



**Overall QC Status:** PASS  
**Sample QC Status:** PASS  
**Fusion QC Status:** PASS  
**Variations QC Status:** PASS  
**Job:** 20211119 AL21001 [4713]  
**Type:** RNA FusionRNA SNP/InDel Targeted Mutations: Archer  
 Comprehensive Targets NIH v1.3.1 2 Include Non-Targeted  
 Variants: No  
**Software Version:** Suite\_Analysis\_v6.2.7  
**Analysis Date:** 19-Nov-2021 5:02  
**Report Creator:** mpvghtpe@gmail.com  
**Report Date:** 20-Jun-2023 21:45

## Statistics

### Molecular Barcode Statistics

Total Fragments	Fragments with Complete Adapter	Number of Reads After Trimming Adapters
658,900	609,832	574,819

### Read Statistics

Type	Total Fragments (# / %)	Mapped (# / %)	Pass Alignment Filter (%)	On Target (%)
All Fragments	572,903 / 100.0	572,903 / 100.0	100.0	98.0
Unique Fragments	151,531 / 26.4	151,531 / 100.0	100.0	98.4

### DNA/RNA Statistics

Type	DNA Reads (# / %)	RNA Reads (# / %)	Ambiguous Reads (# / %)
All Fragments	25,603.0 / 4.6	459,478.0 / 81.9	76,143.0 / 13.6
Molecular Bins	9,279.0 / 6.2	111,323.0 / 74.7	28,468.0 / 19.1
Average Molecular Bins per GSP2	56.93	682.96	174.65
Unique Start Sites	4,527.0 / 24.8	12,043.0 / 65.9	4,325.0 / 23.7
Average Unique Start Sites per GSP2	28.24	92.83	27.89
Average Unique Start Sites per GSP2 Control	42.75	208.5	83.25

QC Statistics

Avg. Unique DNA And Ambiguous Start Sites Per GSP2	Avg. Unique RNA Start Sites Per GSP2 Control
55.69	208.5

Miscellaneous Statistics

On Target Deduplication Ratio
3.76:1

DNA/RNA Fragment Lengths

DNA Median Fragment Length	DNA Mean Fragment Length	RNA Median Fragment Length	RNA Mean Fragment Length
131.0	142.6	140.0	146.8

**Reportable Variants**

*None Found*


# Reportable Isoforms


- ☒ Passed all strong-evidence filters


☒ Likely off-target mispriming event


☒ Exact breakpoint known



☒ Cross contamination


☒ User-annotated false positive
-  Known fusion partners in Archer Quiver™


 Percent GSP2 reads below threshold


 Fusion expression imbalance

 Low confidence

 User-annotated true positive
-  Intronic fusion

 Not enough unique start sites

 Transcriptional readthrough event

 Known ensembl paralogue

Fusion: TPR → ROS1		
<div>Filters: <input checked="" type="checkbox"/></div> <div>GSP2: ROS1_chr6_117642501_21+_A1_GSP2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 67 (60.91%)</div> <div>Start Sites: 38</div>	<div>Segments</div> <div>chr1:186337114→186337018 TPR(-) NM_003292.2, exon:4</div> <div>chr6:117642557→117642502 ROS1(-) NM_002944.2, exon:35</div>