



Overall QC Status: PASS
Sample QC Status: PASS
Fusion QC Status: PASS
Variations QC Status: PASS

Job:
20230804_AS23036_AS23037_AS23038_AS23040_AS23041
[3486]

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: 03-Aug-2023 20:42

Report Creator: mpvghtpe@gmail.com

Report Date: 04-Aug-2023 0:00

Statistics

Molecular Barcode Statistics

Total Fragments	Fragments with Complete Adapter	Number of Reads After Trimming Adapters
1,413,016	1,298,045	1,207,302

Read Statistics

Type	Total Fragments (# / %)	Mapped (# / %)	Pass Alignment Filter (%)	On Target (%)
All Fragments	1,201,925 / 100.0	1,201,925 / 100.0	100.0	99.0
Unique Fragments	497,530 / 41.4	497,530 / 100.0	100.0	99.1

DNA/RNA Statistics

Type	DNA Reads (# / %)	RNA Reads (# / %)	Ambiguous Reads (# / %)
All Fragments	114,659.0 / 9.6	897,715.0 / 75.4	177,839.0 / 14.9
Molecular Bins	53,138.0 / 10.8	348,623.0 / 70.7	91,331.0 / 18.5
Average Molecular Bins per GSP2	80.63	529.02	138.59
Unique Start Sites	19,501.0 / 32.2	35,842.0 / 59.2	14,672.0 / 24.2
Average Unique Start Sites per GSP2	30.79	73.78	25.45

Average Unique Start Sites per GSP2 Control	23.08	118.83	16.42
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QC Statistics

Avg. Unique DNA And Ambiguous Start Sites Per GSP2	Avg. Unique RNA Start Sites Per GSP2 Control
55.83	118.83

Miscellaneous Statistics

On Target Deduplication Ratio
2.41:1

DNA/RNA Fragment Lengths

DNA Median Fragment Length	DNA Mean Fragment Length	RNA Median Fragment Length	RNA Mean Fragment Length
129.0	140.1	126.0	131.8

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: EWSR1 → WT1		
<div>Filters: </div> <div>GSP2: EWSR1_chr22_29683087_28+_A1_GSP 2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 27 (1.38%)</div> <div>Start Sites: 22</div>	<div>Segments</div> <div>chr22:29683088→29683123 EWSR1(+) NM_005243.3, exon:7</div> <div>chr11:32414301→32414212 WT1(-) NM_000378.4, exon:7</div>