



Overall QC Status: PASS  
Sample QC Status: PASS  
Fusion QC Status: PASS  
Variations QC Status: PASS  
Job: 20240516\_AS24007\_AS24008 [3535]  
Type: RNA FusionRNA SNP/InDel Targeted Mutations: Archer  
Comprehensive Targets NIH v1.3.1 Include Non-Targeted  
Variants: No  
Software Version: Suite\_Analysis\_v6.2.7  
Analysis Date: 16-May-2024 5:16  
Report Creator: mpvghtpe@gmail.com  
Report Date: 16-May-2024 19:18

Statistics

Molecular Barcode Statistics

Total Fragments	Fragments with Complete Adapter	Number of Reads After Trimming Adapters
3,500,000	3,235,706	3,091,946

Read Statistics

Type	Total Fragments (# / %)	Mapped (# / %)	Pass Alignment Filter (%)	On Target (%)
All Fragments	3,072,993 / 100.0	3,072,993 / 100.0	100.0	98.2
Unique Fragments	555,722 / 18.1	555,722 / 100.0	100.0	98.5

DNA/RNA Statistics

Type	DNA Reads (# / %)	RNA Reads (# / %)	Ambiguous Reads (# / %)
All Fragments	233,977.0 / 7.8	2,416,829.0 / 80.1	367,197.0 / 12.2
Molecular Bins	53,651.0 / 9.8	391,484.0 / 71.5	102,477.0 / 18.7
Average Molecular Bins per GSP2	49.4	360.48	94.36
Unique Start Sites	24,693.0 / 27.1	56,787.0 / 62.3	21,032.0 / 23.1
Average Unique Start Sites per GSP2	23.33	68.14	21.59
Average Unique Start Sites per GSP2 Control	26.67	156.67	27.25

QC Statistics

Avg. Unique DNA And Ambiguous Start Sites Per GSP2	Avg. Unique RNA Start Sites Per GSP2 Control
44.73	156.67

Miscellaneous Statistics

On Target Deduplication Ratio
5.51:1

DNA/RNA Fragment Lengths

DNA Median Fragment Length	DNA Mean Fragment Length	RNA Median Fragment Length	RNA Mean Fragment Length
126.0	140.7	123.0	132.6

**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: COL1A1 → PDGFB		
<div>Filters: <input checked="" type="checkbox"/> <input checked="" type="checkbox"/></div> <div>GSP2: PDGFB_chr22_39631845_24+_A1_GSP 2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 3638 (88.82%)</div> <div>Start Sites: 203</div>	<div>Segments</div> <div>chr17:48266636→48266529 COL1A1(-) NM_000088.3, exon:40</div> <div>chr22:39631879→39631783 PDGFB(-) NM_002608.3, exon:2</div>

Fusion: SEMA5A → TERT		
<div>Filters: <input checked="" type="checkbox"/></div> <div>GSP2: TERT_chr5_1282672_20+_A1_GSP2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 380 (88.79%)</div> <div>Start Sites: 112</div>	<div>Segments</div> <div>chr5:9224999→9224786 SEMA5A(-) NM_003966.2, exon:8</div> <div>chr5:1282739→1282673 TERT(-) NM_198253.2, exon:3</div>