



**Overall QC Status:** PASS  
**Sample QC Status:** PASS  
**Fusion QC Status:** PASS  
**Variations QC Status:** PASS  
**Job:** 20230609\_AS23027 [3476]  
**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:** 08-Jun-2023 23:28  
**Report Creator:** mpvghtpe@gmail.com  
**Report Date:** 08-Jun-2023 23:34

## Statistics

### Molecular Barcode Statistics

Total Fragments	Fragments with Complete Adapter	Number of Reads After Trimming Adapters
3,500,000	3,279,137	3,087,027

### Read Statistics

Type	Total Fragments (# / %)	Mapped (# / %)	Pass Alignment Filter (%)	On Target (%)
All Fragments	3,070,112 / 100.0	3,070,112 / 100.0	100.0	99.0
Unique Fragments	707,422 / 23.0	707,422 / 100.0	100.0	99.1

### DNA/RNA Statistics

Type	DNA Reads (# / %)	RNA Reads (# / %)	Ambiguous Reads (# / %)
All Fragments	341,437.0 / 11.2	2,227,629.0 / 73.3	471,245.0 / 15.5
Molecular Bins	87,583.0 / 12.5	454,064.0 / 64.8	159,107.0 / 22.7
Average Molecular Bins per GSP2	80.65	418.11	146.51
Unique Start Sites	33,677.0 / 31.3	63,041.0 / 58.6	25,748.0 / 24.0
Average Unique Start Sites per GSP2	32.42	76.42	27.49
Average Unique Start Sites per GSP2 Control	38.25	161.25	28.17

QC Statistics

Avg. Unique DNA And Ambiguous Start Sites Per GSP2	Avg. Unique RNA Start Sites Per GSP2 Control
59.5	161.25

Miscellaneous Statistics

On Target Deduplication Ratio
4.34:1

DNA/RNA Fragment Lengths

DNA Median Fragment Length	DNA Mean Fragment Length	RNA Median Fragment Length	RNA Mean Fragment Length
126.0	138.9	127.0	136.1

**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 → NFATC2		
<div>Filters: </div> <div>GSP2: NFATC2_chr20_50133429_23+_A1_GS P2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 2634 (59.58%)</div> <div>Start Sites: 318</div>	<div>Segments</div> <div>chr22:29684595→29684775 EWSR1(+) NM_005243.3, exon:8</div> <div>chr20:50133494→50133323 NFATC2(-) NM_012340.4, exon:3</div>

Fusion: EWSR1 → NFATC2		
<div>Filters: </div> <div>GSP2: NFATC2_chr20_50133429_23+_A1_GS P2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 1173 (99.66%)</div> <div>Start Sites: 169</div>	<div>Segments</div> <div>chr22:29686714→29686954 EWSR1(+) NM_005243.3, intron:8</div> <div>chr20:50133608→50133495 NFATC2(-) NM_012340.4, intron:2</div>

Fusion: EWSR1 → NFATC2		
<div>Filters: </div> <div>GSP2: NFATC2_chr20_50133429_23+_A1_GS P2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 79 (0.89%)</div> <div>Start Sites: 48</div>	<div>Segments</div> <div>chr22:29682951→29683123 EWSR1(+) NM_005243.3, exon:7</div> <div>chr20:50133494→50133430 NFATC2(-) NM_012340.4, exon:3</div>

Fusion: EWSR1 → NFATC2		
<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: 28 (0.31%)</div> <div>Start Sites: 15</div>	<div>Segments</div> <div>chr22:29683088→29683122 EWSR1(+) NM_005243.3, exon:7</div> <div>chr20:50133494→50133390 NFATC2(-) NM_012340.4, exon:3</div>