

Overall QC Status: PASS**Sample QC Status:** PASS**Fusion QC Status:** PASS**Variations QC Status:** PASS**Job:** 20230330_AL23002 [3453]**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:** 29-Mar-2023 20:15**Report Creator:** mpvghtpe@gmail.com**Report Date:** 29-Mar-2023 22:01

Statistics

Molecular Barcode Statistics

Total Fragments	Fragments with Complete Adapter	Number of Reads After Trimming Adapters
1,038,846	964,579	913,504

Read Statistics

Type	Total Fragments (# / %)	Mapped (# / %)	Pass Alignment Filter (%)	On Target (%)
All Fragments	909,738 / 100.0	909,738 / 100.0	100.0	96.3
Unique Fragments	241,182 / 26.5	241,182 / 100.0	100.0	97.3

DNA/RNA Statistics

Type	DNA Reads (# / %)	RNA Reads (# / %)	Ambiguous Reads (# / %)
All Fragments	36,704.0 / 4.2	738,550.0 / 84.3	100,560.0 / 11.5
Molecular Bins	12,501.0 / 5.3	180,698.0 / 77.0	41,473.0 / 17.7
Average Molecular Bins per GSP2	76.69	1,108.58	254.44
Unique Start Sites	5,920.0 / 25.9	15,176.0 / 66.5	5,792.0 / 25.4
Average Unique Start Sites per GSP2	36.93	125.23	37.79
Average Unique Start Sites per GSP2 Control	41.12	207.62	84.38

QC Statistics

Avg. Unique DNA And Ambiguous Start Sites Per GSP2	Avg. Unique RNA Start Sites Per GSP2 Control
73.7	207.62

Miscellaneous Statistics

On Target Deduplication Ratio
3.73:1

DNA/RNA Fragment Lengths

DNA Median Fragment Length	DNA Mean Fragment Length	RNA Median Fragment Length	RNA Mean Fragment Length
138.0	147.4	145.0	150.6

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: TPM3 → NTRK1		
<div>Filters:  </div> <div>GSP2: NTRK1_chr1_156844367_21_-_A1_GSP 2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 2022 (86.23%)</div> <div>Start Sites: 219</div>	<div>Segments</div> <div>chr1:154142945→154142876 TPM3(-) NM_153649.3, exon:7</div> <div>chr1:156844363→156844418 NTRK1(+) NM_002529.3, exon:10</div>

Fusion: TPM3 → NTRK1		
<div>Filters:  </div> <div>GSP2: NTRK1_chr1_156844367_21_-_A1_GSP 2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 34 (97.14%)</div> <div>Start Sites: 26</div>	<div>Segments</div> <div>chr1:154142875→154142819 TPM3(-) NM_152263.3, intron:8</div> <div>chr1:156844312→156844362 NTRK1(+) NM_002529.3, intron:9</div>

Fusion: TPM3 → NTRK1		
<div>Filters:  </div> <div>GSP2: NTRK1_chr1_156845331_19_-_A1_GSP 2</div> <div>Mutation Classification: Undefined</div> <div>Is Artifact: no</div>	<div>Reads: 16 (0.74%)</div> <div>Start Sites: 11</div>	<div>Segments</div> <div>chr1:154142944→154142876 TPM3(-) NM_152263.3, exon:8</div> <div>chr1:156844363→156844418 NTRK1(+) NM_002529.3, exon:10</div>