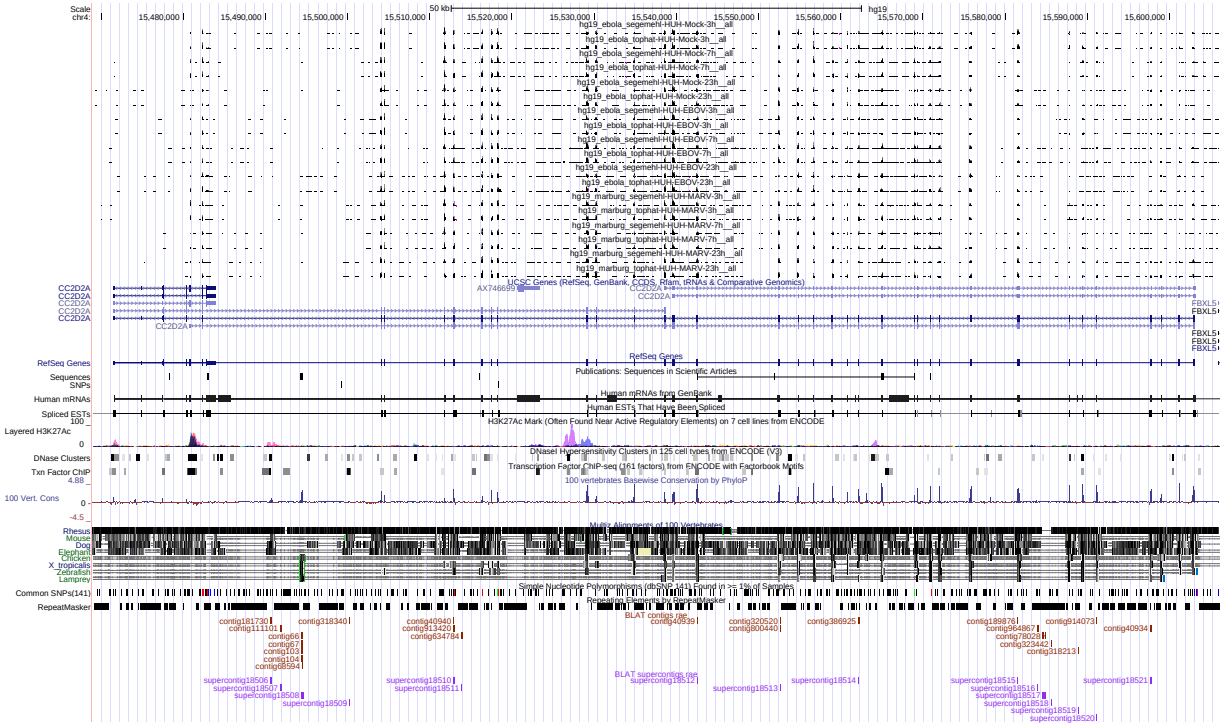

Figure 3: UCSC Genome Browser screenshot of gene CC2D2A.