

Overall QC Status: PASS
Sample QC Status: PASS
Fusion QC Status: PASS
Variations QC Status: PASS
Job: 20211021 AS21004 AS21005 [3440]
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: 21-Oct-2021 6:34
Report Creator: mpvghtpe@gmail.com
Report Date: 19-Jun-2023 2:33

Statistics

Molecular Barcode Statistics

Total Fragments	Fragments with Complete Adapter	Number of Reads After Trimming Adapters
3,500,000	3,280,575	3,073,182

Read Statistics

Type	Total Fragments (# / %)	Mapped (# / %)	Pass Alignment Filter (%)	On Target (%)
All Fragments	3,056,874 / 100.0	3,056,874 / 100.0	100.0	97.8
Unique Fragments	754,179 / 24.7	754,179 / 100.0	100.0	98.6

DNA/RNA Statistics

Type	DNA Reads (# / %)	RNA Reads (# / %)	Ambiguous Reads (# / %)
All Fragments	258,484.0 / 8.6	2,222,282.0 / 74.3	509,188.0 / 17.0
Molecular Bins	82,358.0 / 11.1	497,104.0 / 66.9	163,801.0 / 22.0
Average Molecular Bins per GSP2	140.3	846.86	279.05
Unique Start Sites	28,404.0 / 34.7	48,258.0 / 58.9	19,391.0 / 23.7
Average Unique Start Sites per GSP2	51.79	119.97	39.76
Average Unique Start Sites per GSP2 Control	55.67	242.17	45.83

QC Statistics

Avg. Unique DNA And Ambiguous Start Sites Per GSP2	Avg. Unique RNA Start Sites Per GSP2 Control
90.31	242.17

Miscellaneous Statistics

On Target Deduplication Ratio
4.02:1

DNA/RNA Fragment Lengths

DNA Median Fragment Length	DNA Mean Fragment Length	RNA Median Fragment Length	RNA Mean Fragment Length
146.0	154.5	139.0	146.7

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: SS18 → SSX2		
<div>Filters: </div> <div>GSP2: SS18_chr18_23612369_25_-_A1_GSP2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 936 (21.01%)</div> <div>Start Sites: 247</div>	<div>Segments</div> <div>chr18:23612496→23612363 SS18(-) NM_005637.3, exon:9</div> <div>chrX:52729628→52729493 SSX2(-) NM_003147.5, exon:6</div>