

Overall QC Status: PASS

Sample QC Status: PASS

Fusion QC Status: PASS

Variations QC Status: PASS

Job: 20230628_AS23025_AS23031_AS23032 [3478]

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: 28-Jun-2023 4:07

Report Creator: mpvghtpe@gmail.com

Report Date: 29-Jun-2023 0:22

Statistics

Molecular Barcode Statistics

Total Fragments	Fragments with Complete Adapter	Number of Reads After Trimming Adapters
3,460,043	3,279,846	3,110,884

Read Statistics

Type	Total Fragments (# / %)	Mapped (# / %)	Pass Alignment Filter (%)	On Target (%)
All Fragments	3,090,262 / 100.0	3,090,262 / 100.0	100.0	98.5
Unique Fragments	491,230 / 15.9	491,230 / 100.0	100.0	98.6

DNA/RNA Statistics

Type	DNA Reads (# / %)	RNA Reads (# / %)	Ambiguous Reads (# / %)
All Fragments	185,803.0 / 6.1	2,351,986.0 / 77.3	505,028.0 / 16.6
Molecular Bins	37,187.0 / 7.7	340,184.0 / 70.3	106,864.0 / 22.1
Average Molecular Bins per GSP2	34.24	313.24	98.4
Unique Start Sites	20,767.0 / 22.0	60,807.0 / 64.5	24,861.0 / 26.4
Average Unique Start Sites per GSP2	19.61	71.2	26.2
Average Unique Start Sites per GSP2 Control	31.0	193.92	39.5

QC Statistics

Avg. Unique DNA And Ambiguous Start Sites Per GSP2	Avg. Unique RNA Start Sites Per GSP2 Control
45.56	193.92

Miscellaneous Statistics

On Target Deduplication Ratio
6.28:1

DNA/RNA Fragment Lengths

DNA Median Fragment Length	DNA Mean Fragment Length	RNA Median Fragment Length	RNA Mean Fragment Length
131.0	142.4	133.0	141.4

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: WWTR1 → CAMTA1		
<div>Filters:  </div> <div>GSP2: WWTR1_chr3_149290656_21_-_A1_GSP 2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 1134 (25.78%)</div> <div>Start Sites: 249</div>	<div>Segments</div> <div>chr3:149290787→149290651 WWTR1(-) NM_015472.4, exon:3</div> <div>chr1:7723413→7723644 CAMTA1(+) NM_015215.3, exon:9</div>

Fusion: WWTR1 → CAMTA1		
<div>Filters:  </div> <div>GSP2: WWTR1_chr3_149290656_21_-_A1_GSP 2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 55 (1.22%)</div> <div>Start Sites: 40</div>	<div>Segments</div> <div>chr3:149290787→149290651 WWTR1(-) NM_015472.4, exon:3</div> <div>chr1:7730971→7731097 CAMTA1(+) NM_015215.3, exon:10</div>